



The University of Manchester

The University of Manchester Research

Improving the Quality of Dentistry (IQuaD): a cluster factorial randomised controlled trial comparing the effectiveness and cost-benefit of oral hygiene advice and/or periodontal instrumentation with routine care for the prevention and management of periodontal disease in dentate adults attending dental primary care

10.3310/hta22380

**Document Version** 

Accepted author manuscript

Link to publication record in Manchester Research Explorer

Citation for published version (APA):
Ramsay, C. R., Clarkson, J. E., Duncan, A., Lamont, T. J., Heasman, P. A., Boyers, D., Goulão, B., Bonetti, D., Bruce, R., Gouick, J., Heasman, L., Lovelock-hempleman, L. A., Macpherson, L. E., Mccracken, G. I., Mcdonald, A. M., Mclaren-neil, F., Mitchell, F. E., Norrie, J. D., Van Der Pol, M., ... Young, L. (2018). Improving the Quality of Dentity (IQuaD): a cluster factorial radiation mixed controlled trial company for the prevention and management of oral hygiene advice and/or periodontal instrumentation with routine care for the prevention and management of periodontal disease in dentate adults attending dental primary care. Health Technology Assessment. 22(38), 1-144. https://doi.org/10.3310/hta22380

Published in:

Health Technology Assessment

Citing this paper

Please note that where the full-text provided on Manchester Research Explorer is the Author Accepted Manuscript or Proof version this may differ from the final Published version. If citing, it is advised that you check and use the publisher's definitive version.

**ACCESS** 

General rights

Copyright and moral rights for the publications made accessible in the Research Explorer are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

Takedown policy

If you believe that this document breaches copyright please refer to the University of Manchester's Takedown Procedures [http://man.ac.uk/04Y6Bo] or contact uml.scholarlycommunications@manchester.ac.uk providing relevant details, so we can investigate your claim.

## **HEALTH TECHNOLOGY ASSESSMENT**

VOLUME 22 ISSUE 38 JULY 2018 ISSN 1366-5278

Improving the Quality of Dentistry (IQuaD): a cluster factorial randomised controlled trial comparing the effectiveness and cost-benefit of oral hygiene advice and/or periodontal instrumentation with routine care for the prevention and management of periodontal disease in dentate adults attending dental primary care

Craig R Ramsay, Jan E Clarkson, Anne Duncan, Thomas J Lamont, Peter A Heasman, Dwayne Boyers, Beatriz Goulão, Debbie Bonetti, Rebecca Bruce, Jill Gouick, Lynne Heasman, Laura A Lovelock-Hempleman, Lorna E Macpherson, Giles I McCracken, Alison M McDonald, Fiona McLaren-Neil, Fiona E Mitchell, John DT Norrie, Marjon van der Pol, Kirsty Sim, James G Steele, Alex Sharp, Graeme Watt, Helen V Worthington and Linda Young



Improving the Quality of Dentistry (IQuaD): a cluster factorial randomised controlled trial comparing the effectiveness and cost-benefit of oral hygiene advice and/or periodontal instrumentation with routine care for the prevention and management of periodontal disease in dentate adults attending dental primary care

Craig R Ramsay, 1\* Jan E Clarkson, 2 Anne Duncan, 1
Thomas J Lamont, 3 Peter A Heasman, 4
Dwayne Boyers, 5 Beatriz Goulão, 1 Debbie Bonetti, 2
Rebecca Bruce, 1 Jill Gouick, 2 Lynne Heasman, 4
Laura A Lovelock-Hempleman, 2 Lorna E Macpherson, 2
Giles I McCracken, 4 Alison M McDonald, 1
Fiona McLaren-Neil, 2 Fiona E Mitchell, 2
John DT Norrie, 1 Marjon van der Pol, 5 Kirsty Sim, 2
James G Steele, 4 Alex Sharp, 4 Graeme Watt, 2
Helen V Worthington 6 and Linda Young 7

<sup>1</sup>Health Services Research Unit, University of Aberdeen, Aberdeen, UK
<sup>2</sup>Dental Health Services Research Unit, University of Dundee, Dundee, UK
<sup>3</sup>Dundee Dental School, University of Dundee, Dundee, UK
<sup>4</sup>The Dental School, Newcastle University, Newcastle upon Tyne, UK
<sup>5</sup>Health Economics Research Unit, University of Aberdeen, Aberdeen, UK
<sup>6</sup>The School of Dentistry, University of Manchester, Manchester, UK
<sup>7</sup>NHS Education for Scotland, Edinburgh, UK

<sup>\*</sup>Corresponding author

**Declared competing interests of authors:** Jan E Clarkson reports grants from the National Institute for Health Research (NIHR) Health Technology Assessment (HTA) programme during the conduct of the study. Alison M McDonald reports grants from the NIHR HTA programme during the conduct of the study. John DT Norrie reports grants from the University of Aberdeen and grants from the University of Glasgow outside the submitted work. He was a member of the NIHR HTA Commissioning Board (2010–16), is currently a member of the NIHR Editorial Board (2015–present) and is currently the deputy chairperson of the NIHR HTA General Board (2016–present). Marjon van der Pol reports grants from the NIHR HTA programme during the conduct of the study.

Published July 2018 DOI: 10.3310/hta22380

This report should be referenced as follows:

Ramsay CR, Clarkson JE, Duncan A, Lamont TJ, Heasman PA, Boyers D, *et al.* Improving the Quality of Dentistry (IQuaD): a cluster factorial randomised controlled trial comparing the effectiveness and cost–benefit of oral hygiene advice and/or periodontal instrumentation with routine care for the prevention and management of periodontal disease in dentate adults attending dental primary care. *Health Technol Assess* 2018;**22**(38).

Health Technology Assessment is indexed and abstracted in Index Medicus/MEDLINE, Excerpta Medica/EMBASE, Science Citation Index Expanded (SciSearch®) and Current Contents®/ Clinical Medicine.

#### HTA/HTA TAR

## **Health Technology Assessment**

ISSN 1366-5278 (Print)

ISSN 2046-4924 (Online)

Impact factor: 4.236

Health Technology Assessment is indexed in MEDLINE, CINAHL, EMBASE, The Cochrane Library and the Clarivate Analytics Science Citation Index

This journal is a member of and subscribes to the principles of the Committee on Publication Ethics (COPE) (www.publicationethics.org/).

Editorial contact: journals.library@nihr.ac.uk

The full HTA archive is freely available to view online at www.journalslibrary.nihr.ac.uk/hta. Print-on-demand copies can be purchased from the report pages of the NIHR Journals Library website: www.journalslibrary.nihr.ac.uk

#### Criteria for inclusion in the Health Technology Assessment journal

Reports are published in *Health Technology Assessment* (HTA) if (1) they have resulted from work for the HTA programme, and (2) they are of a sufficiently high scientific quality as assessed by the reviewers and editors.

Reviews in *Health Technology Assessment* are termed 'systematic' when the account of the search appraisal and synthesis methods (to minimise biases and random errors) would, in theory, permit the replication of the review by others.

#### **HTA** programme

The HTA programme, part of the National Institute for Health Research (NIHR), was set up in 1993. It produces high-quality research information on the effectiveness, costs and broader impact of health technologies for those who use, manage and provide care in the NHS. 'Health technologies' are broadly defined as all interventions used to promote health, prevent and treat disease, and improve rehabilitation and long-term care.

The journal is indexed in NHS Evidence via its abstracts included in MEDLINE and its Technology Assessment Reports inform National Institute for Health and Care Excellence (NICE) guidance. HTA research is also an important source of evidence for National Screening Committee (NSC) policy decisions.

For more information about the HTA programme please visit the website: http://www.nets.nihr.ac.uk/programmes/hta

#### This report

The research reported in this issue of the journal was funded by the HTA programme as project number 09/01/45. The contractual start date was in April 2011. The draft report began editorial review in February 2017 and was accepted for publication in September 2017. The authors have been wholly responsible for all data collection, analysis and interpretation, and for writing up their work. The HTA editors and publisher have tried to ensure the accuracy of the authors' report and would like to thank the reviewers for their constructive comments on the draft document. However, they do not accept liability for damages or losses arising from material published in this report.

This report presents independent research funded by the National Institute for Health Research (NIHR). The views and opinions expressed by authors in this publication are those of the authors and do not necessarily reflect those of the NHS, the NIHR, NETSCC, the HTA programme or the Department of Health and Social Care. If there are verbatim quotations included in this publication the views and opinions expressed by the interviewees are those of the interviewees and do not necessarily reflect those of the authors, those of the NHS, the NIHR, NETSCC, the HTA programme or the Department of Health and Social Care.

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

Published by the NIHR Journals Library (www.journalslibrary.nihr.ac.uk), produced by Prepress Projects Ltd, Perth, Scotland (www.prepress-projects.co.uk).

## **NIHR Journals Library Editor-in-Chief**

Professor Tom Walley Director, NIHR Evaluation, Trials and Studies and Director of the EME Programme, UK

## **NIHR Journals Library Editors**

**Professor Ken Stein** Chair of HTA and EME Editorial Board and Professor of Public Health, University of Exeter Medical School, UK

Professor Andrée Le May Chair of NIHR Journals Library Editorial Group (HS&DR, PGfAR, PHR journals)

Dr Martin Ashton-Key Consultant in Public Health Medicine/Consultant Advisor, NETSCC, UK

**Professor Matthias Beck** Professor of Management, Cork University Business School, Department of Management and Marketing, University College Cork, Ireland

Dr Tessa Crilly Director, Crystal Blue Consulting Ltd, UK

Dr Eugenia Cronin Senior Scientific Advisor, Wessex Institute, UK

Dr Peter Davidson Director of the NIHR Dissemination Centre, University of Southampton, UK

Ms Tara Lamont Scientific Advisor, NETSCC, UK

**Dr Catriona McDaid** Senior Research Fellow, York Trials Unit, Department of Health Sciences, University of York, UK

Professor William McGuire Professor of Child Health, Hull York Medical School, University of York, UK

Professor Geoffrey Meads Professor of Wellbeing Research, University of Winchester, UK

Professor John Norrie Chair in Medical Statistics, University of Edinburgh, UK

Professor John Powell Consultant Clinical Adviser, National Institute for Health and Care Excellence (NICE), UK

**Professor James Raftery** Professor of Health Technology Assessment, Wessex Institute, Faculty of Medicine, University of Southampton, UK

Dr Rob Riemsma Reviews Manager, Kleijnen Systematic Reviews Ltd, UK

Professor Helen Roberts Professor of Child Health Research, UCL Great Ormond Street Institute of Child Health, UK

**Professor Jonathan Ross** Professor of Sexual Health and HIV, University Hospital Birmingham, UK

**Professor Helen Snooks** Professor of Health Services Research, Institute of Life Science, College of Medicine, Swansea University, UK

**Professor Jim Thornton** Professor of Obstetrics and Gynaecology, Faculty of Medicine and Health Sciences, University of Nottingham, UK

**Professor Martin Underwood** Director, Warwick Clinical Trials Unit, Warwick Medical School, University of Warwick, UK

Please visit the website for a list of editors: www.journalslibrary.nihr.ac.uk/about/editors

Editorial contact: journals.library@nihr.ac.uk

## **Abstract**

Improving the Quality of Dentistry (IQuaD): a cluster factorial randomised controlled trial comparing the effectiveness and cost-benefit of oral hygiene advice and/or periodontal instrumentation with routine care for the prevention and management of periodontal disease in dentate adults attending dental primary care

Craig R Ramsay, 1\* Jan E Clarkson, 2 Anne Duncan, 1 Thomas J Lamont, 3 Peter A Heasman, 4 Dwayne Boyers, 5 Beatriz Goulão, 1 Debbie Bonetti, 2 Rebecca Bruce, 1 Jill Gouick, 2 Lynne Heasman, 4 Laura A Lovelock-Hempleman, 2 Lorna E Macpherson, 2 Giles I McCracken, 4 Alison M McDonald, 1 Fiona McLaren-Neil, 2 Fiona E Mitchell, 2 John DT Norrie, 1 Marjon van der Pol, 5 Kirsty Sim, 2 James G Steele, 4 Alex Sharp, 4 Graeme Watt, 2 Helen V Worthington 6 and Linda Young 7

<sup>1</sup>Health Services Research Unit, University of Aberdeen, Aberdeen, UK
<sup>2</sup>Dental Health Services Research Unit, University of Dundee, Dundee, UK
<sup>3</sup>Dundee Dental School, University of Dundee, Dundee, UK
<sup>4</sup>The Dental School, Newcastle University, Newcastle upon Tyne, UK
<sup>5</sup>Health Economics Research Unit, University of Aberdeen, Aberdeen, UK
<sup>6</sup>The School of Dentistry, University of Manchester, Manchester, UK
<sup>7</sup>NHS Education for Scotland, Edinburgh, UK

**Background:** Periodontal disease is preventable but remains the most common oral disease worldwide, with major health and economic implications. Stakeholders lack reliable evidence of the relative clinical effectiveness and cost-effectiveness of different types of oral hygiene advice (OHA) and the optimal frequency of periodontal instrumentation (PI).

**Objectives:** To test clinical effectiveness and assess the economic value of the following strategies: personalised OHA versus routine OHA, 12-monthly PI (scale and polish) compared with 6-monthly PI, and no PI compared with 6-monthly PI.

**Design:** Multicentre, pragmatic split-plot, randomised open trial with a cluster factorial design and blinded outcome evaluation with 3 years' follow-up and a within-trial cost–benefit analysis. NHS and participant costs were combined with benefits [willingness to pay (WTP)] estimated from a discrete choice experiment (DCE).

**Setting:** UK dental practices.

**Participants:** Adult dentate NHS patients, regular attenders, with Basic Periodontal Examination (BPE) scores of 0, 1, 2 or 3.

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

<sup>\*</sup>Corresponding author c.r.ramsay@abdn.ac.uk

**Intervention:** Practices were randomised to provide routine or personalised OHA. Within each practice, participants were randomised to the following groups: no PI, 12-monthly PI or 6-monthly PI (current practice).

Main outcome measures: Clinical – gingival inflammation/bleeding on probing at the gingival margin (3 years). Patient – oral hygiene self-efficacy (3 years). Economic – net benefits (mean WTP minus mean costs).

**Results:** A total of 63 dental practices and 1877 participants were recruited. The mean number of teeth and percentage of bleeding sites was 24 and 33%, respectively. Two-thirds of participants had BPE scores of ≤ 2. Under intention-to-treat analysis, there was no evidence of a difference in gingival inflammation/ bleeding between the 6-monthly PI group and the no-PI group [difference 0.87%, 95% confidence interval (CI) −1.6% to 3.3%; p = 0.481] or between the 6-monthly PI group and the 12-monthly PI group (difference 0.11%, 95% CI −2.3% to 2.5%; p = 0.929). There was also no evidence of a difference between personalised and routine OHA (difference −2.5%, 95% CI −8.3% to 3.3%; p = 0.393). There was no evidence of a difference in self-efficacy between the 6-monthly PI group and the no-PI group (difference −0.028, 95% CI −0.119 to 0.063; p = 0.543) and no evidence of a clinically important difference between the 6-monthly PI group and the 12-monthly PI group (difference −0.097, 95% CI −0.188 to −0.006; p = 0.037). Compared with standard care, no PI with personalised OHA had the greatest cost savings: NHS perspective −£15 (95% CI −£34 to £4) and participant perspective −£64 (95% CI −£112 to −£16). The DCE shows that the general population value these services greatly. Personalised OHA with 6-monthly PI had the greatest incremental net benefit [£48 (95% CI £22 to £74)]. Sensitivity analyses did not change conclusions.

**Limitations:** Being a pragmatic trial, we did not deny PIs to the no-PI group; there was clear separation in the mean number of PIs between groups.

**Conclusions:** There was no additional benefit from scheduling 6-monthly or 12-monthly PIs over not providing this treatment unless desired or recommended, and no difference between OHA delivery for gingival inflammation/bleeding and patient-centred outcomes. However, participants valued, and were willing to pay for, both interventions, with greater financial value placed on PI than on OHA.

**Future work:** Assess the clinical effectiveness and cost-effectiveness of providing multifaceted periodontal care packages in primary dental care for those with periodontitis.

Trial registration: Current Controlled Trials ISRCTN56465715.

**Funding:** This project was funded by the National Institute for Health Research (NIHR) Health Technology Assessment programme and will be published in full in *Health Technology Assessment*; Vol. 22, No. 38. See the NIHR Journals Library website for further project information.

## **Contents**

List of tables	xiii
List of figures	xvii
List of abbreviations	xix
Plain English summary	xxi
Scientific summary	xxiii
Chapter 1 Introduction	1
The reason for the trial	1
Background	1
The questions the IQuaD trial addressed	3
Aim	3
Objectives	3
Chapter 2 Methods of the study	5
Study design	5
Ethics approval and consent	5
Participants and procedure	5
Recruitment and consent of dental practices	5
Recruitment and consent of participants	6
Inclusion/exclusion criteria	6
Trial outcome assessor training	7
Trial interventions	7
Oral hygiene advice	8
Periodontal instrumentation	9
Randomisation	9
Practice allocation to oral hygiene advice group	9
Participant allocation to periodontal instrumentation group	9
Descriptive measures	10
Outcome measures	12
Number of periodontal instrumentations received (routine data and self-report)	14
Data collection	14
Baseline	15
Annual follow-up	15
Three-year follow-up	15
Sample size	16
Oral hygiene advice	16
Periodontal instrumentation	16
Interaction	16
Collection of providers outcomes	16
Statistical analyses of outcomes	17
Non-response analysis	17
Missing items	17
Sensitivity analysis	17

Trial oversight	17
Patient and public involvement	18
Protocol amendments after trial initiation	18
Chapter 3 Health economic methods	19
Resource use and costs	19
Health services costs	19
Participant costs	22
Statistical analysis of cost data	22
Benefits (willingness to pay)	24
Selection of attributes and levels	24
Experimental design	24
Questionnaire development	27
Data collection	27
Data analysis	27
Estimating benefits (willingness to pay) for each trial participant	28
Cost–benefit analysis	28
Statistical analysis of (incremental) net benefit	29
Presentation of cost–benefit analysis results	29
Analysis of uncertainty	29
Cost–consequences analysis	29
Chapter 4 Trial results and clinical effectiveness	31
Recruitment to the study	31
Dental practices	31
Participants	31
Description of the groups at trial entry	31
Practice characteristics	31
Participant characteristics	34
Clinical characteristics	34
Patient-reported outcomes	34
Cognition variables	34
Behaviour	34
Intention	36
Quality of life	36
Trial follow-up	36
Attendance at 3-year clinical examination	36
Annual questionnaire returns at years 1, 2 and 3	36
Participant dental characteristics at 3 years	36
Statistical analyses	38
Primary outcomes	38
Secondary outcomes	42
Tertiary outcomes	52
Fidelity of interventions	52
Subgroup analyses	53 53
Sensitivity analyses	55
Chapter 5 Results of economic analysis	57
Resource use and costs	57 57
NHS dental care resource use	57
NHS dental care costs	57
Other NHS-incurred costs	62
Participant-incurred costs	64
raracipant incanca costs	04

Benefits	68
Discrete choice experiment	68
Results of analysis models	70
Validity of responses	73
Analysis of subgroups	73
Benefits: willingness to pay	73
Cost-benefit analysis	78
Cost–consequences analysis	81
Chapter 6 Cohort	85
Introduction	85
Methods	85
Ethics approval	85
Recruitment and consent	85
Outcomes	85
Statistical analyses	85
Results	86
Baseline characteristics	86
Clinical outcomes	86
Patient-centred outcomes	86
Cognitions	86
Behaviour	86
Quality of life	87
Follow-up characteristics	87
Clinical	87
Patient-centred outcomes	87
Discussion	87
Chapter 7 Discussion/conclusions	89
Comparisons with other randomised clinical trials	91
Strengths	92
Limitations	95
Patient and public involvement and engagement	97
Generalisability	97
Recommendations for research	97
Acknowledgements	99
References	103
Appendix 1 Clinical effectiveness outcomes	109
Appendix 1 clinical effectiveness outcomes	109
Appendix 2 Health economics	119
Appendix 3 Recruitment	141

## **List of tables**

TABLE 1 Participant dental characteristics at baseline, by randomised group	10
TABLE 2 Summary of UK dental charges, by region	21
TABLE 3 Unit costs for the opportunity costs of time and travel	23
TABLE 4 Attributes and levels included in the DCE	25
TABLE 5 Deterministic sensitivity analyses considered	30
TABLE 6 Screened/declined	33
TABLE 7 Practice characteristics/beliefs	33
TABLE 8 Clinical characteristics at baseline, by randomised group	35
TABLE 9 Patient-reported outcomes at baseline, by randomised group	35
<b>TABLE 10</b> Comparison of baseline characteristics of 3-year follow-up assessment of attenders and non-attenders	37
TABLE 11 Patient and dental characteristics at 3 years, by randomised group	37
TABLE 12 Treatment effects for primary outcomes	40
TABLE 13 Treatment effects for secondary outcomes	42
TABLE 14 Clinician belief questionnaire at follow-up	51
TABLE 15 Number of PIs received	52
<b>TABLE 16</b> Interaction mean for 6-monthly PI vs. other PI and personalised OHA vs. routine OHA, by subgroup	55
TABLE 17 Number of treatment claims	58
TABLE 18 Number of participants having PI in the routine data set	59
TABLE 19 NHS costs (£) of dental care	60
TABLE 20 Exemptions from NHS dental charges	63
TABLE 21 Validation of the rates of other NHS secondary care resource use	64
TABLE 22 Participant-incurred co-charges (£) for dental care	65
TABLE 23 Participant-incurred costs (£)	66
TABLE 24 Characteristics of DCE participants	69

TABLE 25 Participant dental care experience	71
TABLE 26 Results from general population DCE: mixed logit model	72
TABLE 27 Analysis of DCE subgroups	74
TABLE 28 Benefits	76
TABLE 29 Cost-benefit analysis results: base case – NHS dental perspective costs (£)	79
TABLE 30 Cost-benefit analysis results: NHS and participant perspective costs (£)	80
TABLE 31 Main sensitivity analysis results	81
TABLE 32 Cost–consequences analysis: narrative summary	82
TABLE 33 Calculations for outcomes in the CBQ at baseline	109
TABLE 34 Calculations for outcomes in the CBQ at follow-up	109
TABLE 35 Calculations for patient-reported outcomes	110
TABLE 36 Participant dental characteristics at year 1	111
TABLE 37 Participant dental characteristics at year 2	112
TABLE 38 Attendance rates by randomised allocation	114
TABLE 39 Clinical outcomes for participants who attended clinical follow-up only	114
TABLE 40 Questionnaire response rates by randomised group	114
TABLE 41 Patient-reported outcomes of those who replied to the follow-up questionnaire at 3 years	115
TABLE 42 Tertiary outcomes at baseline	116
TABLE 43 Tertiary outcomes at 3 years	117
TABLE 44 Cost of consumer dental care products	119
TABLE 45 Experimental design, by segmented group	120
TABLE 46 Mapping DCE WTP values to trial outcomes	121
TABLE 47 Details of dental claims (Scotland)	124
TABLE 48 Descriptive statistics for participant-reported contact with non-dental health services	125
TABLE 49 Calculation of unit opportunity cost of time and travel to dental           appointments	126

TABLE 50 Sensitivity analysis: including baseline and final clinical visits	132
TABLE 51 Sensitivity analysis: discounting 0% for costs and benefits	133
TABLE 52 Sensitivity analysis: discounting 6% for costs and benefits	134
TABLE 53 Sensitivity analysis: all PI and OHA delivered by a dentist	135
TABLE 54 Sensitivity analysis: all PI and OHA delivered by a hygienist	136
TABLE 55 Sensitivity analysis: adding an additional cost for personalised OHA to           all respondents in this cluster	137
TABLE 56 Sensitivity analysis: decreasing value of UDA (-20%)	138
TABLE 57 Sensitivity analysis: increasing value of UDA (+ 20%)	139
TABLE 58 Alternative method to map from DCE to trial for aesthetic outcome           (mapping to 'after brushing')	140
TABLE 59 Recruitment by centre	142

# **List of figures**

FIGURE 1 Study design	5
FIGURE 2 The OHA intervention behavioural framework	8
FIGURE 3 Example choice task	26
FIGURE 4 The Consolidated Standards of Reporting Trials (CONSORT) diagram	32
FIGURE 5 Gingival inflammation/bleeding (mean and 95% CI), by randomised allocation	39
FIGURE 6 Self-efficacy score (mean and 95% CI), by randomised allocation	41
FIGURE 7 Calculus percentage (mean and 95% CI), by randomised allocation	43
FIGURE 8 Mean clinical probing depth (mean and 95% CI), by randomised allocation	44
FIGURE 9 Perceived behaviour control score (mean and 95% CI), by randomised allocation	45
FIGURE 10 Attitude score (mean and 95% CI), by randomised allocation	46
FIGURE 11 Subjective norm score (mean and 95% CI), by randomised allocation	47
FIGURE 12 Behaviour score (mean and 95% CI), by randomised allocation	48
FIGURE 13 Intention score (mean and 95% CI), by randomised allocation	49
FIGURE 14 The OHIP-14 score (mean and 95% CI), by randomised allocation	50
FIGURE 15 Subgroup results for PI allocation: difference between arms, by subgroup	54
FIGURE 16 Subgroup results for OHA allocation: difference between arms, by subgroup	54
FIGURE 17 Routine PI for periodontal health in adults	93
FIGURE 18 One-to-one OHA provided in a dental setting for oral health	94
FIGURE 19 Confidence ellipses for base-case health economic results	130
FIGURE 20 Target vs. actual practice accrual	141
FIGURE 21 Target vs. actual participant accrual	141

# **List of abbreviations**

*	furcation involvement	ITT	intention to treat
ADHS	Adult Dental Health Survey	NICE	National Institute for Health and
ASC	alternative-specific constant		Care Excellence
BPE	Basic Periodontal Examination	OA	outcome assessor
BSA	Business Services Authority	ОНА	oral hygiene advice
СВА	cost–benefit analysis	OHIP-14	Oral Health Impact Profile-14
CBQ	clinician belief questionnaire	PBC	perceived behaviour control
CCA	cost–consequences analysis	PI	periodontal instrumentation
CHaRT	Centre for Healthcare Randomised Trials	PMG	Project Management Committee
		PPI	patient and public involvement
CI	confidence interval	QoL	quality of life
DCE	discrete choice experiment	RCT	randomised controlled trial
DMC	Data Monitoring Committee	REC	Research Ethics Committee
FG	focus group	SD	standard deviation
GDS	General Dental Service	TCOD	Trial Coordinating Office in Dundee
HTA	Health Technology Assessment	TMC	Trial Management Committee
ICC	intracluster correlation coefficient	TSC	Trial Steering Committee
IQuaD	Improving the Quality of Dentistry	UDA	Unit of Dental Activity
ISD	Information Services Division	UNC	University of North Carolina
IT	information technology	WTP	willingness to pay

# **Plain English summary**

A lmost half of adults in the UK show signs of gum disease. It is a largely preventable disease, with treatment usually taking the form of self-care, including tooth brushing and dentist cleaning, which is often called 'scale and polish'. In this study, we wanted to find out whether or not the advice patients are given and the cleaning they receive from the dentist work and offer good value for money.

A total of 1877 adults who regularly attended 63 different NHS dental practices across Scotland and north-east England were involved. The adults had early signs of gum disease and were randomly chosen to be offered a scale and polish every 6 months, every 12 months or not at all. They received normal advice or personalised advice depending on what dental practice they attended. The recruited adults completed study surveys every year. They also had clinical measurements of their mouth taken by study staff at the start of the study and 3 years later.

After 3 years, there was no evidence of differences in gum health, whether a scale and polish was carried out every 6 or 12 months or only when either the patient requested it or the dentist recommended it for clinical reasons. There was also no added benefit of providing personalised oral health advice. However, patients value, and are willing to pay for, cleaning advice and scale and polish, but are willing to pay more for scale and polish than for advice.

# **Scientific summary**

### **Background**

Periodontal disease is an inflammatory disease that affects the soft and hard tissues supporting teeth. This disease is largely preventable, yet it remains the major cause of poor oral health worldwide and is the primary cause of tooth loss in older adults. Severe periodontitis is the sixth most prevalent human disease, with a standardised prevalence of 11.2%.

The categorisation of periodontal disease is based on which of the tissues surrounding and supporting the teeth are affected and is classified into two broad categories: (1) gingivitis and (2) periodontitis. Gingivitis is a reversible condition characterised by inflammation and bleeding at the gingival margin. The gum becomes swollen and red because of the inflammation and will bleed easily on probing. It is a prerequisite for periodontitis and a risk indicator for caries progression. Periodontitis is the irreversible destruction and loss of the supporting periodontal structures (periodontal ligament, cementum and alveolar bone). The result can be unsightly gingival recession, sensitivity of the exposed root surface, root caries (decay), mobility and drifting of teeth and, ultimately, tooth loss.

Individuals and dental care professionals have different roles to play in the prevention and management of periodontal disease. Effective individual self-care (tooth brushing and interdental aids) for plaque control is considered the foundation stone of successful periodontal prevention and therapy of disease. The current annual public spend on oral care products in the UK alone is £950M. Dental care professionals' role in prevention and periodontal treatment involves providing patients with oral hygiene advice (OHA) (self-care) and periodontal instrumentation (PI), often known as 'scale and polish'. There is no agreed published content of OHA but the overall aim is to encourage effective self-care. PI comprises removal of plaque and plaque retentive factors [e.g. calculus (tartar) deposits] which, together with the removal of overhanging restorations (poorly adapted dental fillings), facilitates adequate patient-performed oral self-care. In the UK, almost all of this treatment is provided by general dental practitioners and dental hygienists/therapists in primary care.

Despite evidence of an association between sustained, good oral hygiene and a low incidence of periodontal disease and caries in adults, there is a lack of strong and reliable evidence to inform clinicians of the relative clinical effectiveness (if any) of different types of OHA that can be delivered in a dental setting.

The evidence to inform clinicians of the effectiveness and optimal frequency of PI is mixed. A Cochrane systematic review (Worthington HV, Clarkson JE, Bryan G, Beirne PV. Routine scale and polish for periodontal health in adults. *Cochrane Database Syst Rev* 2013;**11**:CD004625) of routine PI for periodontal health in adults found insufficient evidence to determine the effects of routine PI treatments, providing little guidance for policy-makers, dental professionals or patients.

There was therefore an urgent need to assess the relative clinical effectiveness of OHA and PI in a robust, sufficiently powered randomised controlled trial (RCT) in primary dental care.

The trial protocol was published in *BMC Oral Health* (Clarkson JE, Ramsay CR, Averley P, Bonetti D, Boyers D, Campbell L, *et al.* IQuaD dental trial; improving the quality of dentistry: a multicentre randomised controlled trial comparing oral hygiene advice and periodontal instrumentation for the prevention and management of periodontal disease in dentate adults attending dental primary care. *BMC Oral Health* 2013;**13**:58).

## **Objectives**

The aim of this study was to compare the clinical effectiveness and cost-effectiveness of theory-based, personalised OHA or PI at different time intervals (no PI, 12-monthly PI or 6-monthly PI), or their combination (OHA and 6-monthly PI), with routine care for improving periodontal health in dentate adults attending general dental practice.

The primary objectives were to test the clinical effectiveness and cost-effectiveness of the following dental management strategies:

- 1. personalised OHA versus routine OHA
- 2. 6-monthly PI versus 12-monthly PI
- 3. 6-monthly PI versus no PI.

The secondary objectives were to:

- 1. test the clinical effectiveness and cost-effectiveness of a combination of personalised OHA with different time intervals for PI
- 2. measure dentist/hygienist beliefs relating to giving OHA, PI and maintenance of periodontal health.

#### **Methods**

#### Design

Improving the Quality of Dentistry (IQuaD) was a 5-year multicentre, pragmatic split-plot, cluster randomised, open trial with blinded outcome evaluation. The comparisons were made within a factorial design using a combination of cluster and individual participant randomisation. As personalised OHA was given by the dentist or hygienist, there was a theoretical risk of 'contamination' between patient participants seen within the same dental practice. To minimise this potential risk, dental practices were randomised to deliver routine or personalised OHA. All patient participants seen by the same dental practice (a 'cluster') received either routine (current practice) or personalised OHA, depending on their dental practice allocation. To test the effects of PI, each individual patient participant was randomised to no PI, 12-monthly PI or 6-monthly PI (current practice).

#### Setting

The trial recruited dental practitioners from general dental practices in Scotland and north-east England (Newcastle upon Tyne). Participating dentists represented a cross-section of practitioners operating in a range of different circumstances (e.g. urban or rural, high- to low-income communities, employing a dental hygienist or not).

#### **Dentist participants**

### Inclusion criteria

- NHS provider for adult patients.
- Primary care provider.
- Willing to follow protocol.

#### **Exclusion criteria**

- Providing only private dental care.
- Unwilling to follow protocol.

### Patient participants

#### Inclusion criteria

Adult patients (aged  $\geq$  18 years) with gingivitis or moderate periodontitis [a Basic Periodontal Examination (BPE) score of 0, 1, 2 or 3] who:

- were dentate
- had attended for a check-up at least twice in the previous 2 years
- received their dental care in part or in full as a NHS patient.

#### **Exclusion** criteria

- Patients with a BPE score of 4 (clinical probing depth of > 6 mm and/or furcation involvements or attachment loss of ≥ 7 mm) in any sextant on the basis that more extensive periodontal care was indicated.
- Patients with an uncontrolled chronic medical condition (e.g. diabetes mellitus, immunocompromised).

#### **Interventions**

Routine OHA was defined as the OHA currently being provided by the practices. There is no published information describing 'routine' OHA, but anecdotal evidence suggests that this is often the provision of minimal OHA (e.g. 'you need to brush your teeth more frequently') or no OHA.

The personalised OHA intervention was based on social cognitive theory and implementation intention theory. The content of the OHA delivered was personalised according to the dentist's/hygienist's assessment of the needs of the patient. At a minimum, the content included advice and instruction in self-diagnosis (e.g. bleeding gums on brushing indicates the presence of reversible gingival inflammation) and advice and instruction on tooth brushing and flossing (frequency and technique). On completion of the OHA, the dentist agreed an action plan with the patient.

The definition of PI was as used in standard practice and could include the removal of plaque and calculus from the crown and root surfaces using manual or ultrasonic scalers, with no adjunctive subgingival therapy (e.g. local delivery antibiotics), and the appropriate management of plaque retention factors.

Experimental groups received a PI at 6- or 12-monthly intervals according to the individual participant-level randomisation. Participants allocated to the no-PI groups attended their dentist at time intervals determined by current practice. However, participants and dental practices were advised that every trial participant should be invited to attend for a routine examination appointment at least every 12 months.

#### Main outcome measures

#### **Primary outcomes**

- Clinical: gingival inflammation/bleeding on probing at the gingival margin at the 3-year follow-up.
- Patient centred: oral hygiene self-efficacy at the 3-year follow-up.
- Economic: net benefits [mean willingness to pay (WTP) minus mean costs].

### Secondary outcomes

- Clinical: (1) calculus, (2) clinical probing depth, (3) additional PI and (4) referral. (All of which were collected at the 3-year follow-up.)
- Patient centred: (1) dental quality of life, (2) oral health behaviour and (3) knowledge. (All of which were collected during 3 years' annual follow-up.)
- Economic: costs to the NHS and patients; WTP.
- Provider: beliefs relating to giving OHA and maintenance of periodontal health.

Note: the Periodontal Advisory Group considered that Clinical Attachment Loss and plaque cannot be measured reliably and so neither was included as outcomes.

Clinical outcomes were measured at baseline by trained outcome assessors (OAs) who were blinded to allocation. Gingival inflammation/bleeding scores, calculus, clinical probing depth and BPE scores were measured by the OAs and recorded on the baseline clinical chart by the dental research nurse, who was a member of the trial team. Patient-centred outcomes were measured at baseline and annually by self-administered postal questionnaire.

### **Economic evaluation**

A within-trial cost—benefit analysis assessed the costs and benefits (in monetary terms) of each policy compared with standard care (routine OHA with 6-monthly PI). NHS and wider (NHS and participant) perspectives were considered.

Routinely collected dental claims data were linked to trial data to determine the costs of NHS-provided care (including participant co-charges). Additional participant costs, including private care, self-purchased products, and time and travel costs, were sourced from participant annual questionnaires.

A discrete choice experiment (DCE), administered to a nationally representative online sample of the UK general population, was used to estimate WTP. The design was pivoted and segmented to improve realism. DCE data were analysed using mixed logit regression models. WTP tariffs from the DCE were mapped to treatments received (PI and OHA), self-reported bleeding and aesthetics outcomes to calculate benefits.

The discount rate was 3.5%. Multilevel hierarchical models accounted for clustering, correlation between benefits and costs, and minimisation covariates. Results were presented as incremental net benefits, using confidence ellipses to illustrate uncertainty. Deterministic sensitivity analyses tested the impact of key assumptions on results. The fully approved protocol for the IQuaD trial can be accessed online via www.journalslibrary.nihr.ac.uk/programmes/hta/090145/#/ (accessed October 2017).

### **Results**

A total of 2341 patients were screened for trial entry and 1877 participants were recruited. Of the screened patients, a total of 183 (8%) were found to be ineligible. The primary reason for ineligibility was a BPE score of 4 or \* (furcation involvement), affecting 160 patients. From those ineligible because of a BPE score of 4 or \*, 144 (90%) patients agreed to join a separate cohort group. There were 281 patients potentially eligible for the study who were not recruited.

At baseline, the mean number of teeth per participant was 24. Two-thirds of participants had a BPE score of  $\leq$  2. The mean proportion of sites affected by bleeding was 33% and 35% of teeth had calculus present. The mean clinical probing depth was 1.8 mm. Between 10% and 12% of participants in each

group had four or more pockets with a clinical probing depth of  $\geq 4$  mm. There were no important imbalances across the randomised groups.

The pre-chosen clinical primary outcome was mean gingival inflammation/bleeding at 3 years' follow-up; 71% of the participants attended the 3-year clinical follow-up appointment. Under intention-to-treat (ITT) analysis, there was no evidence of a difference between those randomised to receive 6-monthly PI and those randomised to receive no PI [difference 0.87%, 95% confidence interval (CI) -1.6% to 3.3%; p = 0.481] (i.e. there was a < 1% difference in the average number of sites with gingival bleeding between the randomised groups). Similarly, there was no evidence of a difference between 6-monthly PI and 12-monthly PI (difference 0.11%, 95% CI -2.3% to 2.5%; p = 0.929). The 95% CIs were small enough to exclude the prespecified clinically important difference of 7.5% in bleeding. There was also no evidence of a difference between participants randomised to personalised OHA and those randomised to routine OHA (difference -2.5%, 95% CI -8.3% to 3.3%; p = 0.393). The results were robust to other adjusted/unadjusted models. The interaction between personalised OHA and 6-monthly PI was 1.7 (95% CI -3.8 to 7.3) (i.e. neither statistically nor clinically significant).

The pre-chosen patient-centred primary outcome was self-efficacy at the 3-year follow-up; 77% of the participants completed a 3-year questionnaire. Under ITT analysis, there was no evidence of a difference between those randomised to receive 6-monthly PI and those randomised to receive no PI (difference -0.028, 95% CI -0.119 to 0.063; p = 0.543). Between those randomised to receive 6-monthly and 12-monthly PI, there was a statistically significant difference at the 5% level favouring the 6-monthly PI (difference -0.097, 95% CI -0.188 to -0.006; p = 0.037); however, the size of the difference was not clinically important.

There were no significant differences in NHS dental costs. No PI with personalised OHA was the least costly policy (£62.42) [mean difference vs. standard care -£14.91 (95% CI -£15.70 to £28.16)], followed by no PI and routine OHA (£72.23) [mean difference -£3.12 (95% CI -£18.18 to £11.93)]. From a wider perspective, no PI with personalised OHA was significantly less costly [mean difference -£64.11 (95% CI -£112.33 to -£15.88)], followed by no PI with routine OHA [mean difference -£39.80 (95% CI -£83.94 to £4.33)].

The DCE showed that the general population valued both PI and personalised OHA even when controlling for bleeding and aesthetic outcomes. Therefore, 6-monthly PI with personalised OHA has the greatest benefit [mean difference vs. standard care £61.67 (95% CI £40.19 to £83.14)]. A 12-monthly PI with personalised OHA also had positive, but not significant, incremental benefits [mean difference £19.70 (95% CI –£1.64 to £41.04)]. This suggests that, in terms of WTP, a reduction in PI frequency can, in part, be compensated for by introducing personalised OHA.

Six-monthly PI with personalised OHA had the largest incremental net benefit compared with standard care from a NHS perspective [mean difference £48 (95% CI £22 to £74)] and a wider perspective [mean difference £68 (95% CI £15 to £120)]. The overall health economic results were broadly consistent across the UK countries and findings were robust to the sensitivity analyses undertaken.

#### Limitations

Being a pragmatic trial, we did not deny Pls to the no-Pl group, although we did not collect detailed information about the reasons for additional Pls. However, there was clear separation in the mean number of Pls between groups. The economic evaluation was based on current NHS contracts that may change over time. A lifetime decision model was not conducted; however, given the lack of difference in clinical outcomes, extrapolation of trial results would be unlikely to change conclusions.

#### **Conclusions**

The IQuaD trial, involving regular adult NHS dental attenders (with no or early signs of periodontitis), has shown that, over a 3-year period, there is no additional benefit from scheduling 6-monthly or 12-monthly Pls over not providing this treatment unless desired or recommended, and that there is no difference between personalised or routine OHA (current practice) for the trial's primary clinical (gingival inflammation/bleeding) and patient-centred (self-reported) outcomes. However, patients value, and are willing to pay for, both interventions, with greater financial value placed on PI than on OHA.

#### **Recommendations for research**

- Research is needed to assess the clinical effectiveness and cost-effectiveness of providing multifaceted periodontal care packages (e.g. OHA, oral care-products, PI) in primary dental care for those with periodontitis.
- Research is required to better understand the source of WTP values and the extent to which this is influenced by perceptions and current practice.
- Research is needed to explore the relative value of different data sources for estimating resource use in dentistry including routine data, patient-reported data and practice records.

## **Trial registration**

This trial is registered as ISRCTN56465715.

## **Funding**

Funding for this study was provided by the Health Technology Assessment programme of the National Institute for Health Research.

# **Chapter 1** Introduction

The subsequent chapters of this monograph describe Improving the Quality of Dentistry (IQuaD), a National Institute for Health Research (NIHR) Health Technology Assessment (HTA) programme-funded trial testing the relative effectiveness of different types of oral hygiene advice (OHA) and the optimal frequency of periodontal instrumentation (PI). The trial protocol has been published.<sup>1</sup>

### The reason for the trial

### **Background**

#### **Epidemiology**

Periodontal disease is an inflammatory disease that affects the soft and hard tissues supporting teeth. This disease is largely preventable, yet it remains the major cause of poor oral health worldwide and is the primary cause of tooth loss in older adults.<sup>2,3</sup> Severe periodontitis is the sixth most prevalent human disease, with a standardised prevalence of 11.2%.<sup>4</sup>

The categorisation of periodontal disease is based on which of the tissues surrounding and supporting the teeth are affected and is classified into two broad categories: (1) gingivitis and (2) periodontitis. Gingivitis is a reversible condition characterised by inflammation and bleeding at the gingival margin. The gum becomes swollen and red because of the inflammation and will bleed easily on probing. It is a prerequisite for periodontitis and is also a risk indicator for caries progression. Periodontitis is the irreversible destruction and loss of the supporting periodontal structures (periodontal ligament, cementum and alveolar bone).<sup>5</sup> The result can be unsightly gingival recession, sensitivity of the exposed root surface, root caries (decay), mobility and drifting of teeth and, ultimately, tooth loss.

Gingivitis and periodontitis are a continuum of the same inflammatory disease<sup>6</sup> but it is currently not possible to predict if and/or when an individual will progress from the reversible gingivitis to irreversible periodontitis. Accumulation of microbial dental plaque is the primary aetiological factor for gingivitis and periodontitis, as well as dental caries. However, progression of the disease is known to be affected by other factors including genetics (host's defence mechanisms to bacterial infection), calculus, smoking and systemic comorbidities including type 2 diabetes mellitus.<sup>7–10</sup> Although several risk factors have been identified, the lack of certainty as to whether or not, and when, they may cause progression to irreversible periodontitis in an individual causes a challenge for the profession.

The 2009 UK Adult Dental Health Survey (ADHS)<sup>11</sup> provides evidence that the majority of UK adults might be at risk of developing periodontal disease: 66% of dentate adults had visible plaque, indicating that tooth brushing was ineffective, and 68% had calculus in at least one sextant of the mouth. Gingival bleeding was demonstrated in 54% of dentate adults and 45% of dentate adults had periodontitis (defined by at least one site with a clinical probing depth of  $\geq 4$  mm), increasing with age from 19% in those aged 16-24 years to 61% in those aged 75-84 years. Indicators of severe disease (at least one site with a clinical probing depth of  $\geq 6$  mm) also increased with age, affecting 14% of those aged  $\geq 65$  years. Only 17% of dentate adults had excellent periodontal health, which was defined by the ADHS  $2009^{11}$  as 'no bleeding, no calculus, no periodontal probing depths of 4 mm or more and in the case of adults aged 55 or above, no loss of periodontal attachment over 4 mm or more anywhere in their mouth'.

As microbial dental plaque accumulation is considered the most important risk factor for periodontitis, its disruption or removal is a key component of prevention, combined with the control/management of other risk factors (e.g. smoking).<sup>12,13</sup>

Prevention strategies aim to prevent the inflammatory process from destroying the periodontal attachment as well as prevent the recurrence of inflammation in successfully treated patients.<sup>14</sup>

Individuals and dental care professionals have different roles to play in prevention. Effective individual self-care (tooth brushing and interdental aids) for plaque control is considered the foundation stone of successful periodontal prevention and therapy of disease.<sup>13</sup> The current annual public spend on oral care products in the UK alone is £950M.<sup>15</sup> The dental care professionals' role in prevention and periodontal treatment involves providing patients with OHA (self-care) and Pl. There is no agreed published content of OHA, but the overall aim of this intervention is to encourage effective self-care. Pl (or 'scale and polish') comprises removal of plaque and plaque retentive factors [e.g. calculus (tartar) deposits] which, together with the removal of overhanging restorations (poorly adapted dental fillings), facilitate adequate patient-performed oral self-care. In the UK, almost all of this treatment is provided by general dental practitioners and dental hygienists/therapists in the primary care setting. The British Society of Periodontology<sup>16</sup> advises that consideration be given to referral to a specialist periodontist of patients exhibiting severe or aggressive forms of periodontal diseases.

Periodontal instrumentation is one of the most frequently provided treatments in general dental practices. In the year 2014/15, there were 2.2 million claims for this simple periodontal treatment in Scotland, costing £31.5M.<sup>17</sup> This represents 8% of the total NHS budget spend of general and public dental services and 25% of the monetary value of all fee-per-item treatments provided in these settings. There were also 2910 claims for intensive scaling, at a cost of £200,000. During this time frame in England, it was estimated that 12.9 million courses of treatment involved PI for adults and approximately 0.9 million courses for children.<sup>18</sup>

Dental reimbursement in Scotland is a retrospective 'fee-per-item' service for which dentists are primarily reimbursed for the number of individual treatments provided following completion of a course of treatment. In 2006, England's dental health service moved from a similar model to a prospective 'Unit of Dental Activity' (UDA) system. Dentists agree contracts in advance with health boards to provide a prespecified number of UDAs. In both health-care systems, non-exempt patients face a patient copayment. In Scotland, the copayment equals 80% of the fee-per-item payment up to a prespecified upper limit. In England, there is a fixed copayment according to the band of treatment provided.

#### Evidence base

Despite evidence of an association between sustained good oral hygiene and a low incidence of periodontal disease and caries in adults, <sup>19</sup> there is a lack of strong and reliable evidence to inform clinicians of the relative clinical effectiveness (if any) of different types of OHA that can be delivered in a dental setting.

Prior to the start of the IQuaD trial, a number of relevant systematic reviews<sup>20–22</sup> evaluating OHA had been conducted, with some inconsistency in their findings. A recent systematic review<sup>23</sup> of psychological approaches to behaviour change for improved plaque control in periodontal management reported benefits of using goal-setting, self-monitoring and planning for improving oral hygiene. Patient understanding of the benefits of behaviour change and the seriousness of periodontal disease were also considered important by the authors of the review.<sup>23</sup> A number of the included trials were of short duration, had non-experimental designs and were rated as being at high risk of bias. A meta-analysis of the available trials was not possible and the results and recommendations should be interpreted with caution.

The evidence to inform clinicians of the clinical effectiveness and optimal frequency of PI is mixed. A Cochrane systematic review<sup>24</sup> of routine scale and polish (PI) for periodontal health in adults found insufficient evidence to determine the effects of routine PI treatments, providing little guidance for policy-makers, dental professionals or patients. Only three trials<sup>25–27</sup> were eligible for inclusion and all were rated as being at an unclear risk of bias. Given that PI is routinely provided in general dental practice, it is noteworthy that only one of the eligible trials<sup>25</sup> was conducted in primary care. Following baseline PI, Jones *et al.*<sup>25</sup> compared no PI delivery with 6-monthly PI or 12-monthly PI, with a 24-month clinical follow-up. The trial<sup>25</sup> recruited only periodontally healthy participants. The Cochrane systematic review<sup>24</sup> assessed the body of evidence for PI as

being of low quality. The need for further well-conducted trials of sufficient duration, including research investigating patients' willingness to pay (WTP), was highlighted.<sup>28</sup>

The relative effectiveness of OHA and PI was assessed in the IQuaD trial, a robust, adequately powered randomised controlled trial (RCT) in primary dental care.

## The questions the IQuaD trial addressed

#### Aim

The aim of this study was to compare the clinical effectiveness and cost-effectiveness of theory-based, personalised OHA or PI at different time intervals (no PI, 12-monthly PI or 6-monthly PI), or their combination, with routine care in improving periodontal health in dentate adults attending general dental practice.

### **Objectives**

The primary objectives were to test the clinical effectiveness and cost-effectiveness of the following dental management strategies:

- 1. personalised OHA versus routine OHA
- 2. 6-monthly PI versus 12-monthly PI
- 3. 6-monthly PI versus no PI.

The secondary objectives were to:

- 1. test the clinical effectiveness and cost-effectiveness of a combination of personalised OHA and PI at different time intervals
- 2. measure dentist/hygienist beliefs relating to giving OHA, PI and maintenance of periodontal health.

# Chapter 2 Methods of the study

## Study design

This study was a multicentre, cluster randomised controlled, open trial with blinded outcome evaluations and 3-year follow-ups. The comparisons were made within a factorial design using a combination of cluster and individual participant randomisation. Dental practices were randomised to either a routine or personalised OHA group. All participants were seen in the same dental practice by a dentist or hygienist (a 'cluster') and received either routine (current practice) or personalised OHA depending on their dental practice's allocation. This was the optimal design to address the concern that contamination could happen if each dentist delivered routine OHA to some participants and personalised OHA to others. To test the effects of PI, each individual participant was randomised to one of three groups: (1) no PI, (2) 6-monthly PI (current practice) or (3) 12-monthly PI (Figure 1).

## **Ethics approval and consent**

Favourable ethics opinion for the IQuaD trial was confirmed by the East of Scotland Research Ethics Service on 24 March 2011 [Research Ethics Committee (REC) reference number 10/S0501/65].

## **Participants and procedure**

#### Recruitment and consent of dental practices

Recruitment of dental practices was achieved by a variety of methods. The IQuaD trial was presented at a number of national conferences and promoted within dental professional publications to raise awareness of the trial within the dental profession. In addition, the Scottish Dental Practice Based Research Network and the Faculty of General Dental Practitioners provided trial information to their members. A series of trial information and recruitment evenings were organised across Scotland and north-east England. Potential dentist participants were sent personalised invitation letters to these events in which the reasons for the trial, design of the trial and practice involvement were described, and dental professionals were given the opportunity to discuss participation with the trial team. Information packs about the trial were posted to dentists who were unable to attend a meeting.

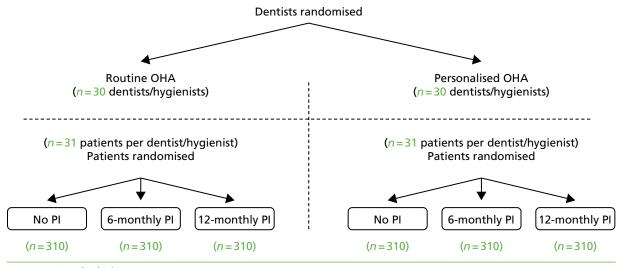


FIGURE 1 Study design.

Trial team members telephoned dental practices to follow up the trial information letters and noted interest of involvement. Principal dentists (usually the owner of the practice) who were interested in the trial were asked to provide written consent for the dental practice (cluster) to participate. All participating dentists and hygienists within a dental practice were individually invited to complete a consent form and the clinician belief questionnaire (CBQ) prior to cluster randomisation. Participating dental practices were asked to identify dates in advance for training in trial processes and at least three dates for the screening and recruitment of potentially eligible participants.

## Recruitment and consent of participants

The identification of potential participants in each dental practice was supported by the Scottish Primary Care Research Network in Scotland and the UK Clinical Research Network in England. A variety of patient appointment management strategies are utilised within dental practices across the UK. Therefore, the IQuaD trial developed a flexible and pragmatic participant recruitment strategy that aimed to adapt to each dental practice's usual appointment management system. Some dental practices arrange routine check-up appointments for their patients up to 6 months or 1 year in advance, while other practices send letters or mobile phone text message reminders to their patients when their routine dental examinations are due, asking them to contact the dental practice to make an appointment. Dental practices were asked to send an invitation letter to attend a trial screening session, along with a patient information sheet and baseline questionnaire to potentially eligible participants who were due to attend for a routine dental examination and who were regular attenders and did not have severe periodontitis [Basic Periodontal Examination (BPE) score of 4]. This was sent, at most, 6 weeks in advance of the patient's routine dental examination appointment. Patients who were not interested in taking part were asked to contact their practice for an alternative appointment.

Trial outcome assessor (OA) teams, consisting of qualified dental hygienists/dental therapists and dental nurses employed by the trial, attended the screening and recruitment sessions in participating dental practices to obtain consent from potentially eligible participants and collect the baseline clinical measurements and questionnaires of consented participants.

At the screening appointment, the dentist was available to discuss the trial with potential participants and answer any questions. The OA was present at this appointment and answered any questions specifically related to the trial. Patients who did not wish to take part were seen by their dentist/hygienist, who provided an examination, OHA and/or PI as normal. Potentially eligible participants provided written informed consent to the trial before the baseline clinical outcomes were measured by the OA to confirm a patient's eligibility for the trial. Reasons for ineligibility or declining to participate were recorded at this session. Participants excluded from the trial solely because of a BPE score of 4 or \* (furcation involvement) were not randomised to a PI allocation and were given the opportunity to consent to follow-up by annual postal questionnaires and clinical follow-up in a separate cohort group (see study documentation, available at www.journalslibrary.nihr.ac.uk/programmes/hta/090145/#/; accessed October 2017). All participants received a baseline PI and OHA from their own dentist or hygienist after the baseline outcome measures and questionnaires were collected by the OA team prior to participants being informed of their randomised PI allocation.

## Inclusion/exclusion criteria

# **Dental practices**

## Inclusion criteria

- NHS provider for adult patients.
- Primary care provider.
- Willing to follow protocol.

### Exclusion criteria

- Providing only private dental care to adults.
- Unwilling to follow protocol.

## **Participants**

### Inclusion criteria

Adult patients (aged  $\geq$  18 years) with periodontal health, gingivitis or moderate periodontitis (BPE score of 0, 1, 2 or 3) who:

- were dentate
- had attended for a check-up at least twice in the previous 2 years
- received their dental care in part or in full as a NHS patient.

### Exclusion criteria

- Patients with periodontal disease with a BPE score of 4 (clinical probing depth of > 6 mm and/or furcation involvements or attachment loss of ≥ 7 mm) in any sextant on the basis that more extensive periodontal care was indicated.
- Patients with an uncontrolled chronic medical condition (e.g. diabetes mellitus, immunocompromised).

### Trial outcome assessor training

Before participant recruitment and 3 years' clinical follow-up, the OA teams were trained in the recording of the trial clinical outcomes. The training was delivered by trial collaborators who have extensive experience of clinical periodontal research (PH, GM). The emphasis of the training was on the consistency of the scoring, both intra- and interassessor, to achieve assessor alignment.

The practical steps of clinical outcome assessment were agreed on in advance, including sequence of outcome assessment; time allocation; sequence around the mouth; isolation as well as the angulation, positioning and pressure of the University of North Carolina (UNC)-15 probe to ensure a standardised approach across the OA teams.

It is widely accepted that methods of assessing gingival inflammation that provoke gingival bleeding do not allow for repeated assessment.<sup>29</sup> The training for the primary outcome of gingival inflammation as bleeding therefore involved OA group discussion of the technique of assessment, scoring definition, as well as demonstrated photograph and clinical examples. This was repeated for the outcome of calculus.

Periodontal probing depths were recorded at six sites on all erupted teeth using a probing force of approximately 25 g on an independent, but similar, cohort of patients to those recruited to the study at the Newcastle upon Tyne and Dundee dental schools. Assessments by an experienced clinical periodontal researcher (LH) were used the reference standard. Intra- and interoutcome assessor scores were recorded and kappa scores of  $\geq 0.60$  achieved, with over 95% of scores within 1 mm.

### **Trial interventions**

Both cluster-level intervention (OHA) and individual participant-level intervention (PI) were delivered by the dental practice dentist or dental hygienist in line with each individual dental practice's usual practice.

# Oral hygiene advice

## Routine oral hygiene advice

Routine OHA was defined as the OHA currently being provided by the dental practices. There is no published information describing 'routine' OHA, but anecdotal evidence suggests that it is often minimal (e.g. 'you need to brush your teeth more frequently') or not provided at all.

## Personalised oral hygiene advice

The personalised OHA intervention used two well-established psychological models for explaining and influencing behaviour (Figure 2). The first was a pre-motivational model, social cognitive theory.<sup>30</sup> This theory proposes that a key variable influencing motivation and behaviour is self-efficacy, assessed as a person's confidence in their ability to perform behaviour. According to this model, the sources of self-efficacy are doing the behaviour (performance: cleaning their teeth), seeing someone else do it (vicarious example/modelling: the dentist demonstrated to patients how to use the oral health-care tools to clean their teeth on a model of the mouth), being encouraged to perform it (verbal persuasion: the dentist helped the patient make oral hygiene a habit with the right plan) and how we feel afterwards (physical state: the dentist discussed biofeedback, highlighting what clean teeth would feel like). Based on this model, an intervention to influence oral hygiene behaviour should target oral hygiene self-efficacy via these sources. The second model was implementation intention theory,<sup>31</sup> a post-motivational model designed to help people put their intentions into practice, bridging the gap between motivation and behaviour. This theory proposes that making an explicit action plan about where and when a behaviour will be performed increases the likelihood of performing it. Action plans work by setting up a cue to remind the individual to perform the behaviour. Therefore, oral hygiene behaviours are likely to be very sensitive to action planning, as they can easily be linked to other behaviours most people do every day, for example tooth brushing after the cue of eating a meal or before going to bed.

The intervention was tested in a study<sup>32</sup> consisting of 84 dental practices and 799 patients. The results of this pilot study<sup>32</sup> supported the theoretically framed intervention as an effective method of influencing oral hygiene beliefs and behaviour. Participants who received the intervention had significantly better behavioural (timing, duration, method), cognitive (confidence, planning) and clinical (plaque, gingival bleeding) outcomes than the participants in the control group receiving routine care.

# Training in the delivery of the personalised oral hygiene advice

Training in the delivery of the personalised OHA intervention was provided by a clinical member of the trial team to all dentists/hygienists within a dental practice randomised to this allocation. The content and the delivery of the intervention were standardised as a series of steps (see study documentation, available at

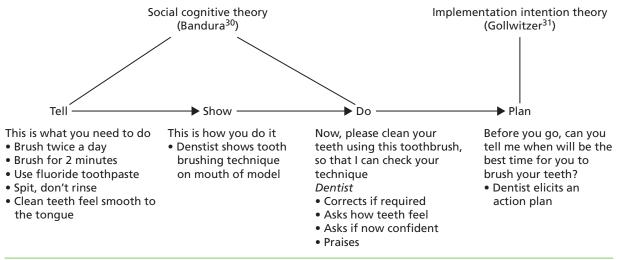


FIGURE 2 The OHA intervention behavioural framework.

www.journalslibrary.nihr.ac.uk/programmes/hta/090145/#/; accessed October 2017) designed to take place within an average primary care consultation, taking approximately 5 minutes in total.

A training DVD demonstrating a consultation using the steps was also developed and provided on a memory stick to all practices assigned to personalised OHA intervention (available at http://dentistry.dundee.ac.uk/nihr-hta-iquad-trial; accessed October 2017). The practices were also provided with laminated instruction sheets for use in the dental surgery. Dentists/hygienists retained these training resources in order to be able to undertake self-directed training as required throughout the trial. New clinicians appointed to any of the IQuaD trial dental practices (clusters) throughout the trial were invited to take part and provide consent to IQuaD if they had taken over the care of any IQuaD trial participants. Full training in trial methodology and intervention delivery was provided, as detailed above.

## Frequency of oral hygiene advice

At baseline, all participants received OHA in accordance with the cluster-level randomisation allocation. Reinforcement of OHA was provided at the discretion of the dentist/hygienist during the trial.

### Periodontal instrumentation

The definition of PI was as used in standard practice and would include the removal of plaque and calculus from the crown and root surfaces using manual or ultrasonic scalers, with no adjunctive subgingival therapy (e.g. antibiotics)<sup>33</sup> and the appropriate management of plaque retention factors.

## Baseline periodontal instrumentation

A full mouth supra- and subgingival PI was carried out by the practice dentist/hygienist on all participants prior to the participant or dental clinician being aware of the trial allocation. No time limit was set on this treatment and dentists/hygienists were instructed to scale the teeth and root surfaces until they were free of all deposits and were smooth to probing.

## Experimental periodontal instrumentation

Experimental groups received a PI at 6- or 12-monthly intervals in accordance with the individual participant-level randomisation. Participants allocated to the no-PI groups attended their dentist at time intervals determined by current practice. However, participants and dental practices were advised that every trial participant should be invited to attend their dentist for a routine examination appointment at least every 12 months.

# **Randomisation**

## Practice allocation to oral hygiene advice group

Recruited dental practices were allocated to routine or personalised OHA by minimisation on two factors: (1) practice employed a dental hygienist (yes/no) and (2) practice size (one or two dentists in practice/three or more dentists). This cluster-level randomisation was conducted using the automated, central randomisation service at the Centre for Healthcare Randomised Trials (CHaRT), University of Aberdeen, after the dental practice consent form was received at the Trial Coordinating Office in Dundee (TCOD) and before any potential participant had been approached.

## Participant allocation to periodontal instrumentation group

Participants were allocated to the PI trial arms using the automated, central randomisation service at the CHaRT, University of Aberdeen, with access by both telephone and web. Allocation took place once the OA had completed the baseline outcome assessment and was minimised on (1) absence of gingival bleeding on probing (yes/no), (2) highest sextant BPE score (BPE score of < 3/BPE score of 3) and (3) currently smoking (yes/no).

The OAs were informed that allocation had taken place but were blinded to allocation, with the actual allocation transmitted to the TCOD. A letter was sent to all participants to inform them of their Pl allocation and all participants received a £25 gift voucher in recognition of their contribution to the study. The TCOD provided each individual recruited dental practice with a list of projected dates for Pl and review (no-Pl group), according to the individual participant-level allocation, to be delivered throughout the trial for all their trial participants. The list was updated and re-sent by TCOD on an annual basis to each practice.

# **Descriptive measures**

Participant descriptive measures were collected at baseline and annually by self-administered postal questionnaire. They included the time since the last visit at the dental practice, the type of attendee (regular vs. non-regular), the self-reported number of PIs and OHA received in the last 12 months and by whom, smoking status and the type of toothbrush (manual vs. electric). Descriptive measures are presented in *Table 1* by year and randomised group, using either mean, standard deviation (SD) or n (%), as appropriate.

TABLE 1 Participant dental characteristics at baseline, by randomised group

	Randomised group, <i>n</i> (%)						
	PI			ОНА			
Participant dental characteristics	No (N = 623)	12-monthly (N = 625)	6-monthly ( <i>N</i> = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)		
Age (years), mean (SD), n	47.8 (15.8), 623	47.9 (15.7), 625	47.8 (15.8), 626	47.4 (16.1), 1008	48.3 (15.3), 866		
Male	223 (36)	229 (37)	225 (36)	387 (38)	290 (33)		
Date of last visit to the dental practice							
< 1 year ago	569 (91)	547 (88)	553 (88)	893 (89)	776 (90)		
1–2 years ago	38 (6)	57 (9)	49 (8)	79 (8)	65 (8)		
> 2 years ago	5 (1)	4 (1)	6 (1)	9 (1)	6 (1)		
Missing	11 (2)	17 (3)	18 (3)	27 (3)	19 (2)		
Last course of treatment was							
NHS	557 (89)	573 (92)	575 (92)	918 (91)	787 (91)		
Private	10 (2)	9 (1)	8 (1)	15 (1)	12 (1)		
Combination	40 (6)	23 (4)	24 (4)	41 (4)	46 (5)		
Missing	16 (3)	20 (3)	19 (3)	34 (3)	21 (2)		
Do you think of yourself as							
A regular attendee	565 (91)	561 (90)	574 (92)	903 (90)	797 (92)		
Someone who sees a dentist when in pain or trouble	43 (7)	47 (8)	31 (5)	76 (8)	45 (5)		
Missing	15 (2)	17 (3)	21 (3)	29 (3)	24 (3)		
Last time you went to the dental praction	ce were you giv	en OHA?					
Yes	410 (66)	411 (66)	420 (67)	688 (68)	553 (64)		
No	192 (31)	187 (30)	179 (29)	275 (27)	283 (33)		
Missing	21 (3)	27 (4)	27 (4)	45 (4)	30 (3)		

TABLE 1 Participant dental characteristics at baseline, by randomised group (continued)

	Randomise	d group, <i>n</i> (%)			
	PI			ОНА	
Participant dental characteristics	No ( <i>N</i> = 623)	12-monthly ( <i>N</i> = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)
By whom?					
Dentist	298 (48)	293 (47)	291 (46)	484 (48)	398 (46)
Hygienist	81 (13)	85 (14)	97 (15)	141 (14)	122 (14)
Both	29 (5)	27 (4)	27 (4)	53 (5)	30 (3)
Missing	215 (35)	220 (35)	211 (34)	330 (33)	316 (36)
Last time you went to the dental pract	cice were you giv	ven a scale and po	lish?		
Yes	371 (60)	364 (58)	381 (61)	619 (61)	497 (57)
No	230 (37)	236 (38)	217 (35)	341 (34)	342 (39)
Missing	22 (4)	25 (4)	28 (4)	48 (5)	27 (3)
By whom?					
Hygienist	106 (17)	108 (17)	114 (18)	182 (18)	146 (17)
Dentist	247 (40)	236 (38)	244 (39)	394 (39)	333 (38)
Missing	270 (43)	281 (45)	268 (43)	432 (43)	387 (45)
What type of toothbrush do you norm	nally use?				
Manual	413 (66)	397 (64)	382 (61)	647 (64)	545 (63)
Electric	180 (29)	192 (31)	205 (33)	300 (30)	277 (32)
Do not use toothbrush	14 (2)	18 (3)	16 (3)	28 (3)	20 (2)
Missing	16 (3)	18 (3)	23 (4)	33 (3)	24 (3)
Do you normally pay for dental treatm	ents?				
Yes	453 (73)	436 (70)	452 (72)	716 (71)	625 (72)
No	155 (25)	170 (27)	150 (24)	260 (26)	215 (25)
Missing	15 (2)	19 (3)	24 (4)	32 (3)	26 (3)
Do you have dental insurance?					
Yes	19 (3)	20 (3)	24 (4)	32 (3)	31 (4)
No	581 (93)	583 (93)	573 (92)	932 (92)	805 (93)
Missing	23 (4)	22 (4)	29 (5)	44 (4)	30 (3)
How often do you prefer to have a sca	ale and polish?				
Never	13 (2)	13 (2)	8 (1)	21 (2)	13 (2)
Once every 2 years	25 (4)	20 (3)	13 (2)	38 (4)	20 (2)
Once a year	117 (19)	122 (20)	121 (19)	193 (19)	167 (19)
Twice a year	269 (43)	268 (43)	289 (46)	451 (45)	375 (43)
Three times a year	65 (10)	65 (10)	59 (9)	89 (9)	100 (12)
Four times a year	67 (11)	71 (11)	56 (9)	103 (10)	91 (11)
More often	26 (4)	19 (3)	25 (4)	32 (3)	38 (4)
Missing	41 (7)	47 (8)	55 (9)	81 (8)	62 (7)

### **Outcome measures**

## **Primary outcomes**

- Clinical: gingival inflammation/bleeding on probing at the gingival margin at the 3-year follow-up.
- Patient centred: oral hygiene self-efficacy at the 3-year follow-up.
- Economic: net benefits (mean WTP minus mean costs).

# Secondary outcomes

- Clinical: (1) calculus, (2) clinical probing depth, (3) additional PI and (4) referral. (All of which were collected at 3-year follow-up.)
- Patient-centred: (1) dental quality of life (QoL), (2) oral health behaviour and (3) knowledge.
   (All of which were collected during 3 years' annual follow-up.)
- Economic: costs to the NHS and patients; WTP.
- Providers: beliefs relating to giving OHA and maintenance of periodontal health.

Note: the Periodontal Advisory Group considered that clinical attachment loss and plaque cannot be measured reliably; therefore, neither was included as outcomes.

# Collection of clinical outcome measures

Gingival inflammation was measured according to the Gingival Index of Löe<sup>34</sup> by running the UNC probe circumferentially around each tooth just within the gingival sulcus or pocket. After 30 seconds, bleeding was recorded as being present or absent on the buccal and lingual surfaces. The primary outcome (gingival inflammation/bleeding) was calculated by adding all the sites at which bleeding was observed and dividing these sites by the number of sites (twice the number of teeth) and presented as a percentage.

The colour-coded UNC periodontal probe was used to measure clinical probing depth and presence of calculus. Clinical probing depths were measured for all teeth (excluding third molars) at six sites per tooth: (1) mesiobuccal, (2) midbuccal, (3) distobuccal, (4) mesiolingual/palatal, (5) mid-lingual/palatal and (6) distolingual/palatal. Clinical probing depth was calculated as the mean of the six different sites measured per tooth and it is presented in mm.

Calculus was calculated by adding all the sites where calculus was observed and dividing it by the number of teeth and presented as a percentage.

The sequence of scoring was gingival inflammation/bleeding, clinical probing depths and calculus.

## Collection of patient-centred outcome measures

Patient-centred outcomes were measured at baseline and annually by self-administered postal questionnaire.

The full details of the calculations used to generate each patient-reported outcome are available in *Appendix 1*, *Section 1*: methods for computing patient-reported outcomes.

## Oral health belief outcomes

The questions for measuring patient-centred oral health belief outcomes were derived from social cognitive theory<sup>30</sup> and the theory of planned behaviour.<sup>31</sup>

The primary patient-centred outcome was self-efficacy, assessed as a person's confidence in their ability to perform several different oral health behaviours. Each behaviour was measured using a 7-point scale scored from 1 (not at all confident) to 7 (extremely confident), with 7 being the best outcome.

Perceived behaviour control (PBC) outcome, assessed in terms of perceived ease or difficulty of performing several different oral health behaviours, was measured using a 7-point scale varying from 1 (strongly agree) to 7 (strongly disagree), with 7 being the best outcome.

Attitude outcome (perceived consequences of the behaviour) was measured using a 7-point scale varying from 1 to 7 (strongly agree to strongly disagree), with 7 being the best outcome.

Subjective norm outcome (perceptions of social pressure to perform the behaviour) was measured using three questions with points varying from 1 (strongly agree) to 7 (strongly disagree), with 7 being the best outcome.

### Oral health behaviour outcome

Patient-reported oral health behaviour outcome was measured using four questions (about duration and frequency of brushing, frequency of flossing and frequency of interdental brushes use). Each response varied from 0 to 3, with a score of 3 being the best possible behaviour. The best value between flossing and interdental brushes was used as a measure of interdental cleaning behaviour. The responses for each question were summed to produce a summary score ranging from 0 to 9, with 9 being the best outcome.

### Intention outcome

Intention (motivation to perform a behaviour) was measured using three questions (about duration and frequency of brushing and frequency of flossing). Each response varied from 0 to 3, with a score of 3 being the best possible intention. The responses were summed to produce a summary score ranging from 0 to 9, with 9 being the best outcome.

## Quality of life

Quality of life was measured using the Oral Health Impact Profile-14 (OHIP-14).<sup>35</sup> The OHIP-14 is a 14-question, oral health-specific, patient-centred measure of symptoms experienced in the previous 12 months. The questions are scored from 0 (never) to 4 (very often) and summed to produce a summary score ranging from 0 to 56, with 56 being the worst outcome.

## Additional periodontal instrumentation outcome

The additional PI outcome assessed at the 3-year follow-up was the number of participants self-reporting in their annual questionnaire that they had received any private PI at any time during the trial. It was based on the question, 'In the last 12 months have you received a private scale and polish?'.

### Having a plan to floss or brush

Participants were asked in all questionnaires if they had a plan to brush better and if they had a plan to floss better. If they answered 'yes' to either question, they were considered to have a plan. This outcome was also used as a measure of fidelity of the personalised OHA intervention.

## **Dentist beliefs**

Clinician belief questionnaires were collected at baseline and at the 3-year follow-up. At baseline, the following variables were measured: self-efficacy, attitude, PBC, intention, subjective norm and action planning. All of the variables are measured on a scale of 1 to 7, except subjective norm (measured from 1 to 49) and intention (measured from 0 to 100). At follow-up, the variables measured were the same. Again, all variables varied from 1 to 7, except for subjective norm (varies from 1 to 49). The full details of the calculations used to generate each dentist belief outcome are available in *Appendix 1*, *Section 1: methods for computing patient-reported outcomes*.

## Post hoc outcomes

Self-reported bleeding was assessed using one question with the scale varying from 1 (never) to 5 (very often) at the 3-year follow-up in the annual questionnaire.

## Tertiary measures

Dental appearance was measured using four different questions (e.g. 'How clean and pleasant do your teeth look and feel after you brush and after a scale and polish'?). Each scale varied from 1 (not at all clean/not at all pleasant) to 7 (could not get any cleaner/extremely pleasant) and were summarised separately using the mean responses, with 7 being the best outcome.

Sensitivity was measured using four different questions. First, it was measured through a binary (yes/no) question: 'Do you experience sensitivity in your teeth?'. Second, it was measured through three different scales, the first varying from rarely sensitive to always sensitive and the other two scales varying from 1 to 7 (never to all the time, and no pain to worst imaginable pain). These tertiary outcomes were self-reported at the 3-year follow-up.

## Fidelity measures

Fidelity measures were collected at baseline and annually by self-administered postal questionnaire. These included whether or not participants had a plan to start brushing and/or flossing better and whether or not, after brushing, the patients did, or intended to, 'spit, but not rinse'.

Dental practice compliance with the protocol was monitored in two ways: (1) through annual face-to-face visits by a member of the trial office team and (2) through an annual audit of six participants. All practices received at least one face-to-face visit between baseline and follow-up. This was to ensure practice compliance with the protocol and an understanding of their role, to answer any queries the practice staff had and to build and maintain a rapport with the practices to ensure a smooth transition into the follow-up phase of the trial. The annual audit of six participants (two from each PI allocation) was conducted with each practice to check if participants had been contacted to attend an appointment according to their allocated treatment group. If  $\geq 50\%$  of these six random participants had not been contacted or invited to attend, this triggered a telephone call to the practice to check the trial processes and, if required, a visit to review protocol was arranged.

# Number of periodontal instrumentations received (routine data and self-report)

The self-reported number of Pls received in the previous 12 months was collected annually by postal questionnaire. If participants replied that they had not received any Pl in the previous 12 months, this response was recorded as a zero for those 12 months. If participants replied that they received Pl in the previous 12 months but did not indicate how many times, these responses were set as missing. The total number of received Pls over the 3 years was calculated by summing the number of times the participant self-reported receiving Pl over the course of the trial.

Routine treatment data were obtained from Information Services Division (ISD) Scotland and the NHS Business Services Authority (BSA) in England for the time period of 2010 to 2016. The routine data informed on the number of PI received throughout the trial, by counting the number of claims for PIs made by dentists for each participant. Further details regarding how these variables were calculated can be found in *Chapter 3*.

### **Data collection**

Assessment for eligibility and informed consent was achieved at the screening stage. Clinical outcome assessment was carried out at baseline and the 3-year follow-up. Participants were sent a trial questionnaire at baseline, 12 months, 24 months and 36 months following recruitment. Clinicians were asked to complete a belief questionnaire at baseline and at the 3-year follow-up.

### **Baseline**

Dentists and/or hygienists were asked to complete the CBQ at baseline, after each individual dental practice had consented to take part in the IQuaD trial.

Patient-centred outcomes were collected at baseline by self-administered questionnaire.

Arrangement of clinical assessment appointments has been outlined in the recruitment section above.

Clinical outcomes were measured at baseline by trained OAs who were blinded to allocation. Gingival inflammation/bleeding scores, calculus, clinical probing depth and BPE scores were measured by the OA and recorded on the baseline clinical chart by the dental research nurse who was a member of the trial team.

At the baseline appointment, the OAs also collected personal details from participants, including preferred contact method (telephone, e-mail) and contact details.

## Annual follow-up

Patient-centred outcomes were collected annually by self-administered postal questionnaires.

Like the baseline questionnaire, the annual follow-up questionnaire contained questions on self-efficacy, PBC, attitude, subjective norm, behaviour, OHIP-14, dental appearance and sensitivity. Questions designed to collect the descriptive measures were also included in this questionnaire, as were questions on dental costs for the health economic outcomes. Annual follow-up questionnaires at year 1 and year 2 post randomisation were posted from the CHaRT trial office to participants' home addresses, along with a covering letter and reply-paid envelope for return of the completed questionnaire. Those participants who failed to return their questionnaire within 3 weeks were sent a reminder letter, a further copy of the questionnaire and a reply-paid envelope. If questionnaires were returned to the trial office marked 'return to sender', every effort was made to obtain updated contact details from the participant's dental practice.

## Three-year follow-up

As at baseline, dentists and hygienists were asked to complete the clinician CBQ. These were collected by the OAs when they visited the dental practices to collect participant clinical outcomes.

Patient-centred outcomes were collected at the 3-year follow-up using the annual follow-up questionnaire, with an additional question on self-reported bleeding. The questionnaire was sent to participants at least 3 weeks before the date of the first 3-year follow-up appointment made at the dental practice where they were recruited. Participants who had not returned a questionnaire by the time of their follow-up appointment were asked to complete a shortened one-page version of the follow-up questionnaire containing questions on the primary patient-centred outcome (self-efficacy) at that appointment. Participants who did not attend the follow-up appointment and who had not returned a questionnaire were sent a reminder letter, a further copy of the questionnaire and a reply-paid envelope to return the completed questionnaire to the CHaRT trial office.

All trial participants were invited to attend a trial follow-up assessment appointment by their dental practice either at the time of their routine check-up or at a separate trial assessment appointment at which the clinical outcomes were measured by trained OAs who were blinded to allocation. As at baseline, gingival inflammation/bleeding scores, clinical probing depths and calculus were measured by the OAs and recorded on the follow-up clinical chart by the dental research nurse who was a member of the trial team.

Participants who could not attend were contacted and given the option of attending on at least one other day or time.

All participants who attended the follow-up assessment received a certificate of appreciation for their participation in the trial, details of when the trial results would be published and a £25 gift voucher in recognition of their contribution.

# Sample size

The study was powered to detect a difference of 7.5% in the number of gingival sites showing bleeding on probing at 3 years' follow-up between routine and personalised OHA and for each pairwise comparison across both routine and personalised OHA groups. An OHA exploratory trial<sup>32</sup> in the same population demonstrated that, at baseline, 35% of gingival sites were bleeding on probing with a SD of 25%. The PI Cochrane review<sup>24</sup> suggested that a reduction of 15% of sites with bleeding was a plausible reduction for 6-monthly PI. If the effect is assumed to be linear, halving the number of PIs should half the expected difference of 15% of sites. If the effect is non-linear and the difference is > 7.5%, the trial would be adequately powered. A smaller effect would be of questionable clinical significance. There is some evidence that personalised OHA can reduce the number of gingival sites that bleed on probing by approximately 7.5%.<sup>32</sup>

## Oral hygiene advice

Assuming a conservative estimate of the intracluster correlation coefficient (ICC) of 0.05, a cluster RCT of 50 dentists collecting information from 25 patient participants each ( $25 \times 25 = 625$  patient participants per arm) was expected to have 90% power to detect a difference of 7.5%. Should the ICC be 0.1, the trial would still have approximately 80% power to detect a difference of 7.5%.

### Periodontal instrumentation

Given that the comparison of routine versus personalised OHA required 625 participants in each arm, equal randomisation 1:1:1 (no PI, 12-monthly PI, 6-monthly PI) of participants implied 208 in each of the six groups. Assuming no interaction effect, the corresponding PI groups could be combined across both routine and personalised OHA groups, requiring 416 patients allocated to each PI group. Based on a sample size of 416 in each group, the trial has in excess of 95% power for each pairwise comparison to detect a difference of 7.5% in the percentage of gingival sites that bleed on probing.

### **Interaction**

A substantive interaction effect between the PI interventions and the personalised OHA was not expected. Assuming an ICC of 0.05, the trial had 80% power to detect an interaction effect of 7.5%. Should the ICC be 0.1, the trial had approximately 80% power to detect an interaction effect of 10%.

The total number of dentists required was 50, and the total number of participants was 1248 (6  $\times$  208). A previous trial<sup>36</sup> in general dental practice suggests that we should expect to lose a small number of dental practices from the trial for reasons such as practices amalgamating with other practices or restricting NHS patients; a conservative assumption of 17% attrition for dentists and 20% for participants was anticipated, requiring a minimum of 60 dentists and 1860 participants. Each dentist was required to recruit, on average, 31 participants to ensure 25 at follow-up.

## Collection of providers outcomes

Differences in the frequency of PI visits would have an impact on clinicians' costs and benefits. The effect on incomes, job satisfaction and changes to the level of fees on the provision of PI were assessed using self-reported guestionnaires administered to clinicians over the duration of the trial.

## Statistical analyses of outcomes

Statistical analyses were based on all participants randomised. Three principal comparisons and their interactions were tested in the factorial design. The principal comparisons were between (1) all those allocated to routine OHA and all those allocated to personalised OHA, (2) all those allocated to no PI and those allocated to 6-monthly PI and (3) all those allocated to 12-monthly PI and those allocated to 6-monthly PI.

Reflecting the clustered nature of the data, the clinical outcomes were compared using mixed models with dental practice as a random effect, and with adjustment for minimisation variables [practice employs dental hygienist (yes/no) and practice size (one or two dentists in practice/three or more dentists) for the cluster-level randomisation and absence of gingival bleeding on probing (yes/no), highest sextant BPE score (BPE score of < 3/BPE score of 3) and currently smoking (yes/no) for the patient-level randomisation] and participant baseline values (when available) as fixed effects. Patient-centred outcomes were compared using a mixed model with a random effect for the participant (to take into account the three-level nested and correlated data structure: a variable number of observations nested within participants and participants grouped in dental practices) and another for the centre and adjusting for the same variables as before. Statistical significance was at the 5% level, with corresponding confidence intervals (CIs) derived.

Pre-planned subgroup analyses on the primary outcome included exploration of participant age (< 45 years, 45–64 years or  $\geq$  65 years), smoking status (non-smoker or smoker), periodontal disease severity (no clinical signs or presence of gingival bleeding on probing) and intervention provider (dentist or practice hygienist). Post hoc subgroup analyses were undertaken by region (Scotland or England) and clinical pocket depth (four or more sites with clinical pocket depth of  $\geq$  4 mm, or fewer than four sites). These analyses were conducted by including a subgroup by treatment interaction term in the primary outcome model described above. Conservative levels of statistical significance (p < 0.01) were sought, reflecting the exploratory nature of these pre-planned and post hoc subgroup analyses.

## Non-response analysis

Descriptive data are presented, comparing the baseline characteristics of participants who did and who did not attend the 3-year follow-up clinical appointment. The *t*-test (continuous outcomes) and chi-squared test (dichotomous outcomes) were used to estimate the statistical significance of the differences between responders and non-responders.

#### Missing items

Missing items were dealt with according to the relevant published recommendations for the specific instruments.<sup>35</sup> For missing items with no published recommendation, self-efficacy, PBC, attitude and subjective norm were calculated as a mean of the items available. Behaviour and intention were calculated using complete cases.

### Sensitivity analysis

To assess the possible effect of outliers, treatment effects by individual dental practices were explored to assess the influence of between-centre differences in the primary clinical outcome: bleeding. Any practices where the treatment effect excluded zero from the 99% CI and had at least twice the target difference estimate (either –15% or 15%) were excluded from the analysis comparing no PI with 6-monthly PI, as well as from the subgroup analysis.

# **Trial oversight**

The University of Dundee acted as sponsor for the study. The trial was co-ordinated from the TCOD in the Dental Health Services Research Unit, University of Dundee, which provided day-to-day support for the dental practices and OAs/research nurses. The TCOD was responsible for transacting the randomisation of dental practices, collecting trial data (including baseline questionnaires) and co-ordination of participant

follow-up. CHaRT, in the Health Services Research Unit, University of Aberdeen, provided the database applications and information technology (IT) programming for the trial, hosted the randomisation system, co-ordinated the participant follow-up questionnaires, provided experienced trial management guidance and took responsibility for all statistical aspects of the trial [including interim reports to the Trial Steering Committee (TSC) and the Data Monitoring Committee (DMC)]. A Project Management Committee (PMG) met monthly, chaired by the chief investigator, comprising co-investigators in Newcastle upon Tyne and key members of the TCOD and CHaRT. A Trial Management Committee (TMC) met biannually, chaired by the chief investigator, comprising co-investigators and key members of the TCOD and CHaRT. A Periodontal Advisory Group was convened to provide expert clinical advice to the TMC.

Independent TSCs and DMCs were also established. The TSC comprised an independent chairperson (a professor and honorary consultant of dental public health) and two further independent members (both members of the public acting as patient representatives). The TSC met approximately annually over the course of the trial. The DMC comprised an independent chairperson (a professor of restorative dentistry) and two further independent members (a professor of dental public health and a statistician). The DMC met approximately annually.

# Patient and public involvement

Prior to the start of the IQuaD trial, patients were involved with the trial design and provided invaluable feedback on trial recruitment and communication strategies. Patients also contributed to the content and layout of the trial invitation, trial newsletters and the design of patient participant questionnaires. This ensured that trial participants could understand and easily complete these materials. Members of the public also contributed to trial oversight through membership of the TSC, including helping to interpret the trial findings and the preparation of the monograph.

### **Protocol amendments after trial initiation**

A number of protocol revisions were made after trial initiation. These included:

- an amended start date, list of grant holders, TSC members and DMC members
- clarification of the measurement of gingival bleeding and the list of grant holders was updated
- an amended timescale for sending patient information to 'at most 6 weeks', in line with routine practice for scheduling check-up appointments
- an amended end date, in line with approved contract variation.

Adaptations of study administrative processes (e.g. the use of additional letters, revisions to letters) were also implemented.

# **Chapter 3** Health economic methods

The health economics analysis assessed the cost–benefit of theory-based, personalised OHA or PI at different time intervals (no PI, 12-monthly PI or 6-monthly PI), or their combination for improving periodontal health in dentate adults attending general dental practice. A cost–benefit analysis (CBA) reporting incremental net benefit, relative to standard care (routine OHA with 6-monthly PI), was conducted alongside the cluster RCT. Costs were assessed from a NHS perspective and a wider societal perspective including both NHS and participant costs. Benefits were assessed using a general population's WTP for personalised OHA and PI, estimated from a discrete choice experiment (DCE). All analyses were based on the intention-to-treat (ITT) principle.

Cost—benefit analysis is based on the concept of maximising societal welfare including a wider, more holistic measure of value, going beyond narrow measures, such as quality-adjusted life-years, which are not appropriate in the context of dentistry. In CBA, both costs and benefits are measured in monetary terms and net benefits (benefits – costs) can be directly interpreted as either welfare increasing or welfare decreasing. The CBA conducted here is an incremental analysis of alternative policies of OHA with/without PI in terms of costs and benefits (WTP). The analysis focuses on whether moving from standard care to an alternative policy increases or decreases welfare.

## **Resource use and costs**

Health service costs and Participant costs describe the methods for generating NHS and participant perspective costs, respectively.

### Health services costs

### Use of NHS dental services

Resource utilisation data for NHS treatments at dental practices over the trial follow-up period were collected using routine sources held by the ISD of the Scottish Government and the NHS BSA in England. Dentists are paid on the basis of claims and, therefore, relatively rich and reliable routine data on NHS dental care provided are available. Dental claims data were linked to the trial data set on an individual level to each trial participant.

Different remuneration systems are in place across the UK, which has implications for the level of detail that is available on resource use.

In Scotland, NHS payments are based on fee-for-service contracts. There is a high level of detail, as fees are attached to each individual item of service. This includes a separate fee for PI. Although there is a separate fee for intensive hygiene instruction, this cannot be claimed if a PI was claimed within the same course of treatment or within the previous 5 months.

In England, dentists are reimbursed according to contracts negotiated at (1) national level [General Dental Service (GDS) contracts negotiated through the British Dental Association] or (2) local level (personal dental contracts). The former accounts for 85% of dental contracts in England and forms the basis for the trial costing analysis.<sup>37</sup> Under the GDS contracts, each dental practice is contracted to provide an agreed amount of dental treatment, described in terms of UDAs. Treatments provided to participants are grouped into four bands based on the complexity of the treatment. Each band carries a set number of UDAs,<sup>38</sup> as follows:

- 1. Band 1 (check-up, radiography, advice, PI) = 1 UDA.
- 2. Band 2 (band 1 treatments plus fillings, extractions, root canal treatments) = 3 UDAs.
- 3. Band 3 (band 1 and band 2 work plus crowns, dentures, bridges) = 12 UDAs.
- 4. Urgent band = 1.2 UDAs.

A number of services do not fall into any of the patient charge bands in their own right, but are awarded UDAs when provided outside other banded treatments. The number of UDAs for such treatments is as follows:

- Arrest of bleeding = 1.2 UDAs.
- Bridge repair = 1.2 UDAs.
- Denture repair = 1.2 UDAs.
- Issue of prescriptions = 0 UDAs.
- Removal of sutures = 1 UDA.

As dentists in England are reimbursed on the basis of UDAs, the level of detail on resource use is clearly much lower than in Scotland. The claim form (FP17) in England collects some further information on treatments provided, including PI, but these are not associated with additional payments. Given the structure of the contract, it is less likely that there will be a difference in costs across the different policies in England as PIs and personalised OHA are not associated with any additional UDAs if they are provided alongside other treatments such as a dental check-up.

Costs were attached directly to items of service (Scotland) and treatment course bands (England) using the appropriate NHS unit cost. Unlike other NHS services, patients contribute to the cost of their dental care. The cost to the NHS is therefore equal to the unit cost minus the appropriate patient charge. The Scotlish *Statement of Dental Remuneration – Amendment No 130 Letter*<sup>39</sup> was used to attach unit costs to the treatment items in Scotland. In England, the cost of a UDA delivered under the GDS contracts can vary widely by practice. Determining an average cost per UDA across England is not straightforward. The 2009 Steele report<sup>40</sup> suggested that the value of a UDA was approximately £25, but that actual practice-specific values ranged widely from £17 to £40. Work informing National Institute for Health and Care Excellence (NICE) public health guidance on oral health promotion<sup>41</sup> assumed a mean cost of £25 (95% CI £15 to £35). We conducted a descriptive analysis of published data for contract payments in England in 2014/15<sup>37</sup> and found that the median gross value (prior to deduction of participant charges) of a UDA (after removal of payments for units of orthodontic activity) is still approximately £25, ranging from £20 to £33 (5th to 95th percentile). Therefore, we use a unit value of £25 per UDA for the analysis. The NHS cost per UDA is varied in the sensitivity analysis to explore the impact of practice-level variation on results.

Some patients do not pay a co-charge (e.g. those in receipt of income support, pension credit, universal credit or tax credit, as well as those aged < 18 years, in full-time education or who are pregnant or who have given birth in the last 12 months). *Table 2* summarises the most common breakdown of costs in England and Scotland. When participants were exempt from treatment charges, the full cost was attributed to the NHS budget. We tested for any cross-group differences in the proportion of participants exempt from charges, as any differences could bias predictions of incremental NHS costs. We also repeated the cost analysis assuming that all participants were exempt, that is, the NHS bears the full cost.

Data are in the form of courses of treatment that have a commencement date and a claim payment date (Scotland) or a completion date (England). The base-case analysis excludes the baseline and final study visits, as treatments provided at these visits are not part of routine care nor the trial intervention. The exception is the provision of personalised OHA at the baseline visit. However, as explained above, these are typically not associated with additional payments in either region. A sensitivity analysis was conducted to explore the impact of including an additional fee for personalised OHA.

The baseline visit is excluded from the analysis by removing all treatment claims with an acceptance date < 90 days following the baseline examination date. Owing to practice-specific approaches to calling participants for their final study visit, some had their final study visit prior to randomisation and others after 3 years post randomisation. To remove all final visits from the claims data, we excluded all treatment acceptance dates at > 2.5 years following the baseline study visit. This implies that, after exclusion of all baseline and final study visits, the follow-up period for the CBA is 2.25 years. Sensitivity analysis explores the impact on net benefits of including all the baseline and final clinic visits in the analysis.

TABLE 2	Summary	of UK	dental	charges,	by region

Region						
England			Scotland			
Treatment category	Total UDA cost (£)	<sup>a</sup> Patient charge (£) <sup>38</sup>	NHS cost (if charge is paid) (£)	Treatment category	Patient charge <sup>a</sup>	NHS cost
Band 1	25.00	18.50	6.50	Check-ups and case assessments	0% of full cost	100% of full cost
Band 2	75.00	50.50	24.50	Other treatments	80% of full cost	20% of full cost
Band 3	300.00	219.00	81.00	Maximum charge per course of treatment	£384.00	Unlimited
Urgent	30.00	18.50	11.50			

a Exceptions and exemptions apply to treatment in both regions when patients do not pay any of the cost (e.g. receiving income support, aged < 18 years, in full-time education, pregnant or have given birth in the last 12 months, receiving pension credit, receiving universal credit or receiving tax credit exemptions). Patient charges relate to 2014/15 values.

Routine data downloads were taken in August 2016 (ISD, Scotland) and November 2016 (NHS BSA, England). For Scotland, data are a complete and accurate reflection of resource use up to 3 months prior to data download (i.e. May 2016). For England, the data are likely to be an accurate reflection of resource use up to April 2016, reflecting the end of the financial year, by which time dental practices are required to have submitted all relevant information. The final clinic examinations took place in July 2016. Given that the base-case analysis excludes claims with treatment acceptance dates beyond 2.5 years post randomisation, the latest treatment claims in the data set have acceptance dates in February 2016 and, therefore, all data are assumed to provide an accurate and complete reflection of resource use for both regions.

All costs are presented in 2014/15 GBP values and discounted at a rate of 3.5% per annum.<sup>42</sup> A sensitivity analysis varying the discount rate was conducted.

## Costs of other NHS services during follow-up

Annual questionnaires collected data on use of other NHS services for problems related to participants' teeth. These included secondary care resource use [hospital inpatient data, outpatient consultations, day-case procedures and accident and emergency (A&E) attendances] together with general practitioner (GP) appointments and contact with NHS 24 (Scotland) or NHS Direct (England). As is common with participant-reported resource use, a number of conservative assumptions were made to address issues of discrepancy in the reported data:

- When respondents reported that they did not attend a service but entered a number of visits, it was assumed that no resource use was incurred.
- When respondents stated that they attended hospital as an inpatient for problems related to their teeth but provided no further information, it was conservatively assumed that such responses should be treated as day-case admissions. National data<sup>43,44</sup> show that the majority of hospital admissions in the UK for dental problems are day-case procedures.

Rates of secondary care visits, including outpatient, day-case and hospital admissions, were checked against practice records and average rates in the general population to assess their validity. The outcomes of the validation exercise are reported in *Chapter 5*.

# Participant costs

Three sources of participant cost were considered for the analysis: (1) charges for NHS dental care, (2) privately purchased dental care products and treatments and (3) time and travel costs (for participants and companions) associated with visits to the dental practice.

## Charges for NHS dental care

Participant charges were sourced directly from the routine data sets described in *Health services costs*. Participant charges were based on the co-charge that should have been collected at the dental practice, as opposed to the actual value paid by the participant. This reflects the way in which dental practices are paid.

# Self-purchased dental care products and private dental care

Participants reported how often they used interdental brushes, electric toothbrushes and manual toothbrushes in the annual questionnaires, as well as how often they replaced electric toothbrush heads. These resource use data were combined with an assumed average market price for four commonly available products, including a range of high- and low-cost items. Details of unit costs applied are provided in *Appendix 2*, *Section 1: additional detailed methods for costing, the discrete choice experiment and mapping discrete choice experiment valuations to the trial outcomes*. Resource use was multiplied by the average unit cost to generate a cost per participant.

Furthermore, in each annual questionnaire, participants were asked to recall and report the total cost of private PI and any other privately purchased dental care that they incurred over the previous year. It should be noted that, although participants were asked to report how often they received private PI, it may be difficult in some scenarios to determine whether treatment was private or via the NHS because, for example, many NHS dentists may also offer private PI. For this reason, there may be a risk of double-counting from the participant perspective, but only if NHS co-charges were interpreted as private treatment. All reported data were summed across the questionnaires to generate a cost to each participant of self-purchased dental care products and stated private care costs.

### Costs of time and travel

Participants incur costs travelling to, and attending, NHS dental appointments. The number of visits to the dentist was collected from the routine data sets, conservatively assuming that each unique treatment acceptance date (rather than each claimed item) relates to a single dental visit. The true costs may be higher if a course of treatment involved more than one visit to the dentist. The opportunity cost to participants and companions of making a return journey to visit the dental practice was estimated using responses to the participant time and travel cost questions in the baseline questionnaire. It is assumed that the data collected on opportunity cost at baseline are applicable to each subsequent visit to the dental practice. National unit cost sources (*Table 3*) were used to value time commitment of attendance (including travelling to and from) at a dental appointment, as well as the opportunity cost based on the reported activity forgone through attendance. The calculated opportunity cost for a single visit was multiplied by the number of treatment courses from the routine data to generate a total participant level estimate of the opportunity cost of time and travel.

### Combined participant and NHS perspective costs

Data from *Health services costs* and *Participant costs* were combined to generate a wider combined NHS and participant perspective of the costing analysis.

## Statistical analysis of cost data

Cost data were analysed according to best practice methodology for cluster randomised trials.<sup>52–54</sup> We used multilevel mixed-effects models that account for clustering of the data through a cluster-level random effect.<sup>52</sup> Models for cost data were estimated using the *meglm* command in Stata® 14 (StataCorp LP, College Station, TX, USA) specifying a gamma family and log-link function for cost data to account for skewness of the distribution. The decision to use this model for costs was based on a combination of the results of a modified Park's test, which identified both an inverse Gaussian and gamma distribution to be acceptable, and software restriction (only gamma family allowed within Stata's *meglm* command).

TABLE 3 Unit costs for the opportunity costs of time and travel

Activity	Assumptions made	Unit cost (£)	Reference
Paid work	Median weekly wage: £528; 39.1 hours per week	13.50	Annual Survey of Hours and Earnings: 2015 Provisional Results, ONS <sup>45</sup>
Self-employed	Average of two reports, 52 weeks,	15.13	The Boox Report 2014, Boox <sup>46</sup>
	40 hours per week <sup>a</sup>		Cost Converter, EPPI Centre <sup>47</sup>
			Self-employed Workers in the UK – 2014, ONS <sup>48</sup>
Transport	Cost per mile <sup>b</sup>	0.45	Expenses and Benefits: Business Travel Mileage for Employees' Own Vehicles, HMRC (approved millage rate) <sup>49</sup>
Caring for a relative or friend	Median gross weekly pay: £341; 39.2 hours per week <sup>c</sup>	8.70	<i>NHS Pay Review Body</i> , p. 17 (2015 values), DHSC <sup>50</sup>
Leisure activities	Value of non-working time	6.81	WebTAG: TAG data book, July 2016, TAG <sup>51</sup>
Childcare	Assumed as paid work	13.50	Annual Survey of Hours and Earnings: 2015 Provisional Results, ONS <sup>45</sup>
Voluntary work	Assumed as paid work	13.50	Annual Survey of Hours and Earnings: 2015 Provisional Results, ONS <sup>45</sup>
Unemployed	Value of non-working time	6.81	WebTAG: TAG data book, July 2016, TAG <sup>51</sup>
Retired	Value of non-working time	6.81	WebTAG: TAG data book, July 2016, TAG <sup>51</sup>
Parking	Participant's individual costs	Various	Participant questionnaires
Housework	Cost of housework in the NHS, assumed annual salary £21,000 gross, 2012 values inflated to 2015	10.56	NHS Pay Review Body, p. 17 (2015 values), DHSC <sup>50</sup>

DHSC, Department of Health and Social Care; EPPI, Evidence for Policy and Practice Information; HMRC, Her Majesty's Revenue and Customs; ONS, Office for National Statistics; TAG, Transport Analysis Guidance.

The log-link function was decided on as it allowed easier model convergence and had a preferred Akaike information criterion score than an identity link. The models were estimated by maximum likelihood. Incremental costs were estimated as the mean cost difference between randomised policies (relative to the policy most representative of standard care: routine OHA with 6-monthly PI), together with 95% CIs. All models were adjusted for baseline characteristics (age, age-squared and sex) as well as minimisation variables (cluster level: hygienist employed at practice and practice size; individual level: smoker, gingival bleeding at baseline and BPE score band).

Owing to the high level of completeness, descriptive analyses of NHS dental costs and participant co-charges were based on complete cases. Participant costs were presented and analysed using multiple imputation of missing cost data from participant annual questionnaires. For the analysis of total costs within the CBA, the small proportion of missing routine data were also imputed to complete the data set. The imputation process followed best practice guidelines.<sup>55,56</sup> All imputation was completed using Stata's multiple imputation procedure. Missing cost data were imputed at the level of cost component (e.g. missing data on private PI imputed for each annual questionnaire). Costs were imputed using predictive mean matching accounting for the repeated measures nature of the costs from the annual participant questionnaires. The average of the five closest values was used for each imputed data point. Data were imputed separately for

a Average hours worked: 40. Median annual income (two reports): £10,800 + £50,820. 2013/14 prices: £30,810, inflated to £31,476 (2015 values).  $£31,476 \div 52$  weeks  $\div 40$  hours/week = £15.13 per hour.

b Based on government-approved travel mileage reimbursement for use of own vehicles.

c Caring, leisure and other service occupations.

each randomised group to preserve the allocation of participants. Imputation models were adjusted for cluster, age and sex as well as minimisation covariates. Sixty imputed data sets were used to ensure stable results and were combined using Rubin's rules to generate estimates of costs.<sup>57</sup>

# **Benefits (willingness to pay)**

The main outcome measure in the economic evaluation is WTP, estimated using a DCE administered via online survey panels to a representative sample of the UK general population. The DCE was administered to a separate sample from the general population while the trial was ongoing and before the results of the trial were known. A DCE is a survey method for which participants are asked to make hypothetical choices between different goods or, in this case, different packages of dental care provided over 3 years (the follow-up period of the trial). An underlying assumption of all DCEs is that a treatment's or service's value depends on its attributes and the levels of those attributes. By including the price proxy within the DCE (i.e. the annual cost of each dental package), we can obtain a monetary valuation for any given dental treatment package. These WTP estimates are used as a measure of benefit within the CBA. Ethics approval for the conduct of the DCE substudy was obtained from the College Ethics Review Board at the University of Aberdeen (REF 2015/12/1278).

### Selection of attributes and levels

The relevant attributes were, to a large extent, determined by the trial. Preferences needed to be elicited for the specific interventions (PI and personalised OHA) and for the outcomes that may vary across the arms of the trial (i.e. self-reported bleeding gums and perception of appearance and feel of cleanliness). We followed recommended practice to assess whether or not these attributes were important to individuals and whether or not there were any other important attributes that should be included in the DCE. We conducted a literature review and focus groups (FGs) with the general population to develop the final list of attributes and levels to present in the DCE (see *Appendix 2, Section 2: research conducted for questionnaire development*). We also engaged with, and sought advice from, a range of stakeholders, including clinical dental experts, practising dentists and hygienists, and patient representatives on our trial advisory group about the DCE content. *Table 4* shows the final list of attributes and levels included in the DCE.

Additional levels were added for provider of service (dentist/hygienist) as this was found to be an important consideration in the FGs. The FGs showed that respondents would have difficulty understanding and attaching value to the primary clinical trial outcome (gingival inflammation/bleeding). Therefore, we included the self-reported frequency of bleeding measure from the participant questionnaire instead. Respondents were informed that having bleeding gums increases the risk of tooth loss in the future. The look and feel attribute was also based on the measures used within the same questionnaire. The cost attribute was framed as an annual commitment over a 3-year time period (trial follow-up). The levels of the cost attribute were determined from a number of sources, including the FGs, baseline trial data on WTP for PI and a supplementary questionnaire asked to FG respondents. *Figure 3* provides an example choice set included in the DCE.

## Experimental design

Choices presented to respondents in the DCE were drawn from the set of all possible attribute and level combinations. With a single three-level attribute and four five-level attributes, the total number of combinations in the full factorial is given as  $3^1 \times 5^4 = 1875$ , resulting in  $[(1875 \times 1874) \div 2] = 1.76$  million unique choice sets. Following standard practice in DCE literature, a D-efficient design was used. So Ngene version 1.1.2 software (Ngene, ChoiceMetrics Pty Ltd, Sydney, NSW, Australia) was used to create a main effects design consisting of 30 choice set questions. To ensure a manageable number of choice tasks, the tasks were split into three blocks, each with 10 questions. Two further choice questions were added to each block to check the validity of responses as follows: (1) choice 11 repeated choice 5 to test for consistency of preferences and (2) choice 12 was a non-satiation (dominance) test in which one option

TABLE 4 Attributes and levels included in the DCE

Attributes	Levels
Dental advice	No detailed or personalised advice
	Detailed and personalised advice from the dentist
	Detailed and personalised advice from the hygienist
Scale and polish	None
	One per year from the dentist
	One per year from the hygienist
	Two per year from the dentist
	Two per year from the hygienist
Your teeth will look and feel	Very unclean
	Unclean
	Moderately clean
	Clean
	Very clean
In 3 years' time, your gums will bleed	Never
	Hardly ever
	Occasionally
	Fairly often
	Very often
The cost to you	£10 per year (total cost: £30 over 3 years)
	£20 per year (total cost: £60 over 3 years)
	£50 per year (total cost: £150 over 3 years)
	£100 per year (total cost: £300 over 3 years)
	£200 per year (total cost: £600 over 3 years)

should be preferred over its alternatives if individuals value PI polish and personalised OHA. Therefore, each respondent completed a total of 12 choice tasks. The responses to the internal validity choice tasks were not included in the analysis; therefore, a total of 10 choice sets were analysed for each respondent.

As respondents were likely to vary considerably in terms of how many Pls they usually receive and the extent to which they experience bleeding gums, a segmented pivoted design was used to increase realism. For example, it is unrealistic that a respondent who has no bleeding gums and never has a Pl would suddenly experience bleeding very often if they opt out. Respondents were assigned to one of three segments [(1) good, (2) moderate or (3) poor dental health] based on self-reported frequency of bleeding gums and the look and feel of their teeth (both attributes in the DCE). The pivoted design tailored the presented levels based on the segment to which the respondents were assigned. For example, respondents categorised as having 'good' dental health would never be presented with the worst level of bleeding gums and unclean teeth, as it would be highly unlikely for dental health to deteriorate so quickly because of forgoing Pl or detailed OHA. The list of attribute levels included in the experimental design for the good, moderate and poor dental health segments can be found in *Appendix 2*.

NIHR Journals Library www.journalslibrary.nihr.ac.uk

	Dental Care Package A	Dental Care Package B	No Dental Care Package	
Dental Advice	Detailed and personalised Provided by the dentist			
Scale and Polish	None	Two per year Provided by the hygienist	None	
In three years time, your gums will bleed	Very often	Occasionally	Very often	
Your teeth will look and feel	Unclean	Very unclean	Very unclean	
The cost to you	£10 per year (Total cost: £30 over 3 years)	£100 per year (Total cost: £300 over 3 years)	£0 per year (Total cost: £0 over 3 years)	

FIGURE 3 Example choice task. Note: the example choice set shown here is for a respondent categorised as having poor dental health. See Appendix 2, Section 1: additional detailed methods for costing, the discrete choice experiment and mapping discrete choice experiment valuations to the trial outcomes, for more information on segmentation of choice sets based on dental health category. This hypothetical respondent has chosen Dental Care Package B and is willing to pay £100 per year for the package.

## Questionnaire development

Think-aloud interviews were used to test the wording and understanding of the questionnaire. Extensive piloting was used to assess the best way of reducing hypothetical bias and to ensure that attributes and levels were sensible, realistic and tradable to the majority of respondents. Full details of all methods used to select attributes and levels (literature searches and FGs) and test their feasibility in the survey (think aloud and piloting) are outlined in *Appendix 2*, *Section 2*: research conducted for questionnaire development.

The survey was divided into three sections. Section 1 asked about respondents' experiences of dental care, how often they attend their dental practice and who they are treated by. Section 2 included questions about bleeding gums and aesthetic appearance to determine the appropriate version of the DCE to present. Section 3 concluded the survey with demographic questions and a number of debriefing questions about respondents' views and attitudes to dental topics. The questionnaire was designed using Qualtrics (Provo, UT, USA).

### Data collection

Data were collected using Qualtrics online panels. The survey was nationally representative, with population census quotas for age (among adult population), sex and region sought. We oversampled in Scotland (n = 125) to enable subgroup analysis across the main participating regions in the trial (England/ Scotland). We further sought to achieve a mix of respondents with recent experience of dental services and without. During the pilot phase, we attempted to achieve 30% of responses from non-regular attenders, defined as not having seen the dentist in the previous 2 years. However, we were unable to achieve this sample in the pilot and relaxed the target to a more achievable 10% for the main phase of data collection. Responses were anonymised and respondents were free to leave the survey at any point without having to give a reason for doing so. Respondents who completed the survey were reimbursed in a manner determined by the survey panel from which they were partaking.

## Data analysis

The DCE data were analysed using best practice methods<sup>59</sup> and followed random utility maximisation theory.<sup>60</sup> The utility of the option  $(V_j)$  is a linear function of the attributes and levels presented to the respondents, where:

$$V_{j} = \alpha ASC + \beta_{1}Advice D + \beta_{2}Advice H + \beta_{3}One PI per year D + \beta_{4}One PI per year H \\ + \beta_{5}Two PI per year D + \beta_{6}Two PI per year H + \beta_{7}Bleed Hardly Ever + \beta_{8}Bleed Occasionally \\ + \beta_{9}Bleed Fairly Often + \beta_{10}Bleed Very Often + \beta_{11}Look unclean + \beta_{12}Look Moderately Clean \\ + \beta_{13}Look Clean + \beta_{14}Look very clean + \beta_{15}Annual Cost.$$
 (1)

Alternative-specific constant (ASC) is accounting for latent or unobserved utility associated with choosing any package, regardless of the attribute levels,  $\beta$  represents the marginal utilities associated with the attributes and levels. All categorical variables were included in the model using effects coding. The advantage of effects coding is that the reference level of an attribute is uncorrelated with the ASC. The reference categories for the effects coded attributes are no OHA, no PI, never bleeding and having teeth that look and feel very clean. For example, the value for 'no advice',  $\beta_{no \, advice}$ , is retrieved as the negative sum of  $\beta_1$  Advice  $D + \beta_2$  Advice H.

The marginal WTP of an attribute level is equal to the marginal rate of substitution between that level and the cost attribute. For ease of interpretation, we also estimate the difference in WTP between the reference level and the attribute level. For example, the annual marginal WTP for OHA from the hygienist compared with the reference level (no OHA) is equal to:

$$-\left(\frac{\beta_2 - \beta_{\text{no advice}}}{\beta_{15}}\right). \tag{2}$$

Mixed logistic regression models in preference space were used to analyse the data to determine the impact of the attributes and levels on preferences for dental care services and outcomes. The ASC was assumed random and normally distributed for all models, reporting mean and SD. All other attribute levels were fixed effects, reporting mean coefficients only. The final model was estimated using the maximum simulated likelihood method, with 2000 Halton draws. All models were estimated using Stata 14 software.

The model was re-estimated separately using interaction terms for several pre-determined subgroups including (1) sex; (2) region, to determine if system of reimbursement has an impact on preferences; (3) smoking status, to determine if current or previous smokers' preferences differ from the overall group; and (4) household income, to determine if greater ability to pay (annual income of  $\geq$  £20,800) had an impact on preferences. Further subgroup analyses explored if the preferences of those with experience of PI or of treatment from a hygienist differed from the overall sample. We planned to conduct subgroup analysis by dental attendance (regular/non-regular attenders) but were unable to achieve sufficient numbers of respondents attending the dentist less often than every 2 years to power such an analysis. The impact that subgroups had on the main attribute level effects was assessed by the significance of interaction terms between subgroup identifier and attribute level coefficient. Likelihood ratio tests were used to test the joint significance of the interaction terms.

# Estimating benefits (willingness to pay) for each trial participant

Benefits measured as WTP were estimated for each trial participant by attaching WTP values from the DCE to the relevant treatments provided (PI and personalised OHA) and to the relevant trial outcomes (self-reported bleeding when brushing and aesthetic outcome). The DCE provided estimates of WTP for zero, one and two PIs provided per year or a total of zero, three or six PIs over the trial period. A step-wise linear utility function was used to estimate WTP for the number of PIs other than those presented in the DCE. It was assumed that the provider of the baseline PI (dentist or hygienist) also delivered the intervention over follow-up. The impact that the results of this assumption have is also explored in a sensitivity analysis.

Willingness to pay for personalised OHA was attached based on randomised group, as all participating practices complied with their allocation to provide either personalised or routine OHA at the baseline visit. For the remaining attributes, including aesthetic outcome and frequency of bleeding gums when brushing, WTP values were directly attached at an individual level in the trial. The full mapping process for each attribute to the trial interventions and outcomes can be found in *Appendix 2*.

Discounting was applied at a rate of 3.5% per annum.<sup>42</sup> Discounted marginal WTPs were summed to generate a total WTP for each trial participant. Total WTP data were analysed using multilevel mixed-effects models with a random effect capturing cluster effects to account for the hierarchical nature of the trial. A normal distribution was assumed for benefits data; hence, the analyses were implemented using Stata's *mixed* command. All models were adjusted for individual- and cluster-level minimisation variables, age and sex. The base-case analysis undertakes statistical imputation of missing data on benefits (i.e. WTP mapped to trial outcomes). Data were imputed for annual questionnaires at the level of WTP for each outcome (e.g. bleeding gums), following a similar procedure to that outlined for costs.

# **Cost-benefit analysis**

The CBA compares the costs and benefits of alternative policies of OHA with/without PI with standard care (routine OHA with 6-monthly PI). Both costs and benefits are measured in monetary terms and net benefits can therefore be directly estimated (benefits – cost). The incremental net benefits indicate whether moving from standard care to an alternative policy increases or decreases overall societal welfare.

# Statistical analysis of (incremental) net benefit

The imputed data sets were used to estimate net benefits and incremental net benefits. Net benefit was calculated for each participant in the trial according to *Equation 3*:

Net benefit = total benefits - total costs. (3)

Multilevel mixed-effects models were used to analyse the data, implemented using Stata's 'mi estimate: mixed' command following the approach of Gomes  $et~al.^{52.53}$  These models can explicitly recognise clustering in parameter estimates though the incorporation of cluster-level random effects ( $u_i^c$ ,  $u_j^e$ ), while also adjusting for cluster- and individual-level covariates (minimisation variables: age and sex). The model addresses both individual- and cluster-level correlation between costs and benefits, using a bivariate normal distribution of the error terms. The model assumes linear additive effects for treatments and covariates. Restricted maximum likelihood was used to estimate the model for the base-case analysis. Interaction terms were included to retrieve the true impact of randomisation on costs/benefits. Stata's mimargins command was used to retrieve the values of the reference category of cost/benefit from the imputed data sets and the nlcom command was used to calculate incremental costs, incremental benefits and incremental net benefits across groups, together with 95% Cls.

## Presentation of cost-benefit analysis results

The base-case perspective is that of the UK NHS. A wider perspective, including both NHS and participant costs, was considered as a secondary analysis. All results are presented as incremental costs, incremental benefits and incremental net benefits, relative to standard care. Note that exclusion of dominated policy options is not required in the absence of a threshold value of a ratio between costs and benefits (as would be the case in a cost–utility analysis).

Confidence ellipses, based on parametric calculations using the model variance—covariance matrix, were used to illustrate the uncertainty in each comparison of costs and benefits graphically on the cost—benefit plane. The confidence ellipses illustrate the probability that each intervention is less/more costly and less/more beneficial than standard care. All policies with positive incremental net benefits are associated with higher levels of welfare than standard care. Note that this can be generated by a higher level of benefit and/or a lower level of cost of the policy. The strategy with the greatest incremental net benefit would be considered the preferred policy from a welfare perspective. The analysis is solely based on incremental changes in net benefits between policies and, as is common for economic evaluations alongside trials, makes no statement about the magnitude of the underlying net benefits of specific policies.

## Analysis of uncertainty

The confidence ellipses of the cost–benefit plane illustrate the joint uncertainty surrounding estimates of incremental costs and incremental benefits on results. Further deterministic sensitivity analyses, outlined in *Table 5*, investigated the impact of assumptions around cost data, benefit estimation and model structure on the results.

## Cost-consequences analysis

In addition to the results of the CBA conducted for the primary economic analysis, we further considered all of the available information across all components of the study, using a narrative cost–consequences analysis (CCA) framework. The CCA is a useful approach to summarise all the study information succinctly for decision-makers. We use a balance sheet approach summarising the advantages and disadvantages of each policy option under consideration. The CCA includes the following information:

- the results of clinical effectiveness, using both primary and secondary outcomes defined in the protocol
- the results of all economic analyses, including costs to the NHS and participants, general population preferences for PI/OHA (results from the DCE) and overall incremental net benefits
- a summary of information sourced from the CBQ investigating the practitioner's perception of the advantages and disadvantages of different policy strategies.

**TABLE 5** Deterministic sensitivity analyses considered

Base-case analysis	Sensitivity analysis	Justification for approach
Cost of personalised OHA incorporated according to routine claims database	Requirement of an additional consultation to provide personalised OHA	There was no additional cost of providing personalised OHA for most participants. An additional fee is included in sensitivity analysis according to the SDR for intensive hygiene instruction
Baseline and final visits excluded from analysis	Baseline and final treatment visits included	Base-case analysis pragmatically excludes treatment initiated at trial-driven visits. Sensitivity analysis incorporates all visits, including treatment requirements identified at baseline (treatment claim acceptance dates on or after randomisation date) or at follow-up (treatment acceptance dates included up to 3 years post randomisation) trial visits. Changes applied to both costs and WTP
Mapping aesthetic DCE outcome to a <i>Right now</i> response to the trial data	Mapping aesthetic DCE outcome to a <i>After brushing teeth</i> response to the trial data	Analysis conducted to address uncertainty about the most appropriate trial outcome to map marginal WTP onto
Unit cost per UDA assigned at the national average level (£25)	Vary the cost per UDA $\pm 20\%$	Payment per UDA substantially varies across practices. The base-case analysis gives the best estimate for the average cost per UDA in England, giving the most generalisable results. The sensitivity analysis explores the impact of practice level variation
Costs and benefits discounted at a rate of 3.5% per annum	Varying the discount rate between 0% and 6%	Standard exploration of methodological uncertainty
Professional delivering baseline intervention also delivered follow-up appointments (i.e. dentist or hygienist)	Explore assumptions in which all OHA and PI delivered by dentist; all OHA and PI delivered by hygienist	Anecdotal evidence from the trial suggests that the professional delivering baseline intervention may not have delivered all treatments for a participant over the follow-up period. The sensitivity analysis explores the likely impact of this assumption on results

SDR, Statement of Dental Remuneration (Scotland)

No attempt was made to quantitatively synthesise the information across different categories. The results of the CCA are presented qualitatively as a means to aid decision-makers' understanding of the overall outcomes of the trial as well as the trade-offs between clinical and economic outcomes.

# **Chapter 4** Trial results and clinical effectiveness

This chapter describes the clinical effectiveness results of the IQuaD trial. The chapter starts with an explanation of how the trial groups were derived. It then describes the study groups at trial entry. The results at the annual follow-up points are then reported, followed by a formal statistical analysis of the data for the principal measures of outcome.

# **Recruitment to the study**

Recruitment of dental practices began in October 2011 and continued until May 2013. Participant recruitment to the trial began in February 2012 and continued until July 2013. Data were closed to follow-up on 2 September 2016. The derivation of the main study groups and their progress through the stages of follow-up in the trial is shown in *Figure 4*, in the form of a Consolidated Standards of Reporting Trials (CONSORT) flow diagram.

## **Dental practices**

Sixty-eight dental practices were initially recruited in total, with 49 in Scotland and 19 in England: 34 practices were allocated to give routine OHA and 34 to provide personalised OHA. From the 68 practices, four were excluded post randomisation in the routine OHA arm and one in the personalised OHA arm. The exclusions were due to a trial procedural error in the early recruitment phase of the study whereby practices were randomised when a dentist in a practice had verbally given consent to take part but before the practice principal dentist (usually the owner of the practice) had signed a consent form for the practice to take part in the trial. The principal dentist did not give permission to use their practice. The practice study intervention allocation was not known to any potential dentist, practice principal dentist (or patient) participants at the time of non-consent. Selection bias due to the post-randomisation exclusions was not, therefore, possible. Hence, there were 63 recruiting practices in total.

## **Participants**

A total of 2341 patients were screened for trial entry and 183 (8%) of the screened patients were found to be ineligible (*Table 6*). The primary reason for ineligibility was due to a BPE score of 4 or \* (160 patients). From those ineligible as a result of a BPE score of 4 or \*, 144 (90%) patients agreed to join a separate cohort group. Further details about the cohort participants are given in *Chapter 6*. There were 281 patients eligible for the study who were not recruited. The main reason for non-recruitment was that the patient did not attend the baseline appointment (see *Table 6*).

Participants were recruited in 63 dental practices (see *Appendix 3*). A total of 1877 participants were recruited to the study, with 867 in the routine OHA arm (n = 289 allocated to receive no PI, n = 290 allocated to receive 6-monthly PI and n = 288 allocated to receive 12-monthly PI) and 1010 in the personalised OHA arm (n = 334 allocated to receive no PI, n = 337 allocated to receive 6-monthly PI and n = 339 allocated to receive 12-monthly PI). Of the total participants, 1348 (72%) were recruited across Scotland, including 121 (6%) recruited from the Scottish islands. No dental practice recruited > 2.5% of the participants. The average cluster size was 30.6 and 28.9 participants for personalised OHA and routine OHA practices, respectively. Three participants were subsequently identified to be non-NHS patients at consent (and therefore ineligible) and were thus excluded post randomisation.

# **Description of the groups at trial entry**

### **Practice characteristics**

*Table 7* provides the dental practice characteristics and beliefs. The majority of practices employed hygienists and had three or more dentists. Participating dentists and hygienists completed a questionnaire

NIHR Journals Library www.journalslibrary.nihr.ac.uk

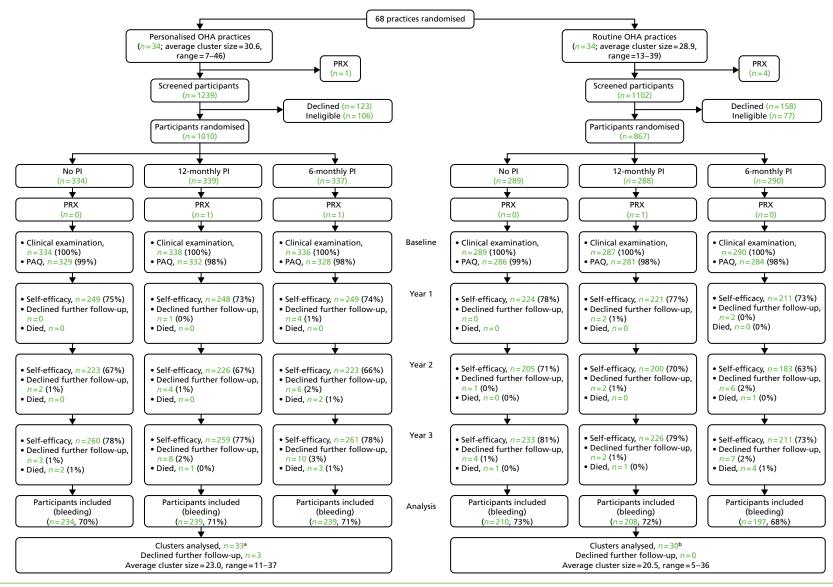


FIGURE 4 The Consolidated Standards of Reporting Trials (CONSORT) diagram. PAQ, patient annual questionnaire; PRX, post-randomisation exclusion. a, one was a post-randomisation exclusion; b, four were post-randomisation exclusions.

**TABLE 6** Screened/declined

	OHA, n (%)	
Total ineligible/total declined	Personalised (N = 1239)	Routine ( <i>N</i> = 1102)
Total declined	123 (10)	158 (14)
Unable to complete the study	4 (< 1)	24 (2)
Did not attend baseline appointment	64 (5)	72 (7)
No reason stated	53 (4)	61 (6)
Observing Ramadan	_	1 (< 1)
Probes locked in safe and patient could not wait	1 (< 1)	-
Did not have time to consider	1 (< 1)	-
Total ineligible	106 (9)	77 (7)
BPE score of 4 or *	92 (7)	68 (6)
Uncontrolled medical condition	13 (1)	5 (1)
Too nervous	1 (< 1)	2 (< 1)
Not a regular attendee	_	2 (< 1)

**TABLE 7** Practice characteristics/beliefs

	OHA, n (%)	
Dental practices	Personalised (N = 33) <sup>a</sup>	Routine (N = 30) <sup>a</sup>
Employs hygienists	24 (73)	24 (80)
Has three or more dentists	25 (76)	24 (80)
Region		
Scotland	24 (72.7)	20 (66.7)
England	9 (27.3)	10 (33.3)
	OHA, n (%)	
Health professional beliefs	Personalised (N = 87) <sup>b</sup>	Routine ( <i>N</i> = 75) <sup>b</sup>
Dentist	70 (81)	63 (84)
Hygienist	17 (20)	12 (16)
Beliefs, mean (SD), n		
Self-efficacy	5.9 (0.7), 81	5.9 (0.7), 67
Attitude	5.4 (0.7), 81	5.4 (0.7), 67
PBC	3.7 (1.0), 81	3.7 (0.8), 67
Intention	90.0 (14.2), 77	85.7 (17.2), 64
Has a plan to give OHA	2.2 (1.4), 79	2.4 (1.4), 67
Has a plan to give Pl	2.2 (1.3), 78	2.3 (1.3), 66
Subjective norm	13.5 (9.2), 78	11.3 (7.3), 67

a Number of practices by cluster randomised allocation.

b Number of responders to the baseline CBQ by cluster randomised allocation.

at baseline (the CBQ) on their beliefs about providing OHA. Scores for belief variables varied from 1 to 7, where 7 represented the most positive outcome (e.g. the highest level of confidence or the most positive attitude). Intention and subjective norm were the exceptions: intention varied from 0 to 100, where 100 represented the highest level of intention, and subjective norm varied from 1 to 49, where 49 represented the highest level of subjective norm. The results suggested that dental professionals had positive attitudes towards, and high levels of intention of, providing OHA. There was little perceived influence of peers in the ability of the respondents to provide OHA (subjective norm) (see *Table 7*). Overall, there were no important differences between groups on any of the practice characteristics or beliefs at baseline.

# Participant characteristics

## Sociodemographic factors and dental characteristics

Participant and sociodemographic characteristics are shown in *Table 1*. *Tables 1* and *8–16* follow a consistent format to aid interpretation of the main cluster and participant comparisons. Columns 2–4 of data represent the participant-level comparison and show the no PI, 12-monthly PI and 6-monthly PI groups, respectively. Columns 5 and 6 of data represent the cluster-level comparison and show personalised OHA and routine OHA, respectively.

The average age of participants was 48 years; the majority (around 65%) were women and regular attendees of the dental practice (> 90% reported that their last visit to the dentist was within 2 years prior to baseline). Around 60% reported having received OHA and a PI at their last dental appointment. For those who reported receiving one treatment or the other, the dentist was most often the provider of the OHA or PI. Overall, 64% of patients used a manual toothbrush and > 90% had no dental insurance. There were no substantive differences between groups on any of the sociodemographic factors or dental characteristics.

## Clinical characteristics

Table 8 displays the results of the clinical assessment at baseline. The mean number of teeth per participant was 24. The highest sextant BPE score was  $\leq 2$  in two-thirds of the participants. The mean per cent of sites bleeding was 33%, and 35% of teeth had calculus present. The mean clinical probing depth was 1.8 mm and between 10% and 12% of participants in each group had four or more sites with a clinical probing depth of  $\geq 4$  mm.

There were no important imbalances across the randomised groups.

### Patient-reported outcomes

The baseline patient-reported outcomes summary is shown in *Table 9*. More details about individual questions that produced the scales are given in *Appendix 1*.

Self-efficacy, the patient-reported primary outcome, was, on average, 5.2 points (on a scale of 1 to 7 points with 7 points meaning very confident).

## Cognition variables

Perceived behaviour control was, on average, 4.5 points; attitude towards oral health behaviour was very positive and was, on average, 5.8 points and subjective norm was 5.3 points (scales of 1 to 7, with 7 meaning very confident).

## **Behaviour**

The mean self-reported oral hygiene behaviour score was 4.7 points. The scale varies from zero (poorest hygiene behaviour) to 9 (perfect hygiene behaviour). The behaviour scale had space for improvement across the three behaviours measured: (1) frequency of brushing, (2) duration of brushing and (3) frequency of interdental cleaning (either flossing or the use of interdental brushes). In both brushing frequency and duration, around 10% of participants scored the maximum of 3 points (brushed more than twice a day, for

TABLE 8 Clinical characteristics at baseline, by randomised group

	Randomised group, mean (SD), <i>n</i>					
	PI	PI			ОНА	
Clinical characteristics	No ( <i>N</i> = 623)	12-monthly (N = 625)	6-monthly ( <i>N</i> = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)	
Present teeth	23.7 (4.5), 623	23.6 (4.5), 625	23.7 (4.7), 626	23.6 (4.6), 1008	23.7 (4.5), 866	
Highest sextant BPE score,	n (%)					
0	_	4 (1)	1 (0)	2 (0)	3 (0)	
1	40 (6)	42 (7)	38 (6)	73 (7)	47 (5)	
2	376 (60)	374 (60)	377 (60)	590 (59)	537 (62)	
3	207 (33)	205 (33)	210 (34)	343 (34)	279 (32)	
Gingival bleeding, n (%)	608 (98)	610 (98)	612 (98)	991 (98)	839 (97)	
Current smoker, n (%)	135 (22)	143 (23)	147 (23)	213 (21)	212 (24)	
% of sites bleeding	33.5 (23.8), 623	32.5 (23.9), 625	32.4 (22.9), 626	34.3 (23.2), 1008	31.0 (23.7), 866	
% of teeth with calculus	35.5 (26.3), 623	35.9 (27.4), 625	34.9 (26.8), 626	33.7 (25.8), 1008	37.4 (27.8), 866	
Mean clinical probing depth (mm)	1.8 (0.3), 623	1.8 (0.3), 625	1.8 (0.3), 626	1.8 (0.3), 1008	1.8 (0.3), 866	
Four or more sites with a clinical probing depth of $\geq 4$ mm, $n$ (%)	64 (10)	75 (12)	70 (11)	117 (12)	92 (11)	

TABLE 9 Patient-reported outcomes at baseline, by randomised group

	Randomised group, mean (SD), <i>n</i>							
	PI		ОНА					
Patient-reported outcomes	No ( <i>N</i> = 623)	12 monthly (N = 625)	6 monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)			
Self-efficacy	5.2 (1.1), 612	5.3 (1.2), 607	5.2 (1.2), 607	5.2 (1.2), 976	5.2 (1.2), 850			
PBC	4.5 (1.2), 607	4.5 (1.2), 603	4.5 (1.2), 606	4.5 (1.2), 972	4.4 (1.2), 844			
Attitude	5.8 (1.2), 611	5.9 (1.1), 605	5.7 (1.3), 607	5.8 (1.2), 977	5.8 (1.2), 846			
Subjective norm	5.3 (1.1), 601	5.3 (1.2), 598	5.2 (1.1), 597	5.3 (1.1), 958	5.2 (1.1), 838			
Behaviour	4.7 (1.7), 608	4.7 (1.8), 604	4.8 (1.7), 605	4.6 (1.7), 971	4.9 (1.7), 846			
Intention	5.5 (1.7), 560	5.5 (1.8), 568	5.6 (1.8), 556	5.4 (1.8), 903	5.6 (1.7), 781			
OHIP-14	6.0 (7.4), 595	6.4 (8.1), 591	6.5 (8.3), 591	5.9 (7.6), 952	6.7 (8.3), 825			

longer than 2 minutes). Regarding interdental cleaning, around 20% of participants scored the maximum of 3 points (floss or use interdental brushes every day).

### Intention

The mean intention to perform good OHA practice was 5.5 points (on a scale from 0 to 9, with 9 being the best outcome).

## Quality of life

The mean score on the oral health QoL scale, OHIP-14, was 6.3 points on a scale form 0 to 56 (with 0 indicating perfect oral health QoL). The OHIP-14 75th percentile was 9 and the 99th percentile was 39, illustrating a generally good oral health-related QoL in this population.

The baseline participant self-reported data suggested that participants were confident that they could perform good toothbrushing practice, thought oral hygiene practice was a positive thing to do, were influenced by their peers to do it and intended to do it. However, the participants did not currently actually perform good oral hygiene practice and reported that there were external factors, PBC, that limited their ability to do so. Overall, across the randomised groups, there were no apparent imbalances in the patient-reported outcomes.

# **Trial follow-up**

## Attendance at 3-year clinical examination

The primary clinical outcome was collected at the 3-year clinical follow-up. Overall, 71% of the participants attended the appointment (see *Appendix 1*). Twelve participants were known to have died by the end of the 3-year follow-up. The main reasons for non-attendance were that the practice was unable to contact the participant (41%), the participant was unable to attend (30%) and the participant did not want to attend (13%). There were no important differences between groups for the reasons of non-attendance.

A comparison of baseline characteristics between those who did and those who did not attend the final appointment was undertaken. The results in *Table 10* show that only a few factors differed between attenders and non-attenders. Attenders were, on average, older (50 vs. 43 years). Although there was a statistically significant difference in the OHIP-14 score, with the mean score reported by non-attenders higher than that reported by attenders, the difference of 1.3 points (0.15 of the SD of the score) was not thought to be clinically important. There was no evidence that non-attenders had different levels of disease, with BPE scores, bleeding, calculus and probing depths all very similar to those seen in attenders.

## Annual questionnaire returns at years 1, 2 and 3

Approximately 77% of participants completed a follow-up questionnaire at 3 years. There were no substantive differences in response rates between the randomised groups. The overall rates of return of follow-up questionnaires at 1, 2 and 3 years are shown in *Appendix 1*.

## Participant dental characteristics at 3 years

The participant dental characteristics at 3 years are shown in *Table 11*. The profile of characteristics is similar across the groups and similar to baseline responses (e.g. smoking status, type of toothbrush used, preferences for two Pls per year; see *Tables 1* and 8). Characteristics for years 1 and 2 are given in *Appendix 1*, *Section 2: participant dental characteristics (years 1 and 2)*.

TABLE 10 Comparison of baseline characteristics of 3-year follow-up assessment of attenders and non-attenders

	Attendance, mean (SD), i		
Baseline characteristics	Attender ( <i>N</i> = 1327)	Non-attender (N = 547)	<i>p</i> -value
Age (years)	49.9 (15.1), 1327	42.7 (16.1), 547	< 0.001
Male, <i>n</i> (%)	490 (37)	187 (34)	0.27
BPE score, n (%)			0.28
0	5 (0)		
1	82 (6)	38 (7)	
2	788 (59)	339 (62)	
3	452 (34)	170 (31)	
Bleeding n (%)			0.78
Yes	1295 (98)	535 (98)	
No	32 (2)	12 (2)	
Self-efficacy	5.2 (1.1), 1306	5.2 (1.2), 520	0.50
OHIP-14	5.9 (7.3), 1276	7.2 (9.2), 501	0.003
% of sites bleeding	32.4 (23.1), 1327	33.9 (24.4), 547	0.20
% of calculus	34.7 (26.3), 1327	37.2 (28.0), 547	0.08
Mean clinical probing depth (mm)	1.8 (0.3), 1327	1.8 (0.3), 547	0.64

TABLE 11 Patient and dental characteristics at 3 years, by randomised group

	Randomise	Randomised group, n (%)				
	PI	PI			ОНА	
Patient and dental characteristics	No (N = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)	
Date of last visit to the dental practice (years ago)						
<1	393 (63)	386 (62)	383 (61)	630 (63)	532 (61)	
1–2	17 (3)	21 (3)	12 (2)	24 (2)	26 (3)	
> 2	3 (0)	2 (0)	_	4 (0)	1 (0)	
Missing	210 (34)	216 (35)	231 (37)	350 (35)	307 (35)	
Do you think of yourself as						
A regular attendee	399 (64)	394 (63)	388 (62)	632 (63)	549 (63)	
Someone who sees a dentist when in pain or trouble	13 (2)	16 (3)	6 (1)	24 (2)	11 (1)	
Missing	211 (34)	215 (34)	232 (37)	352 (35)	306 (35)	
Last time you went to the dental	oractice were	you given OHA?				
Yes	281 (45)	339 (54)	327 (52)	534 (53)	413 (48)	
No	131 (21)	70 (11)	67 (11)	121 (12)	147 (17)	
Missing	211 (34)	216 (35)	232 (37)	353 (35)	306 (35)	
By whom?						
Dentist	210 (34)	217 (35)	178 (28)	357 (35)	248 (29)	
Hygienist	39 (6)	65 (10)	83 (13)	90 (9)	97 (11)	
Both	30 (5)	56 (9)	66 (11)	86 (9)	66 (8)	
Missing	344 (55)	287 (46)	299 (48)	475 (47)	455 (53)	

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

TABLE 11 Patient and dental characteristics at 3 years, by randomised group (continued)

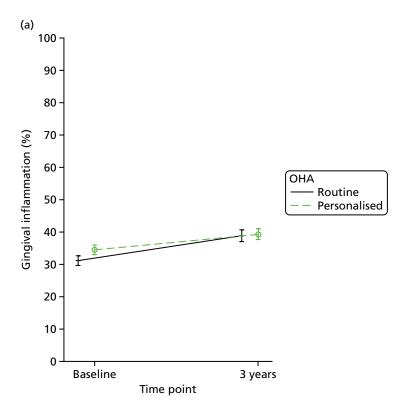
	Randomis	Randomised group, n (%)				
	PI	PI			OHA	
Patient and dental characteristics	No ( <i>N</i> = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)	
In the last 12 months, did you	In the last 12 months, did you receive a scale and polish?					
Yes	210 (34)	334 (53)	354 (57)	491 (49)	407 (47)	
No	199 (32)	72 (12)	40 (6)	161 (16)	150 (17)	
Missing	214 (34)	219 (35)	232 (37)	356 (35)	309 (36)	
By whom?						
Dentist	145 (23)	212 (34)	198 (32)	317 (31)	238 (27)	
Hygienist	54 (9)	111 (18)	138 (22)	151 (15)	152 (18)	
Both	8 (1)	8 (1)	13 (2)	16 (2)	13 (2)	
Missing	416 (67)	294 (47)	277 (44)	524 (52)	463 (53)	
Smoked in the last 12 months	?					
Yes	50 (8)	55 (9)	45 (7)	76 (8)	74 (9)	
No	362 (58)	355 (57)	349 (56)	580 (58)	486 (56)	
Missing	211 (34)	215 (34)	232 (37)	352 (35)	306 (35)	
What type of toothbrush do yo	ou normally use?					
Manual	233 (37)	223 (36)	199 (32)	345 (34)	310 (36)	
Electric	158 (25)	154 (25)	167 (27)	261 (26)	218 (25)	
Do not use toothbrush	21 (3)	31 (5)	27 (4)	48 (5)	31 (4)	
Missing	211 (34)	217 (35)	233 (37)	354 (35)	307 (35)	
How often do you prefer to have a scale and polish?						
Never	19 (3)	12 (2)	11 (2)	24 (2)	18 (2)	
Once every 2 years	20 (3)	9 (1)	15 (2)	20 (2)	24 (3)	
Once a year	108 (17)	113 (18)	50 (8)	144 (14)	127 (15)	
Twice a year	184 (30)	196 (31)	212 (34)	322 (32)	270 (31)	
Three times a year	27 (4)	40 (6)	49 (8)	71 (7)	45 (5)	
Four times a year	41 (7)	25 (4)	46 (7)	47 (5)	65 (8)	
More often	4 (1)	10 (2)	10 (2)	19 (2)	5 (1)	
Missing	220 (35)	220 (35)	233 (37)	361 (36)	312 (36)	

# **Statistical analyses**

# **Primary outcomes**

## Gingival inflammation/bleeding

The pre-chosen clinical primary outcome was mean gingival bleeding at the 3-year follow-up. The means and 95% CI for baseline and follow-up for each randomised group are displayed in *Figure 5*. All randomised groups followed a similar pattern: bleeding increased from baseline (mean 32% of sites) to 3-year follow-up (mean 39% of sites). The differences between groups and corresponding 95% CIs are shown in *Table 12*.



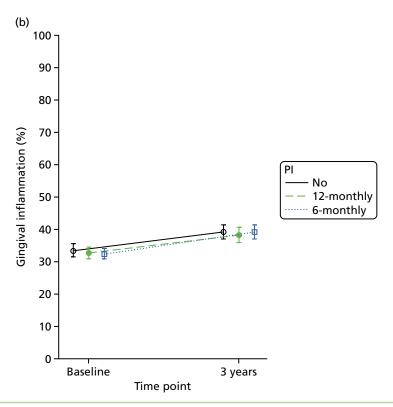


FIGURE 5 Gingival inflammation/bleeding (mean and 95% CI), by randomised allocation. (a) OHA delivery and (b) PI interval.

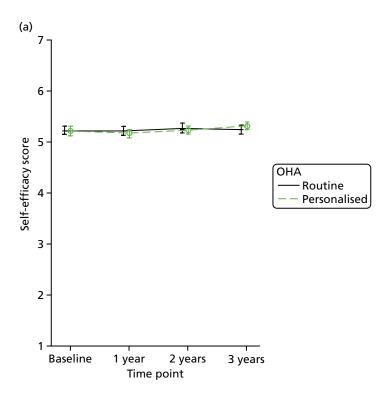
**TABLE 12** Treatment effects for primary outcomes

	Estimate, p-value (95% CI)			
Primary outcomes	No PI vs. 6-monthly PI	12-monthly PI vs. 6-monthly PI	Personalised OHA vs. routine OHA	
Gingival inflammation/bleeding				
Unadjusted	0.71 (-2.0 to 3.7), 0.579	-0.53 (-3.4 to 2.3), 0.715	0.02 (-6.3 to 6.4), 0.985	
Adjusted for baseline bleeding	0.8 (-1.7 to 3.2), 0.534	0.1 (-2.4 to 2.5), 0.947	-1.9 (-8.1 to 4.3), 0.549	
Adjusted for baseline bleeding and minimisation variables	0.87 (-1.6 to 3.3), 0.481	0.11 (-2.3 to 2.5), 0.929	-2.5 (-8.3 to 3.3), 0.393	
Self-efficacy				
Unadjusted	-0.017 (-0.133 to 0.098), 0.767	-0.082 (-0.198 to 0.033), 0.162	0.002 (-0.144 to 0.147), 0.984	
Adjusted for baseline self-efficacy	-0.098 (-0.189 to -0.007), 0.035	-0.027 (-0.118 to 0.064), 0.564	0.003 (-0.110 to 0.115), 0.962	
Adjusted for baseline self-efficacy and minimisation variables	-0.028 (-0.119 to 0.063), 0.543	−0.097 (−0.188 to −0.006), 0.037	0.017 (-0.089 to 0.123), 0.750	

Three types of analyses are reported: (1) an unadjusted analysis, (2) an adjusted-for-baseline bleeding analysis and (3) a fully adjusted model accounting for baseline bleeding and minimisation variables (this is the prespecified primary model). Under ITT analysis, there was no evidence of a statistical or clinical difference between those randomised to receive 6-monthly PI and those randomised to receive no PI (difference 0.87%, 95% CI –1.6% to 3.3%; p = 0.481). Similarly, there was no evidence of a difference between 6-monthly PI and 12-monthly PI (difference 0.11%, 95% CI –2.3% to 2.5%; p = 0.929). The 95% Cls were small enough to exclude the prespecified clinically important difference of 7.5% in bleeding. There was also little evidence of a difference between participants randomised to personalised OHA and those randomised to routine OHA (difference -2.5%, 95% CI -8.3% to 3.3%; p = 0.393), although the 95% CI did not entirely rule out a 7.5% reduction. The results were robust to other adjusted/unadjusted models. The interaction between personalised OHA and 6-monthly PI was 1.7% (95% CI -3.8% to 7.3%). The interaction demonstrated that a participant receiving both personalised OHA and 6-monthly PI would have a further 1.7% bleeding reduction compared with just adding the individual effects of 6-monthly PI with personalised OHA. The interaction term ruled out an additional 7.5% reduction. The ICC at follow-up for bleeding for all participants was 0.23 (95% CI 0.16 to 0.31). A full description of the means (SD) for gingival inflammation at each time point is given in Appendix 1, Section 3: clinical and patient-reported outcomes by year of follow-up.

### Self-efficacy

The pre-chosen patient-centred primary outcome was self-efficacy at the 3-year follow-up. The means and 95% CIs for baseline and follow-up for each randomised group are displayed in *Figure 6*, by randomised allocation. All groups followed a similar pattern. The score was similar from baseline (mean 5.2 points) up to the year 3 questionnaire (mean 5.3 points) in both the PI and the OHA groups. The differences between groups and corresponding 95% CIs are shown in *Table 12*. Under ITT analysis, there was no evidence of a difference between those randomised to receive 6-monthly PI and those randomised to receive no PI (difference -0.028, 95% CI -0.119 to 0.063; p = 0.543). Between those randomised to receive 6-monthly and those randomised to receive 12-monthly PI, there was a statistically significant difference, favouring the 6-monthly PI (difference -0.097, 95% CI -0.188 to -0.006; p = 0.037). However, this result is not clinically significant. There was also no evidence of a difference between participants randomised to personalised OHA and those randomised to routine OHA (difference 0.017, 95% CI -0.089 to 0.123; p = 0.750). The interaction coefficient between personalised OHA and 6-monthly PI was -0.007 (95% -0.22 to 0.20),



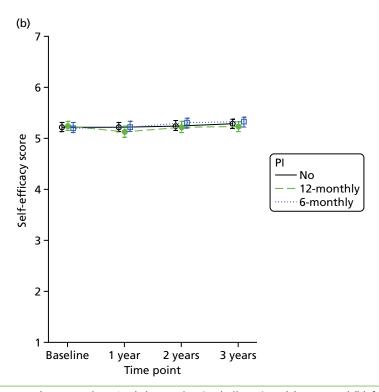


FIGURE 6 Self-efficacy score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

suggesting no evidence of important interaction between interventions. The self-efficacy ICC for all participants was 0.06 (95% CI 0.04 to 0.09). A full description of the means (SD) for self-efficacy at each time point is given in *Appendix 1*, *Section 3*: clinical and patient-reported outcomes by year of follow-up.

## Secondary outcomes

A full description of the means (SD) for the secondary outcomes at each time point is given in *Appendix 1*, *Section 3: clinical and patient-reported outcomes by year of follow-up*.

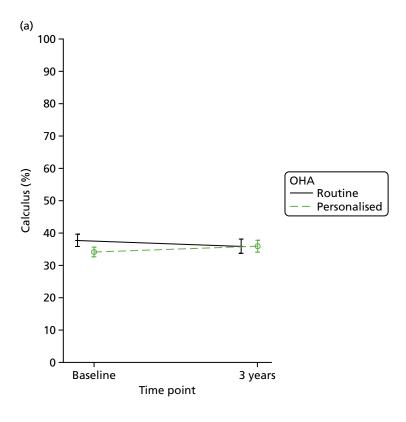
The clinical secondary outcomes were calculus and mean clinical probing depth collected at the 3-year follow-up. The patient-reported secondary outcomes were PBC; attitude; subjective norm; behaviour and intention collected at 1, 2 and 3 years; and private PI received throughout the trial collected at year 3. Self-reported bleeding was a post hoc secondary outcome collected at 3 years. The differences between groups, and corresponding 95% Cls, are shown in *Table 13*.

#### Calculus

The calculus means and 95% CIs for baseline and follow-up for each randomised group are displayed in *Figure 7*. The mean level of calculus decreased across the no PI, 12-monthly and 6-monthly PI groups. The calculus level was statistically significantly higher in the no-PI group than in the 6-monthly PI group (mean 8.0%, 95% CI 5.4% to 10.7%, p < 0.001). There was no evidence of a difference in mean calculus between 12-monthly PI and 6-monthly PI or between personalised and routine OHA (*Table 13*).

**TABLE 13** Treatment effects for secondary outcomes

	Estimate, <i>p</i> -value (95% CI)		
Secondary outcomes	No PI vs. 6-monthly PI	12-monthly PI vs. 6-monthly PI	Personalised OHA vs. routine OHA
Clinical			
% of calculus	8.0 (5.4 to 10.7), < 0.001	1.6 (-1.0 to 4.2), 0.231	2.3 (-5.8 to 10.3), 0.577
Mean clinical probing depth (mm)	0.003 (-0.024 to 0.030),	0.022 (-0.004 to 0.049),	-0.024 (-0.084 to 0.036),
	0.808	0.102	0.433
Patient-centred outcomes			
PBC	0.02 (-0.09 to 0.13),	-0.01 (-0.12 to 0.10),	-0.02 (-0.12 to 0.07),
	0.704	0.838	0.620
Attitude	-0.137 (-0.273 to -0.001),	-0.093 (-0.229 to 0.044),	-0.019 (-0.134 to 0.097),
	0.048	0.183	0.754
Subjective norm	-0.07 (-0.18 to 0.04),	-0.06 (-0.17 to 0.05),	-0.04 (-0.14 to 0.06),
	0.198	0.272	0.473
Behaviour	-0.125 (-0.256 to 0.007),	-0.164 (-0.296 to -0.032),	0.072 (-0.063 to 0.207),
	0.064	0.015	0.297
Intention	-0.10 (-0.25 to 0.05),	-0.14 (-0.29 to 0.00),	0.09 (-0.06 to 0.25),
	0.186	0.054	0.253
OHIP-14	0.35 (-0.23 to 0.93),	0.41 (-0.17 to 0.99),	-0.33 (-0.86 to 0.20),
	0.239	0.169	0.229
Private Pls, odds ratio (95% Cl), <i>p</i> -value	0.5 (0.3 to 0.7), 0.001	0.8 (0.6 to 1.2), 0.302	1.0 (0.7 to 1.6), 0.908
Has a plan to better brush or floss, odds ratio (95% CI), <i>p</i> -value	1.1 (0.9 to 1.3), 0.563	0.94 (0.77 to 1.16), 0.635	1.1 (0.9 to 1.4), 0.579
Self-reported bleeding	-0.01 (-0.14 to 0.12),	-0.03 (-0.16 to 0.10),	-0.06 (-0.17 to 0.06),
	0.886	0.666	0.333



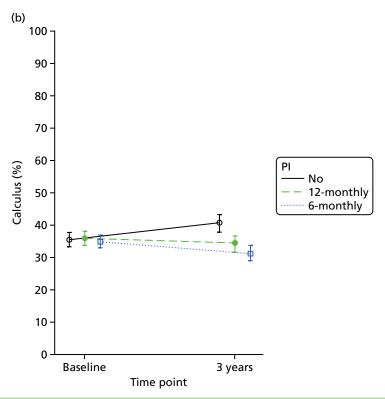
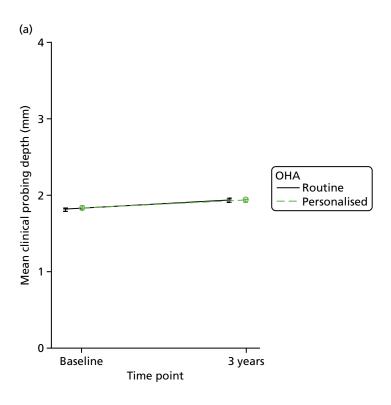


FIGURE 7 Calculus percentage (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

## Clinical probing depth

The mean clinical probing depth and 95% CI for baseline and follow-up for each randomised group are displayed in *Figure 8*. There was no evidence of changes from the baseline clinical pocket depths and no evidence of differences between any of the randomised comparisons (see *Table 13*).



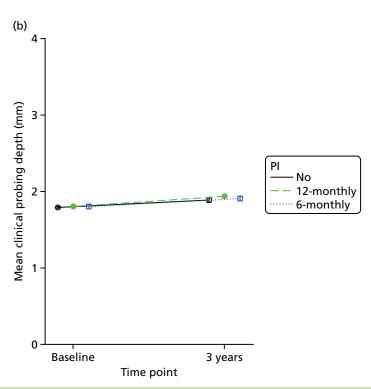


FIGURE 8 Mean clinical probing depth (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

## Patient-reported outcomes

The patient-reported outcomes (PBC, attitude and subjective norm) are shown in *Figures 9–11*. There was no evidence of any clinically important differences between the randomised groups in any of the cognitive variables at any follow-up time (see *Table 13*). Attitude was statistically significantly different between participants randomised to 6-monthly PI and those randomised to no PI, favouring a better attitude to oral hygiene behaviour in the no-PI group (difference -0.137, 95% CI -0.273 to -0.001; p = 0.048). However, the 95% CI excludes the possibility of a clinically important difference.

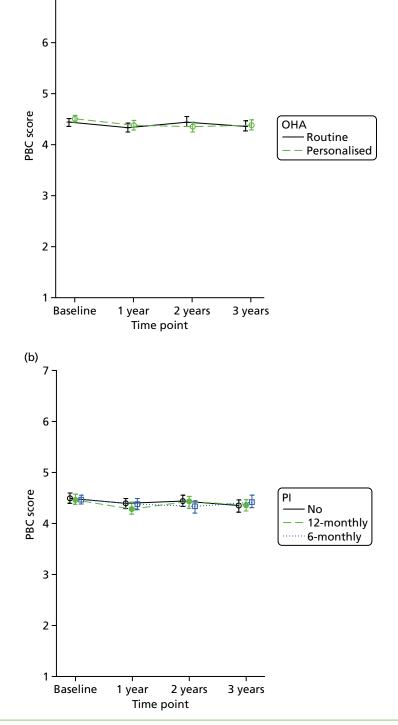
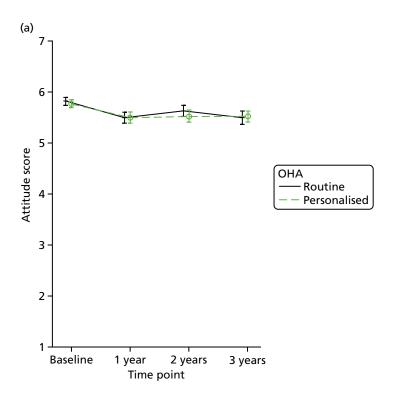


FIGURE 9 Perceived behaviour control score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.



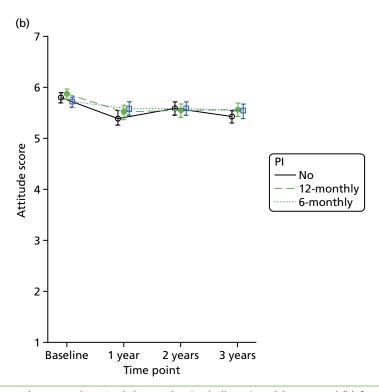
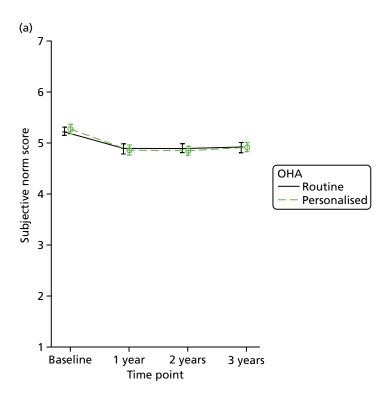


FIGURE 10 Attitude score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.



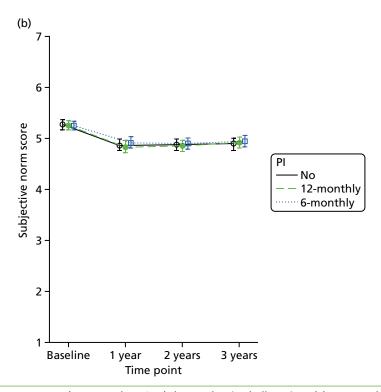


FIGURE 11 Subjective norm score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

### Behaviour score

The mean behaviour score and 95% CI for baseline and follow-up for each randomised group are displayed in *Figure 12*. The behaviour score was significantly different between participants randomised to the 6-monthly PI and those randomised to 12-monthly PI, favouring those in the 6-monthly group (difference -0.164, 95% CI -0.296 to -0.032; p = 0.015) (see *Table 13*). However, the 95% CI excludes the possibility of a clinically important difference. Therefore, there was no evidence of any clinically important differences between the randomised groups.

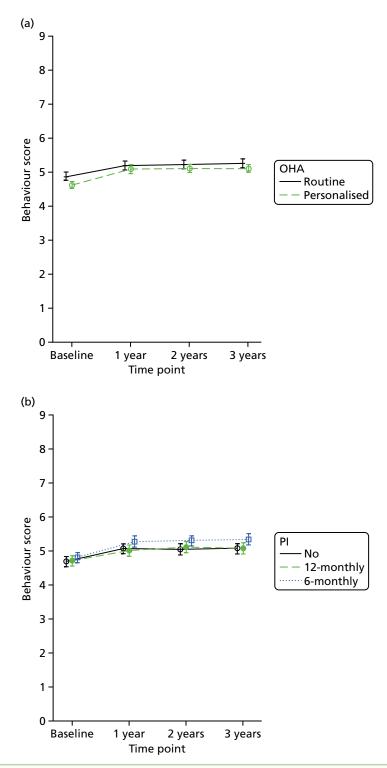
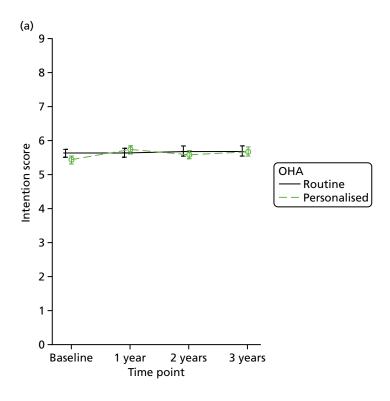


FIGURE 12 Behaviour score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

#### Intention score

The mean intention score and 95% CI for baseline and follow-up for each randomised group are displayed in *Figure 13*. There was no evidence of any clinically or statistically significant differences between the randomised groups for the intention score at any of the follow-up times (see *Table 13*).



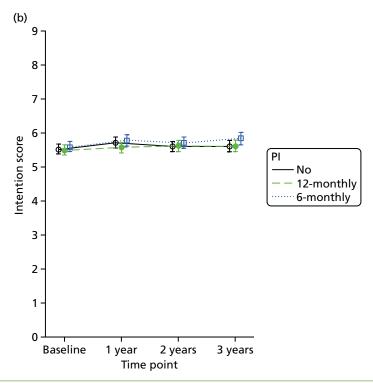
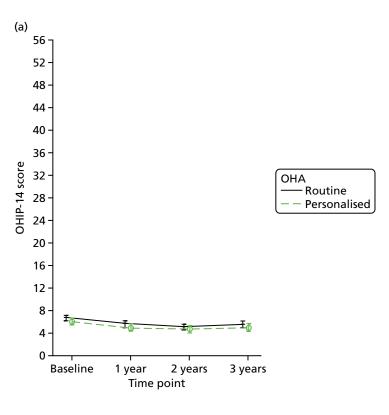


FIGURE 13 Intention score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

## Oral Health Impact Profile-14

The OHIP-14 score was overall very low (representing a good oral QoL) across the randomised groups at all follow-up time points (*Figure 14*), varying from mean 6.0 at baseline to mean 5.0 at 3 years. There was no evidence of any clinically important differences between the randomised groups for OHIP-14 scores at any of the follow-up times (see *Table 13*).



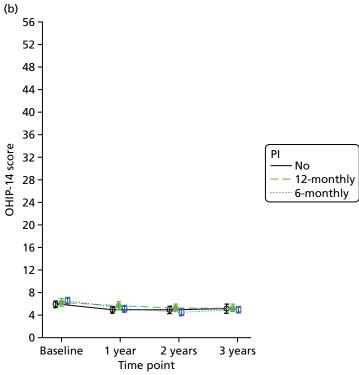


FIGURE 14 The OHIP-14 score (mean and 95% CI), by randomised allocation. (a) OHA and (b) frequency of PI.

## Private periodontal instrumentation received during the trial

There was a significant difference in the self-report of having received at least one private PI during the trial between the participants randomised to 6-monthly PI and the participants randomised to no PI. Participants in the 6-monthly PI had 50% greater odds of reporting a private PI than those in the no-PI group (odds ratio 0.5, 95% CI 0.3 to 0.7; p = 0.001). There was no evidence of a difference between the other randomised groups in private PI received during the trial (see *Table 13*). This is likely to reflect the number of PIs received in each group and a misunderstanding of the difference between a private and NHS PI rather than a real effect.

## Having a plan to brush or floss better

There was no evidence of a difference between PI frequencies or OHA randomised groups in having a plan to brush or floss better over the course of the trial (see *Table 13*). This variable was also used as a measure of fidelity (see *Fidelity of interventions*).

## Self-reported bleeding (post hoc outcome)

There was no evidence of a difference between the randomised groups for self-reported bleeding at 3 years (see *Table 13*).

## Clinician beliefs at follow-up

The majority (around 78%) of respondents to the CBQ at follow-up were dentists. Every scale regarding the provider's beliefs varied from 1 to 7, except for subjective norm, which varied from 1 to 49 (*Table 14*). Self-efficacy was high in both randomised groups (around 6 out of 7) and PBC was low. The overall attitude towards OHA and PI was around 3.5 out of 7, which was worse than at baseline, when the value was around 5 out of 7. The comparability between the follow-up and baseline CBQ is limited since the responders to both questionnaires were not always the same clinicians. Professionals did not report strong feelings of change in self-efficacy regarding both OHA and PI delivery over the course of the trial (mean of around 3 and similar between randomised groups). There was no evidence of a difference between randomised groups for any of the provider's beliefs.

TABLE 14 Clinician belief questionnaire at follow-up

	ОНА		
Clinician beliefs	Personalised (N = 44)	Routine ( <i>N</i> = 40)	Personalised OHA vs. routine OHA, estimate (95% CI; <i>p</i> -value <sup>a</sup> )
Profession, n (%)			
Hygienist	11 (25)	7 (18)	-
Dentist	33 (75)	33 (83)	-
Clinician beliefs, mean (SD), n			
Self-efficacy	6.2 (0.6), 44	6.0 (0.6), 40	0.16 (-0.12 to 0.44; 0.254)
PBC	3.0 (1.0), 44	2.8 (1.0), 40	0.15 (-0.26 to 0.57; 0.472)
Attitude	3.7 (0.5), 44	3.5 (0.7), 40	0.15 (-0.11 to 0.41; 0.270)
Subjective norm	13.2 (7.8), 43	11.3 (8.5), 40	1.87 (-1.49 to 5.22; 0.275)
Intention	5.5 (1.2), 43	5.4 (1.0), 39	0.12 (-0.39 to 0.62; 0.648)
Has the plan about advice changed throughout the trial?	3.7 (1.8), 43	3.3 (2.1), 39	0.36 (-0.39 to 1.11; 0.350)
Has the plan about providing Pl changed throughout the trial?	3.1 (1.8), 43	3.1 (1.8), 39	-0.02 (-0.77 to 0.73; 0.949)

a These analyses adjust for cluster-level minimisation variables.

## **Tertiary outcomes**

Appendix 1, Section 4: tertiary outcomes (at baseline and 3 years) provides the self-reported data on dental sensitivity and appearance for baseline and 3 years. Possible answers to all the questions varied from 1 point (not clean at all/not at all pleasant) to 7 points (could not get any cleaner/extremely pleasant). At baseline, patients reported feeling that their teeth were clean after brushing (on average, 5.8 points), but cleaner after PI (on average, 6.6 points). The same was reported for how clean their teeth look (after brushing 5.3 points vs. after a PI 6.2 points), how pleasant they feel (5.8 points after brushing vs. 6.3 points after PI) and how pleasant they look (5.1 points after brushing vs. 5.7 points after PI). Around half of the patients reported feeling sensitivity in their teeth. Dental sensitivity and appearance characteristics were balanced across randomised groups.

At 3 years, participants reported that their teeth felt and looked clean and pleasant after brushing (on average, around 5.5 points out of 7 points) and even more clean and pleasant after PI (on average, around 6.4 points out of 7 points). The average scores were similar across randomised groups, indicating that having fewer PIs throughout the trial did not influence the participants' perceptions.

## Fidelity of interventions

## Number of periodontal instrumentations received throughout the trial

Table 15 presents descriptive information from routine data and participant self-reported questionnaire data regarding the number of PIs claimed during the course of the trial. It includes participants with a clinical follow-up assessment and routine data available. The number of PIs reported in *Table 15* excluded the baseline PI given to all participants. There was a separation between the number of PIs (claimed or self-reported) in each randomised PI group: the no-PI group claimed on average one PI throughout the trial, the 12-monthly

**TABLE 15** Number of PIs received

	Randomised gr	oup, <i>n</i> (%)			
	PI			ОНА	
Number of PIs received	No ( <i>N</i> = 427)	12-monthly (N = 433)	6-monthly ( <i>N</i> = 426)	Personalised (N = 699)	Routine ( <i>N</i> = 587)
Routine data					
0	197 (46)	61 (14)	41 (10)	168 (24)	131 (22)
1	108 (25)	89 (21)	58 (14)	133 (19)	122 (21)
2	60 (14)	192 (44)	56 (13)	176 (25)	132 (22)
3	42 (10)	52 (12)	103 (24)	118 (17)	79 (13)
4	13 (3)	33 (8)	127 (30)	85 (12)	88 (15)
≥5	7 (2)	6 (1)	41 (10)	19 (3)	35 (6)
Mean (SD), n	1.0 (1.2), 427	1.8 (1.1), 433	2.8 (1.5), 426	1.8 (1.4), 699	2.0 (1.5), 587
Self-report					
0	154 (36)	62 (14)	19 (4)	122 (17)	113 (19)
1	98 (23)	88 (20)	59 (14)	142 (20)	103 (18)
2	66 (15)	94 (22)	81 (19)	128 (18)	113 (19)
3	29 (7)	77 (18)	43 (10)	74 (11)	75 (13)
4	18 (4)	42 (10)	71 (17)	74 (11)	57 (10)
≥5	14 (3)	21 (5)	105 (25)	74 (11)	66 (11)
Missing	48 (11)	49 (11)	48 (11)	85 (12)	60 (10)
Mean (SD), <i>n</i>	1.2 (1.4), 379	2.0 (1.4), 384	3.1 (1.6), 378	2.1 (1.6), 614	2.1 (1.7), 527

PI group claimed around two PIs and the 6-monthly PI group claimed three PIs. Personalised and routine OHA had a similar number of PIs throughout the trial. The self-reported number of PIs was similar to the routinely collected data; however, the similarity was due to participants over- or underestimating PIs compared with the routine data (25% of the participants self-reported the same number of PIs as those claimed in the routine data; 47% self-reported having more PIs than those claimed; and 28% self-reported having fewer PIs than those claimed). There was clear evidence of a separation in the number of PIs given as randomised.

In addition to using the routine and self-reported data, the dental practice data monitoring showed that all practices were compliant. A telephone call was made to each practice on two occasions; however, no practices required further follow-up by the trial manager.

Routine check-ups were similar across the randomised groups, with an overall mean of 3.8 visits and a SD of 1.6 (i.e. participants attended the dental practice with the same frequency across groups).

## Personalised and routine oral hygiene advice

Personalised OHA was given as intended to all participants at the baseline visit. Four questions in the participant annual questionnaire were used to further measure the continued fidelity of the OHA: (1) 'Usually, when you finish brushing your teeth do you . . .?', (2) 'What do you intend to do when you finish brushing your teeth in the future?' (related to the use of mouth wash after brushing), (3) 'Do you have a plan about when you will start brushing your teeth better?' and (4) 'Do you have a plan about when you will start flossing your teeth better?'. Correct answers to the first two questions (i.e. 'I do spit but not rinse after brushing' or 'I should spit but not rinse after brushing'), as well as having a plan about when to start brushing or flossing better, were considered as proxy indicators that the personalised OHA was delivered. At baseline, the routine and personalised groups were balanced for each question (around 28% of participants answered that they spit but do not rinse after brushing, 28% that they intended to do that and 25% that they had a plan to brush or floss better). At year 1, 29% of participants in the routine group said that they spit but do not rinse, or intended to do so, compared with around 40% in the personalised group. At the end of the trial, in the 3-year follow-up questionnaire, around 29% of the participants in the routine arm reported that they spit but do not rinse or intended to do so, compared with 39% in the personalised arm. Around 20% of the participants in either arm had a plan to brush or floss better. This was similar across the years. Participants did not necessarily need to have a plan to brush or floss better if they did not need one (one of the options in the questionnaire). There was evidence, from the answer 'I do spit but not rinse after brushing', that the personalised OHA group retained/received the OHA as intended.

## Subgroup analyses

Prespecified subgroup analyses for gingival bleeding included currently a smoker (yes/no), BPE score (of < 3 or of 3), age (< 45/45-64/>64 years) and whether or not the practice employed a hygienist (yes/no). Post hoc subgroup analyses were undertaken for participants with four or more sites with a clinical probing depth of  $\ge 4$  mm (yes/no) and region (Scotland/England). *Figures 15* and *16* show the means and 99% CIs for the differences in bleeding at 3 years in the subgroups for PI frequency and OHA, respectively. There was no evidence that any of the subgroups were statistically significantly different at the 1% level (*Table 16*).

## Sensitivity analyses

#### Missing data

Possible mechanisms of missingness were investigated by modelling baseline predictors of participants missing the final examination (see *Table 10*). Only age was identified as a predictor. Clinical severity of disease indicators (bleeding, calculus and clinical pocket depths) were not predictors. There were no differences between randomised groups in the proportion of missing data. In addition, the main reason for missing the final examination was that the participant was no longer contactable by the dental practice and, therefore, unlikely to be related to the study interventions. Given these findings, no further missing data sensitivity analyses were undertaken.

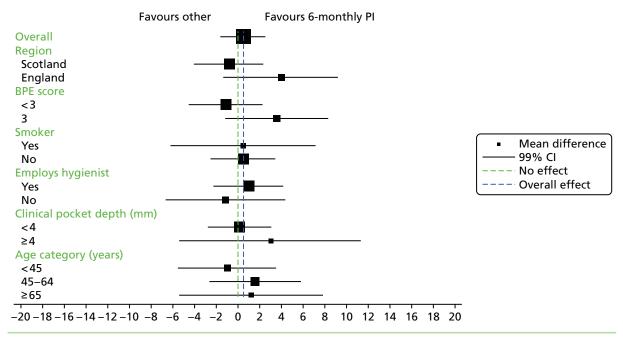


FIGURE 15 Subgroup results for PI allocation: difference between arms, by subgroup.

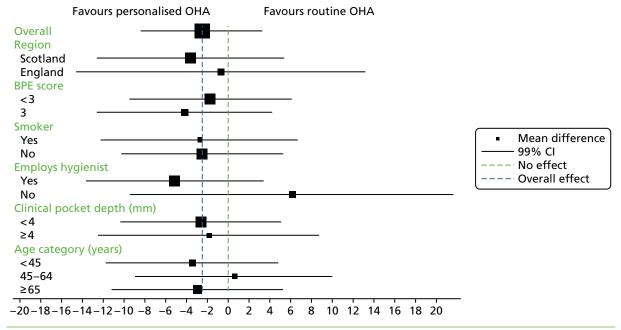


FIGURE 16 Subgroup results for OHA allocation: difference between arms, by subgroup.

## **Outliers**

To test the sensitivity of the primary outcome to outliers, we explored potential between-practice differences influencing the primary analyses. Practices where the treatment difference was twice the target difference (either –15% or +15%) and excluded zero from the 99% CI were excluded from the analysis comparing no PI with 6-monthly PI. A positive treatment difference was found in favour of the 6-monthly PI versus other PI. Three practices in England met the criteria and were excluded as part of the sensitivity analysis. They had the following treatment difference estimates +19.4; +24.2; and +24.7. After excluding the three outlier practices from the main analysis, the treatment effect of 6-monthly PI versus no PI was –0.38 (95% CI –2.9 to 2.1), of 6-monthly PI versus 12-monthly PI was –0.18 (95% CI –2.7 to 2.3) and of

TABLE 16 Interaction mean for 6-monthly PI vs. other PI and personalised OHA vs. routine OHA, by subgroup

	Mean (99% Cl; <i>p</i> -value)	
Subgroups	Other PI vs. 6-monthly PI	Personalised OHA vs. routine OHA
Scotland vs. England	-4.8 (-11.0 to 1.4; 0.048)	-2.9 (-19.5 to 13.6; 0.649)
BPE score of 3 vs. < 3	4.7 (-1.1 to 10.5; 0.038)	-2.5 (-8.1 to 3.1; 0.254)
Smoker vs. non-smoker	0.0 (-7.3 to 7.4; 0.992)	-0.3 (-7.2 to 6.7; 0.922)
Employs hygienist vs. does not	2.1 (-4.3 to 8.5; 0.396)	-11.3 (-29.0 to 6.5; 0.102)
Clinical pocket depth (mm) $\geq$ 4 vs. < 4	2.8 (-6.1 to 11.6; 0.417)	0.8 (-7.7 to 9.2; 0.811)
Age (years) 45–64 vs. < 45	2.5 (-3.7 to 8.8; 0.291)	0.4 (-5.5 to 6.3; 0.855)
Age (years) > 65 vs. < 45	2.2 (-5.9 to 10.2; 0.487)	3.9 (-3.6 to 11.4; 0.177)

personalised OHA versus routine OHA was -2.1 (95% CI -8.1 to 4.0). The overall  $I^2$  for the practice effects in the trial was 27%. When the outliers were removed, the  $I^2$  was 0% and the interaction mean in the subgroup analysis for region changed from -4.8 (95% CI -11 to 1.4; p = 0.048) to -2.3 (95% CI -9.0 to 4.5; p = 0.387).

# **Chapter 5** Results of economic analysis

#### Resource use and costs

#### NHS dental care resource use

Data were linked for 1337 out of 1346 (99%) trial participants randomised in Scotland (after post-randomisation exclusion) and 477 out of 525 (91%) trial participants randomised in England. It was not possible to link data for 11 participants in Scotland, but data were available for two participants randomised in England having some dental treatment in Scotland. Linkage was not possible for 48 participants in England. A further participant had claims in both Scotland and England. For this individual, claims were summed across both data sets, with the participant remaining allocated to their region of randomisation for the purposes of reporting results.

Table 17 describes the total number of treatment claims provided to all randomised participants with linked data over the 2.25-year follow-up period (excluding baseline and final study visits) by randomised group. The majority of claims in England fell into band 1 treatments. There were relatively few emergency claims and free treatments across all groups. The highest number of claims in Scotland was for diagnostic assessments, including clinical examinations and radiographs. There were 2516 periodontal treatments provided, accounting for 23% of claims. Among the remaining categories, the greatest number of claims were for conservative treatments [2506 claims (23%)] such as fillings, with relatively few surgical extractions or more intensive treatments. There were clear differences across groups in terms of numbers of periodontal treatments reflecting the different number of PIs delivered in each randomised trial arm. Overall, owing to the restrictions of the reimbursement system, there were only 11 claims for 'preventative care' (including intensive hygiene instruction), all in the personalised OHA cluster. Detailed information on items claimed in Scotland can be found in *Appendix 2, Section 3: detailed claims for Scottish routine data*.

Table 18 shows that the numbers of PIs delivered to all randomised participants with linked data were evenly balanced across clusters. The mean number of reported PIs was lower in England than in Scotland. It should be noted that the number of PIs in England may be under-reported as only one qualifying procedure for a band 1 treatment needs to be reported when filling in the clinical data set (e.g. a clinical examination).

#### NHS dental care costs

Table 19 presents complete-case NHS dental costs for the UK and at the regional level based on the appropriate region-level unit costs applied to resource use data. Incremental differences are presented for each randomised policy, compared with standard care.

The mean cost per trial participant over the 2.25-year follow-up period (excluding baseline and final study visits) ranged from £62.42 to £85.43 across the randomised groups. The SDs are generally large, suggesting relatively large variation in costs across participants. Data were somewhat skewed to the left, with some outliers incurring very high costs; however, these were spread evenly across groups.

The average cost is lowest for the no-PI groups. However, none of the differences in mean cost between standard care (routine OHA and 6-monthly PI) and the other policies considered was statistically significant at a 5% level. The largest difference in costs is between standard care and a policy of no PI with personalised OHA (mean difference –£14.91; 95% CI –£34.18 to £4.36).

Table 19 also reports the NHS costs by region. This is important given the differences in reimbursement systems and the level of detail on resource use. Average per-participant costs are higher in England than in Scotland. However, the overall results are the same across the two regions: there are no statistically

**TABLE 17** Number of treatment claims

	OHA, n								
	Routine				Personalise	ed			
Region	No PI	12-monthly PI	6-monthly PI	Total	No PI	12-monthly PI	6-monthly PI	Total	Overall total, n
Scotland	(N = 192)	(N = 193)	(N = 195)	(N = 580)	(N = 250)	(N = 255)	(N = 252)	(N = 757)	(N = 1337)
Category of claim <sup>a</sup>									
Diagnosis	679	666	698	2043	873	894	912	2679	4722
Preventative care	0	0	0	0	7	4	0	11	11
Periodontal	268	343	530	1141	251	466	658	1375	2516
Conservative treatments	370	351	361	1082	462	495	467	1424	2506
Surgical treatments	59	86	62	207	83	98	69	250	457
Prostheses	28	36	19	83	39	63	54	156	239
Orthodontic	1	4	0	5	0	0	0	0	5
Other	53	55	72	180	67	110	85	262	442
Total	1458	1541	1742	4741	1782	2130	2245	6157	10,898
England	(N = 79)	(N = 80)	(N = 88)	(N = 247)	(N = 77)	(N = 77)	(N = 76)	(N = 230)	(N = 477)
Band 1 <sup>b</sup>	156	173	204	533	118	161	156	435	968
Band 2	74	75	62	211	52	66	63	181	392
Band 3	15	18	12	45	8	17	13	38	83
Band urgent	24	20	27	71	24	24	20	68	139
Free	1	1	0	2	0	6	0	6	8
Total	270	287	305	862	202	274	252	728	1590

SDR, Statement of Dental Remuneration (Scotland).

a Categories are taken from the respective sections of SDR. Examples include: diagnosis (e.g. examinations/films); preventative (e.g. intensive instruction preventative surface applications); periodontal (e.g. PI); conservative treatments (e.g. fillings/restorations/inlays/crowns); surgical treatments (e.g. extractions); prostheses (e.g. dentures, including fitting, repair and amendments); orthodontic treatments (including construction, repair); other (e.g. stoning or smoothing a tooth surface/treatment of sensitive cementum or dentine/urgent treatment/prescriptions).

b Differences in UDAs across groups would not pick up differences in the number of PIs delivered that are likely to get masked by the banding system, which assigns the same band to a check-up or a check-up plus PI.

TABLE 18 Number of participants having Pla in the routine data set<sup>b</sup>

	OHA, n												
	Routine				Personalised	ŀ							
Region	No PI	12-monthly PI	6-monthly PI	Total	No Pl	12-monthly PI	6-monthly PI	Total	Overall total, <i>n</i>				
Scotland	(N = 192)	(N = 193)	(N = 195)	(N = 580)	(N = 250)	(N = 255)	(N = 252)	(N = 757)	(N = 1337)				
0	68	29	28	125	109	35	21	165	290				
1	50	49	24	123	70	54	45	169	292				
2	28	74	31	133	40	111	40	191	324				
3	30	23	32	85	24	37	66	127	212				
4	11	14	63	88	6	14	70	90	178				
≥ 5°	5	4	17	26	1	4	10	15	41				
Mean (SD)	1.4 (1.4)	1.8 (1.2)	2.7 (1.7)	2.0 (1.6)	1.0 (1.1)	1.8 (1.2)	2.6 (1.4)	1.8 (1.4)	1.9 (1.5)				
England	(N = 79)	(N = 80)	(N = 88)	(N = 247)	(N = 77)	(N = 77)	(N = 76)	(N = 230)	(N = 477)				
0	40	20	25	85	44	27	20	91	176				
1	28	16	18	62	16	15	17	48	110				
2	7	35	14	56	9	22	12	43	99				
3	1	7	10	18	7	4	13	24	42				
4	1	2	9	12	0	8	8	16	28				
≥ 5°	2	0	12	14	1	1	6	8	22				
Mean (SD)	0.8 (1.1)	1.4 (1.0)	2.0 (1.9)	1.4 (1.5)	0.8 (1.2)	1.4 (1.4)	1.9 (1.6)	1.4 (1.5)	1.4 (1.5)				

DOI: 10.3310/hta22380

HEALTH TECHNOLOGY ASSESSMENT 2018 VOL. 22 NO. 38

SDR, Statement of Dental Remuneration (Scotland).

a 'PI' refers to any claim for PI including simple PI (SDR item 10A); treatment of periodontal disease, including PI (SDR item 10B); and more intensive treatment over 3 visits, including deep scaling (SDR item 10C)

b It should be noted that the number of PIs reported here refer to all participants with linked data, whereas Table 15 reports the number of PIs for participants attending the final clinic visit. As such, these numbers should not be directly compared.

c The maximum number of PIs provided to any individual participant over the time frame (90 days to 2.5 years post baseline clinic examination) was nine.

TABLE 19 NHS costs (f) of dental care

	OHA, r	n														
	Routin	e							Person	alised						
	No PI		12-mont	thly PI	6-mont	hly PI	Total		No PI		12-monthly Pl		6-monthly PI		Total	
Region	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
UK	(N = 27	1)	(N = 273)	)	(N = 28	3)	(N = 82	77)	(N = 32	? <b>7</b> )	(N = 332	2)	(N = 328	8)	(N = 987	7)
Total	73.23	118.04	81.37	141.64	73.75	104.78	76.10	122.16	62.42	116.13	85.43	131.62	82.84	134.19	76.94	127.87
Mean (95% CI) cost difference vs. routine OHA with 6-monthly PI	-3.12 (-18.18	s to 11.93)	0.14 (-15.42 t	to 15.69)	-		N/A		-14.91 (-34.18	3 to -4.36)	11.67 (–11.00	to 34.33)	6.23 (–15.70	to 28.16)	N/A	
Scotland	(N = 19	2)	(N = 193)	)	(N = 19	5)	(N = 58	80)	(N = 25	50)	(N = 25	5)	(N = 252	2)	(N = 757	7)
Diagnosis	24.27	10.43	24.30	11.08	24.88	12.36	24.49	11.31	23.24	10.83	23.93	11.90	25.46	11.10	24.21	11.31
Prevention	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.20	1.54	0.17	2.35	0.00	0.00	0.12	1.63
Periodontal	7.72	12.70	9.90	12.76	14.20	16.73	10.63	14.43	5.65	11.03	10.27	13.45	11.53	12.06	9.16	12.47
Conservative	23.40	55.90	22.66	46.77	25.15	61.25	23.74	54.90	25.54	77.32	33.53	86.20	32.02	97.44	30.39	87.35
Surgical	1.06	3.19	2.01	6.36	1.99	11.64	1.69	7.90	1.53	6.34	1.73	7.58	1.33	4.90	1.53	6.37
Prosthesis	5.04	34.19	7.34	37.11	4.87	32.32	5.75	34.55	3.49	23.70	7.65	33.28	7.09	32.13	6.09	30.05
Orthodontic	1.22	16.88	0.44	6.07	0.00	0.00	0.55	10.32	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Other	0.95	3.30	0.68	2.24	1.02	3.33	0.88	3.00	1.58	14.61	1.49	6.32	1.38	5.95	1.49	9.77
Total	63.68	86.38	67.32	82.51	72.12	93.08	67.73	87.37	61.24	102.71	78.76	120.40	78.82	119.90	72.99	114.83
Mean (95% CI) cost difference vs. routine OHA with 6-monthly PI	-9.11 (-23.62	to 5.41)	-6.60 (-21.39 t	to 8.20)	-		N/A		-13.54 (-33.83	3 to 6.76)	5.45 (–17.37	to 28.28)	4.16 (–18.46	to 26.75)	N/A	

DOI: 10.3310/hta22380

	OHA, n	,														
	Routine	e							Person	alised						
	No Pl		12-monthly PI		6-mont	hly Pl	Total	Total		No Pl		thly Pl	6-monthly PI		Total	
Region	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
England	(N = 79)	)	(N = 80)		(N = 88)		(N = 24	17)	(N = 77	)	(N = 77)		(N = 76)		(N = 23	0)
Band 1	19.65	24.90	21.65	22.88	20.40	26.63	20.56	24.82	13.73	£17.37	22.41	24.49	20.11	19.73	18.74	20.98
Band 2	32.31	59.37	42.38	113.02	26.49	46.21	33.50	77.58	23.77	£41.97	36.37	78.65	35.51	57.62	31.87	61.31
Band 3	38.63	130.37	46.57	133.00	25.08	97.55	36.38	120.48	23.39	£111.83	39.70	114.59	35.96	123.45	33.00	116.40
Urgent	5.49	21.48	4.31	11.24	5.42	13.31	5.08	15.81	5.38	£15.13	6.10	15.23	4.58	10.82	5.35	13.84
Free	0.35	3.15	0.36	3.24	0.00	0.00	0.23	2.56	0.00	£0.00	1.84	9.61	0.00	0.00	0.62	5.61
Total	96.44	170.85	115.27	225.57	77.39	127.47	95.75	177.72	66.27	£152.54	106.41	162.10	96.16	173.67	89.59	163.14
Mean (95% CI) cost difference vs. routine OHA with 6-monthly PI	23.24 (–22.22	to 68.70)	22.09 (–23.63 t	o 67.80)	-		N/A		-17.67 (-62.30	) to 26.95)	33.73 (–27.62	to 95.08)	13.87 (–40.83	to 68.60)	N/A	

N/A, not applicable.

#### Note

NHS costs are based on the amount paid to the dentist for each participant, accounting for all exemptions. Cost model analysed using mixed-effects generalised linear model, gamma family, link log with a random effect for centre. Models adjusted for cluster- and individual-level minimisation covariates, age, age<sup>2</sup> and sex.

significant differences in costs between standard care and the other five groups. The CIs for incremental costs are generally wider for England, which is not surprising given the smaller sample sizes.

The Scottish data provide much more detail on resource use than the English data and can, therefore, provide additional insights into differences in different cost categories. The costs of periodontal treatments were expected to be lower for no PI and 12-monthly PI than for standard care, and this was found to be the case. Mean cost differences of periodontal treatments in Scotland for each randomised group, compared with routine OHA and 6-monthly PI were as follows: routine OHA, no PI, –£7.47 (95% CI –£11.22 to –£3.72); routine OHA, 12-monthly PI, –£5.11 (95% CI –£8.96 to –£1.26); personalised OHA, no PI, –£9.31 (95% CI –£13.30 to –£5.32); personalised OHA, 6-monthly PI –£2.00 (95% CI –£6.76 to £2.75); and personalised OHA, 12-monthly PI, –£3.78 (95% CI –£8.29 to £0.73). There were no significant differences across groups in any of the other categories of cost. Given the relatively low average NHS costs of PI, the differences in PI costs did not lead to differences in the total costs.

Participant exemptions from NHS co-charges can have an impact on the split of dental charges between NHS and participant. Therefore, it is important to explore whether or not there are differences in the proportion of participants exempt across the randomised groups. *Table 20* shows that around 24% of participants were exempt from charges. There were no significant differences across randomised groups in terms of the proportion of participants exempt from payment ( $\chi^2$  6.82; p-value of 0.234). As a robustness check, we also analysed the data assuming that all participants are exempt, that is, the NHS pays the full cost of dental care provided. Relative to standard care, for this exploratory analysis, incremental costs were as follows: routine OHA, no PI, -£4.76 (95% CI -£30.53 to £21.01); routine OHA, 12-monthly PI, -£1.89 (95% CI -£28.04 to £24.27); personalised OHA, no PI, -£25.65 (95% CI -£53.15 to £1.85); personalised OHA, 12-monthly PI, £10.95 (95 CI -£20.17 to £42.06); and personalised OHA, 6-monthly PI, £11.21 (95% CI -£19.97 to £42.40). Furthermore, using similar regression models as for the main results, there was no evidence that exemption status had an impact on the number of PIs delivered (p = 0.518). The main conclusion was the same: there was no statistically significant difference in costs across the randomised groups.

#### Other NHS-incurred costs

The analysis protocol set out to measure costs to the NHS of primary and secondary care, based on annual questionnaire data, for problems related to participants' teeth. There were 193 out of 1630 (12%) participants who reported at least one contact with secondary care services over the 3 years of follow-up. This relatively high number of participants reporting secondary care use was unexpected given the relatively good dental health of the trial population. A validation exercise was conducted by comparing secondary care contacts from the trial with the general population using routine data and by checking self-report data against practice records. In terms of the routine data, the average annual rate of outpatient consultations in the trial (1:19) was six times greater than the corresponding number of consultations in the general population in Scotland (1:121) and three times greater than the corresponding rate in England (1:70). A similar result was found when inpatient admissions were compared, with an average annual rate among trial participants (1:70) 10 times higher than the general population in Scotland (1:722) and over three times higher than the general population rate in England (1: 249). It should be noted that the general population rates for England and Scotland should not be directly compared as the data are based on different assumptions. For 80 out of the 193 trial participants reporting resource use (23 in England and 57 in Scotland), data were checked against dental practice records for evidence of referral for secondary care treatment (across all hospitals, dental or otherwise). Fewer than one in four reports were validated against practice records. Table 21 summarises the results of the validation exercise.

Given the poor level of validation of other NHS resource use against routine data, the lack of evidence of dental care referrals in practice records and the fact that the general population may be expected to have, on average, poorer rather than better dental health, the data on use of NHS resources have been excluded from the base-case analysis. *Appendix 2, Section 4: participant-reported contact with non-dental health services* presents descriptive data for primary and secondary care resource use falling outside the dental budget for information only. The data are excluded from all subsequently reported results.

DOI: 10.3310/hta22380

**TABLE 20** Exemptions from NHS dental charges

	Proportion	OHA, n (%	<b>6)</b>							
	charge to	Routine				Personalis	ed			Overall
Region	participant (%)	No Pl	12-monthly Pl	6-monthly PI	Total	No PI	12-monthly PI	6-monthly PI	Total	total
UK		(N = 271)	(N = 273)	(N = 283)	(N = 827)	(N = 327)	(N = 332)	(N = 328)	(N = 987)	(N = 1814)
Exempt	Region-specific	62 (23)	68 (25)	81 (29)	211 (26)	71 (22)	88 (27)	70 (21)	229 (23)	440 (24)
Non-exempt		209 (77)	205 (75)	202 (71)	616 (74)	256 (78)	244 (73)	258 (79)	758 (77)	1374 (76)
Scotland		(N = 192)	(N = 193)	(N = 195)	(N = 580)	(N = 250)	(N = 255)	(N = 252)	(N = 757)	(N = 1337)
Employment support allowance	0	4 (2)	1 (1)	4 (2)	9 (2)	4 (2)	3 (1)	5 (2)	12 (2)	21 (2)
Pay full patient charge	100	152 (79)	141 (73)	134 (69)	427 (74)	193 (77)	186 (73)	201 (80)	580 (77)	1007 (75)
Help with health care	Various	0 (0)	2 (1)	3 (2)	5 (1)	2 (1)	3 (1)	2 (1)	7 (1)	12 (1)
Income support	0	3 (2)	11 (6)	7 (4)	21 (4)	4 (2)	7 (3)	10 (4)	21 (3)	42 (3)
Jobseeker's allowance	0	2 (1)	8 (4)	8 (4)	18 (3)	8 (3)	8 (3)	6 (2)	22 (3)	40 (3)
Maternity/newborn	0	7 (4)	3 (2)	3 (2)	13 (2)	7 (3)	9 (4)	5 (2)	21 (3)	34 (3)
Pension/tax credits	0	24 (13)	27 (14)	36 (18)	87 (15)	32 (13)	39 (15)	23 (9)	94 (12)	181 (14)
England		(N = 79)	(N = 80)	(N = 88)	(N = 247)	(N = 77)	(N = 77)	(N = 76)	(N = 230)	(N = 477)
Child	0	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	1 (1)	1 (1)	2 (1)	2 (0)
Exempt	0	22 (28)	16 (20)	20 (23)	58 (23)	14 (18)	18 (23)	18 (24)	50 (22)	108 (23)
Not exempt	According to band	57 (72)	64 (80)	68 (77)	189 (77)	63 (82)	58 (75)	57 (75)	178 (77)	367 (77)

a Note that exemptions are based on the most commonly reported exemption for each individual participant across the follow-up period. Initially, linked data in Scotland had a small number of claims detailing exemptions for children. However, after checking with the trial database and identifying all participants as aged ≥ 18 years, these were superseded with the full payment of charges for the participant.

TABLE 21 Validation of the rates of other NHS secondary care resource use

Validation against routine ISD data	Trial data (UK)ª	Trial data (England)ª	General population rate (England) <sup>b,c</sup>	Trial data (Scotland)ª	General population rate (Scotland) <sup>c</sup>
Outpatients					
Year 1	1 : 18	1:19	1:70 <sup>b</sup>	1:15	1:121 <sup>d</sup>
Year 2	1 : 18	1:18		1:17	
Year 3	1:23	1:22		1:26	
Average annual rate	1:19	1:20		1:19	
Inpatients					
Year 1	1:40	1:41	1:249 <sup>b</sup>	1:39	1:722 <sup>e</sup>
Year 2	1:42	1:46		1:35	
Year 3	1:129	1:164		1:85	
Average annual rate	1:70	1:84		1:53	
Validation against practice records			% with a referral in practice notes		% with a referral in practice notes
One or more contacts with secondary care over 3 years <sup>f</sup>	1:10	1:11	3:23 (13)	1:10	16:57 (28)

- a All trial data conservatively refer to 'at least one consultation' over 1 year (many will have had more than one).
- b Data for general population in England were sourced from Hospital Episode Statistics data<sup>62,63</sup> for outpatient and inpatient consultations. Data conservatively assumes that all oral surgery procedures were related to dental care for the purposes of the validation exercise. Note that general population routine data should not be compared across regions as assumptions are unlikely to be directly comparable.
- c An estimate of the adult (aged ≥ 18 years) populations of 4.5 million (Scotland) and 43.1 million (England) were used to calculate general population rates.<sup>64</sup>
- d Validation from ISD records (Scotland, 2015 data)<sup>44</sup> for outpatient reports is based on nurse-led clinics (ISD R045; n = 2450 consultations), allied health professionals (ISD R046; n = 9997 consultations) and consultant-led outpatients (ISD R044; n = 24,606 patients). There is likely an overestimate as it assumes each new nurse-led clinic and allied health professionals visit is for a new patient.
- e Validation from ISD records (Scotland, 2015 data)<sup>44</sup> for inpatient reports based on inpatient admissions (ISD R040; n = 129 admissions) and day cases (ISD R042; n = 6104 admissions). There is likely an overestimate as it assumes each admission was for a new patient.
- f Validation against practice notes applies to referrals from the dental practice and any notes the practice may have of a patient's contact with secondary care services for dental-related problems. There may be rare cases in which the practice was not aware of a patient's attendance for secondary care services.

## Participant-incurred costs

### Participant charges for NHS dental care

Table 22 reports complete-case data on participant-incurred charges for NHS dental treatments, provided to all randomised participants with linked routine data. The average charge to participants over the 2.25-year follow-up period ranged from £48.95 to £61.33.

## Opportunity costs of time and travel to dental appointments

Appendix 2, Section 5: unit opportunity cost of time and travel to dental appointments reports the descriptive data used to calculate the unit opportunity costs of time and travel for trial participants attending NHS dental appointments. Overall, 1308 out of 1873 (70%) participants completed the appropriate baseline questions and had linked routine data on the number of visits to the dentist to enable a full calculation of per-participant unit opportunity costs of attending dental appointments (for both participants and any companions they took with them to the dentist). The average opportunity cost of time and travel for a single dental visit ranged from £9.85 to £11.07 across groups. The average companion cost was much lower,

TABLE 22 Participant-incurred co-charges (£) for dental care<sup>a</sup>

	OHA, co	o-charges	(£)													
	Routine								Persona	ilised						
	No PI		12-mon	thly PI	6-mont	hly Pl	Total		No PI		12-mon	thly PI	6-mont	nly Pl	Total	
Region	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
UK	(N = 271	1)	(N = 273	3)	(N = 283	3)	(N = 827	7)	(N = 327	7)	(N = 332	?)	(N = 328	3)	(N = 987	7)
Total	59.96	80.16	53.94	68.50	60.97	70.65	58.32	73.21	48.95	67.12	60.64	99.87	61.33	77.92	56.99	82.95
Scotland <sup>b,c</sup>	(N = 192	?)	(N = 193	3)	(N = 195	5)	(N = 580	))	(N = 250	))	(N = 255	5)	(N = 252	)	(N = 757	7)
Diagnosis	3.15	4.69	2.35	3.25	2.73	3.54	2.74	3.88	3.47	4.31	3.34	6.09	2.95	4.07	3.25	4.91
Prevention	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.06	0.66	0.09	0.79	0.00	0.00	0.05	0.59
Periodontal	11.78	15.34	13.95	13.58	22.08	20.37	15.97	17.25	8.27	11.42	14.54	13.38	23.38	18.71	15.41	16.05
Conservative	30.41	55.62	23.07	38.94	26.23	50.89	26.56	48.99	30.27	59.32	31.92	76.69	25.65	55.99	29.29	64.67
Surgical	1.65	5.51	1.76	6.31	1.42	4.48	1.61	5.48	1.85	7.72	1.48	4.47	1.14	4.68	1.49	5.80
Prosthesis	4.33	22.77	1.19	9.70	1.06	8.47	2.19	15.12	3.32	18.41	5.55	26.21	5.14	27.99	4.68	24.57
Orthodontic	0.00	0.00	1.75	24.28	0.00	0.00	0.58	14.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Other	0.57	2.34	1.00	2.78	0.76	2.51	0.77	2.55	0.70	2.00	0.98	2.79	1.19	4.53	0.96	3.29
Total	51.88	76.16	45.07	55.86	54.27	64.49	50.42	66.01	47.94	71.22	57.89	102.78	59.46	77.36	55.13	85.05
England	(N = 79)		(N = 80)		(N = 88)		(N = 247	7)	(N = 77)		(N = 77)		(N = 76)		(N = 230	))
Treatment band 1	27.44	28.89	30.02	29.07	33.93	30.79	30.59	29.64	22.32	23.63	26.82	24.76	27.82	28.00	25.64	25.52
Treatment band 2	33.06	48.67	24.30	42.46	24.10	37.27	27.03	42.88	23.85	42.42	23.24	39.64	23.03	42.33	23.37	41.30
Treatment band 3	16.17	56.77	18.16	67.66	14.35	53.36	16.16	59.17	2.84	24.96	16.40	66.36	13.64	61.89	10.95	54.38
Urgent	2.93	9.67	2.87	8.15	3.44	9.34	3.09	9.05	3.20	6.82	2.52	7.96	3.05	9.35	2.92	8.08
Free	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Total	79.61	86.53	75.34	88.97	75.82	81.12	76.88	85.13	52.20	51.85	68.97	89.20	67.54	79.97	62.88	75.40

SDR, Statement of Dental Remuneration (Scotland).

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

a Participant charges based on each individual's exemption status.

b All costs are inflated to 2015 (SDR 130) values.

c No participants reached maximum treatment course charge (£384).

TABLE 23 Participant-incurred costs (f)

	OHA, cost	t (£)																
	Routine								Personalise	ed								
	No PI		12-month	ly Pl	6-mont	hly Pl	Total		No PI		12-month	ly PI	6-monthly	/ PI	Total			
Region	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD		
UK	(N = 289)		(N = 287)		(N = 290	))	(N = 86	6)	(N = 334)		(N = 338)		(N = 335)		(N = 100	) <i>7</i> )		
Private Pl	5.75	20.59	5.67	23.10	12.04	34.56	7.83	26.94	4.23	20.88	7.71	26.23	9.18	24.51	7.04	24.06		
Other private dental care	33.26	142.49	31.58	94.19	30.22	109.12	31.68	116.98	20.54	90.63	18.54	80.00	23.78	87.21	20.94	86.01		
Electric toothbrushes	19.24	38.94	18.00	34.67	25.04	44.36	20.77	39.63	17.91	31.86	23.20	52.44	15.91	31.47	19.02	39.95		
Manual toothbrushes	20.59	12.46	19.65	12.51	18.83	12.61	19.69	12.54	18.46	12.47	19.14	12.95	19.10	12.78	18.90	12.73		
Interdental brushes	101.28	191.70	120.06	202.15	129.81	208.14	117.06	200.95	101.17	194.15	104.98	189.31	120.49	197.30	108.88	193.62		
Electric toothbrush (heads)	19.40	21.77	19.95	21.94	21.27	20.92	20.21	21.55	18.74	20.72	21.05	20.98	19.00	22.86	19.60	21.55		
Participant NHS charges	60.30	80.75	54.72	69.17	61.44	71.10	58.83	73.84	49.00	67.19	60.49	99.50	61.20	77.76	56.92	82.76		
Time and travel costs	43.55	51.33	47.31	57.12	46.44	48.35	45.77	52.36	34.57	36.17	42.75	56.46	44.36	50.95	40.57	48.81		
Total participant costs	303.38	281.03	316.95	267.39	345.10	301.61	321.85	284.02	264.62	249.57	297.85	271.66	313.02	263.00	291.87	262.16		
Mean (95% CI) cost difference vs. routine OHA with 6-monthly Pl <sup>a</sup>	-39.80 (-83.94 to	4.33)	-23.48 (-70.00 to	23.04)	-		N/A		-64.11 (-112.33 to	-15.88)	-16.64 (-68.28 to	35.00)	-30.28 (-81.59 to	21.03)	N/A			
Scotland	(N = 196)		(N = 195)		(N = 195	5)	(N = 58	6)	(N = 252)		(N = 255)		(N = 252)		(N = 75	9)		
Private Pl	4.62	13.89	4.92	22.18	5.89	18.76	5.14	18.57	3.01	14.92	6.11	21.41	7.24	18.71	5.45	18.63		
Other private dental care	30.41	135.64	30.89	95.63	22.06	78.27	27.79	105.95	16.03	60.77	15.17	69.50	22.72	87.20	17.96	73.32		
Electric toothbrushes	15.96	30.60	16.96	34.88	25.30	42.44	19.40	36.49	18.01	32.95	19.03	35.54	15.40	29.86	17.49	32.88		
Manual toothbrushes	20.23	12.04	20.18	12.15	19.02	12.55	19.81	12.24	18.87	12.24	18.77	12.74	19.62	12.76	19.08	12.58		
Interdental brushes	79.08	170.38	97.77	182.56	107.58	192.33	94.78	182.11	99.52	193.14	93.75	177.23	107.55	185.92	100.25	185.41		
Electric toothbrush (heads)	18.01	21.53	17.26	20.62	20.99	20.70	18.75	21.00	17.81	20.24	20.44	20.85	17.98	21.98	18.75	21.05		
Participant NHS charges	51.75	75.85	45.07	55.80	54.51	64.53	50.45	65.93	47.91	71.13	58.12	102.86	59.46	77.36	55.17	85.02		
Time and travel costs	39.86	44.44	46.53	56.87	43.28	45.98	43.22	49.41	38.16	37.63	45.27	60.37	48.16	54.86	43.87	52.02		

DOI: 10.3310/hta22380

	OHA, cos	st (£)														
	Routine								Personalis	ed						
	No Pl		12-monthly PI		6-monthly PI		Total		No Pl		12-monthly PI		6-monthly PI		Total	
Region	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Total participant cost	259.92	249.03	279.58	246.81	298.64	245.20	279.34	247.21	259.32	240.28	276.66	259.72	298.11	252.42	278.03	251.19
Mean (95% CI) cost difference vs. routine OHA with 6-monthly PI <sup>a</sup>	-44.10 (-93.67 to	5.47)	-24.78 (-77.38 to	27.81)	-		N/A		-41.13 (-93.09 to	10.82)	–27.65 (–80.89 to	25.60)	−3.78 (−59.28 to	51.72)	N/A	
England	(N = 93)		(N = 92)		(N = 95)		(N = 28	0)	(N = 82)		(N = 83)		(N = 83)		(N = 248	8)
Private PI	8.14	30.15	7.28	24.96	24.68	51.98	13.47	38.47	8.00	32.90	12.62	37.05	15.06	36.41	11.91	35.51
Other private care	39.27	156.39	33.04	91.42	46.98	153.28	39.84	137.02	34.41	148.49	28.88	105.68	26.99	87.56	30.08	116.25
Electric toothbrushes	26.17	51.84	20.19	34.28	24.48	48.22	23.63	45.41	17.57	28.38	36.00	84.64	17.45	36.04	23.70	56.15
Manual toothbrushes	21.35	13.32	18.54	13.22	18.44	12.76	19.44	13.14	17.20	13.13	20.25	13.58	17.53	12.77	18.33	13.20
Interdental brushes	148.08	224.00	167.30	232.34	175.44	231.56	163.68	228.94	106.23	198.18	139.46	219.77	159.80	224.89	135.28	215.02
Electric toothbrush (heads)	22.32	22.05	25.65	23.59	21.85	21.41	23.25	22.37	21.59	21.98	22.92	21.36	22.12	25.20	22.21	22.85
Participant NHS charges	78.32	87.89	75.17	88.16	75.68	81.50	76.39	85.61	52.37	53.53	67.79	88.56	66.51	79.17	62.26	75.34
Time and travel costs	51.34	62.90	48.98	57.86	52.93	52.46	51.10	57.75	23.53	28.69	35.01	41.47	32.83	34.26	30.49	35.48
Total participant costs	394.98	321.23	396.15	292.16	440.48	376.48	410.80	331.94	280.89	277.00	362.93	297.55	358.29	289.51	334.25	289.59
Mean (95% CI) cost difference vs. routine OHA with 6-monthly Pl <sup>b</sup>	-25.41 (-113.77	to 62.95)	-14.18 (-109.65	to 81.29)	-		N/A		–125.59 (–219.85 to	-31.33)	-20.18 (-132.27	to 91.91)	-43.81 (-148.79 t	to 61.16)	N/A	

MEGLM, mixed-effects generalised linear model; N/A, not applicable.

a MEGLM analysis, gamma distribution, log link, with random effect for cluster and adjustment for cluster- and individual-level covariates, age and sex.

b MEGLM with gamma distribution, log link did not converge for the English data. Therefore, a mixed model with assumed normal distribution was used for English data with random effect for cluster and adjustment for cluster- and individual-level covariates, age and sex.

ranging from £0.85 to £1.35, because only a small proportion ( $\approx$ 15%) of participants were accompanied to their dental appointment. The total number of visits to a dental practice for NHS care was between three and four across the groups.

## Other private dental care costs

Participant-reported costs for dental care items include the cost of toothbrushes, replacement heads, interdental brushes and private dental care costs, including PI and other private care. Respondents in the 6-monthly PI group were more likely to self-report having private PI. However, as noted in *Chapter 4*, this potentially reflects a misunderstanding of what constitutes private care. Furthermore, an assessment of the validity of self-reported PI indicated a mismatch between self-reported data and routine information on PIs. Therefore, there is the potential for double-counting of participant-incurred costs as NHS-provided treatments may have been interpreted as private PIs in some cases. For this reason, the reader should exercise caution when interpreting any differences in participant-reported private PI costs.

Twenty respondents reported exceptionally high cost values for 'other private treatment', > £1000 (maximum = £11,100) for private dental care in at least one annual questionnaire (some reported high-cost items in more than one questionnaire). These high-cost items were unevenly spread across the randomised groups (n = 4, 1, 5, 3, 5 and 2) and led to skewed distributions and potentially misleading mean estimates. Such treatments included crowns, dentures and orthodontics. These high-cost items were verified with the practice in 14 out of 22 (64%) cases, with the remainder having no record of private treatments at their NHS practice. These high-cost items (> £1000 per year) were removed from the analysis given that there is no feasible clinical link between these treatments and the provision of PI or OHA.

## Total participant costs

Table 23 details all participant-incurred costs. Overall, charges for NHS care represented a small proportion of total participant costs (< 20%), with the largest cost attributed to the purchase of interdental brushes. Total participant costs ranged from £265 to £345 across the groups. No PI with personalised OHA was less costly from the participant's perspective (mean difference, compared with routine OHA and 6-monthly PI, -£64.11, 95% CI -£112.33 to -£15.88). The between-group difference is driven primarily by differences in the use of interdental brushes in England. There were no other significant differences between groups from the participant perspective. Owing to missing data across specific questions in each of the three annual participant questionnaires, participant costs are reported on the basis of multiple imputation of missing cost component data (at the item and questionnaire level). Missing opportunity costs of time and travel are imputed at the total cost level (number of visits multiplied by the calculated unit cost).

## **Benefits**

## Discrete choice experiment

#### Participant characteristics

The DCE was administered online and 667 respondents completed the full questionnaire. Each respondent completed 10 choice tasks (after exclusion of dominance and consistency checks), leading to a total of 20,010 observations in the data set. The average median survey completion time was 17 minutes (interquartile range 13–24 minutes). *Table 24* shows the characteristics of the DCE respondent sample.

Most respondents were from England, but there was over-representation of the general population in Scotland to enable region-level subgroup analysis. In total, 67% of the sample never smoked. There was an even spread across the sample regarding educational attainment and 77% of the sample were either in paid work or retired. Overall, the majority of respondents reported that they were in fair or better general health and dental health. A total of 37 (6%), 232 (35%) and 398 (60%) respondents completed the worst, moderate and best segmented versions of the DCE, respectively.

**TABLE 24** Characteristics of DCE participants

Characteristic	Mean	SD
Age (years)	51	16
	Number of participants ( $n = 667$ )	% of tot
Sex		
Male	308	46.18
Female	359	53.82
Currently living in		
England	482	72.26
Scotland	125	18.74
Wales	44	6.60
Northern Ireland	14	2.10
Isle of Man	2	0.30
Currently a smoker		
Yes	98	14.69
No	447	67.02
Previously a smoker	122	18.29
Annual gross income (£)		
< 20,800	247	37.03
20,800–41,600	211	31.63
≥ 41,600	119	17.84
Prefer not to answer/blank	90	13.49
Educational attainment		
O levels/SVQ (level 1 or 2)/1 A level	183	27.44
≥ 2 A levels/SVQ (level 3)	101	15.14
Degree	164	24.59
Professional qualifications	81	12.14
Apprentice qualification	13	1.95
Vocational/foreign/other/none	125	18.74
Employment		
Any paid employment	326	48.88
Unemployed or seeking work	24	3.60
Retired	185	27.74
Student	20	3.00
Other	112	16.79

TABLE 24 Characteristics of DCE participants (continued)

	Number of participants ( $n = 667$ )	% of total
Self-reported dental health		
Very poor	6	0.90
Poor	28	4.20
Fair	200	29.99
Good	315	47.23
Very good	118	17.69
Self-reported general health		
Very poor	10	1.50
Poor	47	7.05
Fair	169	25.34
Good	343	51.42
Very good	98	14.69

A level, advanced level; O level, ordinary level; SVQ, Scottish Vocational Qualification.

Table 25 presents descriptive statistics detailing the DCE participants' experience of dental care services. Most respondents were registered with a dental practice. A range of methods of payment for dental care were reported. A total of 62% of the sample had seen a dental hygienist in the past and 87% of the sample had had a PI in the past, with 77% having one at least annually. Respondents in the sample had more experience of contact with dental hygienists and having a PI than the general population: 47% had experienced treatment from a hygienist and 80% had had a PI.<sup>11</sup> Overall, the majority in the sample were regular dental attenders (86%), defined as seeing the dentist every 2 years or more often. By comparison, our sample had fewer regular attenders than the general population (the ADHS 2009<sup>11</sup> found that 71% of respondents attend the dentist at least every 2 years). This may be due to an element of self-selection into the DCE survey.

## Results of analysis models

The results of the mixed logit model are presented in *Table 26*.

The positive value for the ASC indicates a preference among the general population to have any package of dental care as opposed to none. The significance of the SD around the constant term indicates the presence of significant preference heterogeneity for opting into a dental care package, with some respondents more likely to opt out (choosing no dental care package). Across all choice sets, 288 (43%) respondents always opted in to the choice task and 45 (7%) always opted out, choosing no dental care package.

Respondents valued having both personalised OHA and PI regardless of provider. They preferred to receive personalised OHA and 12-monthly PI from the dentist. However, who provided the care was less important for the 6-monthly PI. As expected, respondents prefer to have gums that do not bleed and teeth that look and feel clean. The negative coefficient on the annual cost attribute indicates that respondents prefer dental packages that cost less.

Willingness-to-pay estimates, together with their CIs, should be interpreted as the general population's valuation of each attribute and level. The differences between WTP values indicate how much the general population values moving from one health-care package to another. For example, the general population would be willing to pay an additional £90.53 per year over 3 years to move from having bleeding gums

TABLE 25 Participant dental care experience

Characteristic	Number of participants	% of total
Registered with a dental practice		
Yes	636	95.35
No	26	3.90
Do not know	5	0.75
Normally pay for dental care		
Co-charge	307	46.03
NHS pays all cost	127	19.04
Private: pay full cost	143	21.44
Dental treatment plan	65	9.75
Dental insurance	12	1.80
Never had dental care	6	0.90
Do not know	7	1.05
Ever been to visit a dental hygienist		
Yes	414	62.07
No	213	31.93
Do not know	40	6.00
Normally have a PI		
> every 3 months	10	1.52
Every 3 months	65	9.89
Every 6 months	309	47.03
Every year	121	18.42
Every 2 years	22	3.35
< every 2 years	46	7.00
Never	84	12.79
Regular attendance <sup>a</sup>		
Attends at least every 2 years	570	85.84
Attends less often	94	14.16

a Three respondents had missing data on this question.

very often (very often: -£54.13) to never having bleeding gums (never: £36.40). They would be willing to pay £145.50 per year to move from having teeth that look and feel very unclean to teeth that look and feel very clean. The difference between these valuations indicates that the general population attaches a greater value to shifts in average aesthetic outcome than to changes in frequency of bleeding 3 years after signing up to the dental package.

The WTP estimates can thus be used to calculate the general population's WTP for packages of dental care. Assuming that a participant in the trial received no OHA/6-monthly PI from the dentist, hardly ever had bleeding gums at the final annual questionnaire and had teeth that, on average over the period, looked and felt moderately clean, the general population would be willing to pay £93.93 per year for this package (i.e. £43.95 - £13.50 + £29.83 + £20.55 + £13.10). Following a similar approach, the results of the DCE are mapped to each individual to enable a CBA to be completed alongside the trial.

TABLE 26 Results from general population DCE: mixed logit model

		WTP (£)									
Attribute/level	Model estimates	Mean WTP compared with mean attribute level	95% CI	Mean WTP compared with reference level							
No OHAª	-0.143***	-13.50	-18.56 to -8.45	_							
Personalised OHA from dentist	0.122***	11.50	6.61 to 16.39	25.00							
Personalised OHA from hygienist	0.021	2.00	-2.89 to 6.90	15.50							
No Pl <sup>a</sup>	-0.721***	-68.24 <sup>b</sup>	–76.52 to –59.96	_							
12-month PI from dentist	0.194***	18.31 <sup>b</sup>	10.68 to 25.93	86.55							
12-month PI from hygienist	-0.067*	-6.32 <sup>b</sup>	-13.63 to 0.98	61.92							
6-month PI from dentist	0.315***	29.83 <sup>b</sup>	22.23 to 37.43	98.07							
6-month PI from hygienist	0.279***	26.43 <sup>b</sup>	19.21 to 33.65	94.67							
Bleeding gums											
Never <sup>a</sup>	0.385***	36.40	26.82 to 45.98	_							
Hardly ever	0.307***	29.03	20.55 to 37.52	-7.37							
Occasionally	-0.046	-4.40	-11.40 to 2.61	-40.80							
Fairly often	-0.073	-6.90	-16.61 to 2.80	-43.30							
Very often	-0.572***	-54.13	−73.37 to −34.90	-90.53							
Teeth look and feel											
Very unclean	-0.899***	-85.03	-104.55 to -65.52	-							
Unclean	-0.393***	-37.12	−47.24 to −27.01	47.91							
Moderately clean	0.139***	13.10	5.97 to 20.23	98.13							
Clean	0.514***	48.59	39.91 to 57.26	133.62							
Very clean <sup>a</sup>	0.639***	60.47	50.76 to 70.18	145.50							
Annual cost	-0.011***	_	_	_							
ASC											
Mean	0.465***	43.95	29.49 to 58.41	-86.42°							
SD	1.470***										
Log likelihood	-5174.58										
Number of observations	20,010										
Number of respondents	667										
AIC	10,383										
BIC	10,518										

<sup>\*</sup>p < 0.1, \*\*p < 0.05, \*\*\*p < 0.01.

AIC, Akaike information criterion; BIC, Bayesian information criterion.

a Indicates reference level.

b The WTP tariffs given for no PI, 12-monthly PI and 6-monthly PI can also be interpreted as 0, 3 and 6 PIs over a 3-year period. Following the step-wise linear calculation of intervening PIs outlined in *Chapter 3*, WTP tariffs for 0–9 PI provided by the dentist are –£68.24, –£39.39, –£10.54, £18.31, £22.15, £25.99, £29.83, £33.67, £37.51 and £41.35, respectively. WTP tariffs for 0–9 PI provided by the hygienist are –£68.24, –£47.60, –£26.96, –£6.32, £4.59, £15.51, £26.43, £37.35, £48.26 and £59.18, respectively.

c The WTP for package with all attributes at reference level.

## Validity of responses

The estimates of annual WTP for PI (relative to none) fall within a feasible range of prices charged in the private market for PI (i.e. between £49 and £86 per PI provided by the dentist). In terms of the internal validity checks, 535 (80%) respondents passed the consistency of preferences check (i.e. answering the same way for exactly the same choice twice). A total of 498 (75%) respondents passed the non-satiation (dominance) test. Very few respondents failed both tests (≈5%). The high percentage passing such tests is in line with other DCE studies.<sup>65</sup>

## Analysis of subgroups

Subgroup analyses were conducted for participant characteristics by sex, region and income, as well as experience of PI and dental hygienists. *Table 27* describes the impact of subgroup membership on preferences, through interaction terms between effects coded categorical variables and attribute-level main effects. Subgroup effects were considered significant at a p-value of < 0.01.

The likelihood ratio tests show no overall effect on preferences (jointly across all model parameters) for UK region or smoking status. For subgroups for which the tests indicated an impact of subgroup on preferences, the following findings were observed:

- Respondents with experience of the hygienist valued the dental care packages more highly (disutility
  associated with cost attribute was lower). They also gained less utility from teeth that look and feel
  only moderately clean.
- The significant interaction effect on the ASC with experience of PI indicates that those with experience
  of PI are more likely to commit to a dental care package. They also valued the dental care packages
  more highly.
- Neither experience of seeing the dental hygienist or of having PI affected the preferences of respondents for having these services, or for whether it is the dentist or hygienist who provides the service.
- Females in the sample gained greater utility from dental care packages in which teeth look and feel clean or very clean. Similarly, they experienced greater disutility from teeth that look and feel moderately clean or unclean.
- As expected, the cost attribute had less of an impact on the preferences of higher-income respondents.

The subgroup analyses indicate that, although there were some differences across subgroups, in general, the direction of attribute-level effect remained similar with respondents preferring PI and personalised OHA, less bleeding and teeth that look and feel clean and healthy, as well as preferring lower-cost dental care packages.

#### Benefits: willingness to pay

Table 28 presents the mean WTP values across randomised groups, including the component parts of total WTP based on the mapping of DCE results to treatment received and trial outcomes. The average total benefits in terms of WTP ranged from £79.09 to £207.65 across the groups. The policy of 6-monthly Pl and personalised OHA is associated with the highest average benefit. The difference in benefits between this policy and standard care (routine OHA and 6-monthly Pl) is statistically significant. Compared with standard care, having no Pl or a 12-monthly Pl and routine OHA is associated with significantly lower benefits. There was no significant difference between the benefits for personalised OHA with either 12-monthly Pl or no Pl and standard care, implying that a reduction in Pl can be compensated for by the provision of personalised OHA. The findings were consistent across regions. The average benefit was generally lower in England, as Pls were more likely to be provided by hygienists in England than in Scotland (following the assumption that the provider of baseline OHA/Pl would consistently provide the trial intervention over follow-up).

The differences in benefits observed between groups were driven primarily by the differences in WTP for PI and personalised OHA. For example, the 6-monthly PI with routine OHA had a higher WTP (–£27.43) associated with PIs than no PI with routine OHA (–£82.92), a difference of £55.49. Note that the negative

NIHR Journals Library www.journalslibrary.nihr.ac.uk

TABLE 27 Analysis of DCE subgroups

Attribute levels	Base case	Hygienist experience <sup>a</sup>	Region (Scotland) <sup>b</sup>	Sex (f) <sup>c</sup>	Income (> £20,800) <sup>d</sup>	Smoker (ever) <sup>e</sup>	Experience of PI <sup>f</sup>
Main effects							
Personalised OHA from dentist	0.122***	0.133***	0.143***	0.124***	0.121***	0.140***	0.144***
Personalised OHA from hygienist	0.021	0.016	0.027	0.019	0.022	0.011	-0.027
12-monthly PI from dentist	0.194***	0.199***	0.232***	0.194***	0.193***	0.208***	0.170***
12-monthly PI from hygienist	-0.067*	-0.065	-0.089*	-0.071*	-0.065*	-0.100**	-0.171***
6-monthly PI from dentist	0.315***	0.309***	0.316***	0.323***	0.315***	0.319***	0.280***
6-monthly PI from hygienist	0.279***	0.274***	0.281***	0.283***	0.277***	0.294***	0.261***
Bleeding gums							
Hardly ever	0.307***	0.313***	0.337***	0.303***	0.312***	0.301***	0.280***
Occasionally	-0.046	-0.032	-0.094**	-0.055	-0.043	-0.053	-0.143**
Fairly often	-0.073	-0.058	-0.012	-0.073	-0.084	-0.055	-0.057
Very often	-0.572***	-0.599***	-0.591***	-0.556***	-0.573***	-0.608***	-0.429**
Teeth look and feel							
Very unclean	-0.899***	-0.989***	-0.887***	-0.845***	-0.868***	-0.901***	-0.719***
Unclean	-0.393***	-0.368***	-0.308***	-0.392***	-0.400***	-0.398***	-0.311***
Moderately clean	0.139***	0.183***	0.109**	0.120***	0.134***	0.138***	0.049
Clean	0.514***	0.497***	0.448***	0.494***	0.508***	0.506***	0.402***
Annual cost	-0.011***	-0.011***	-0.011***	-0.011***	-0.011***	-0.011***	-0.012***
ASC							
Mean	0.465***	0.402***	0.448***	0.459***	0.472***	0.417***	0.239**
SD	1.470***	1.435***	1.476***	1.473***	1.459***	1.464***	1.420***
Interaction terms for subgroup	with main attr	ibute level effects					
Personalised OHA from dentist		-0.034	0.032	0.011	-0.018	0.044	-0.023
Personalised OHA from hygienist		0.017	0.010	0.022	0.021	-0.028	0.058
12-monthly PI from dentist		-0.007	0.062	-0.021	-0.009	0.036	0.030

DOI: 10.3310/hta22380

Attribute levels	Base case	Hygienist experience <sup>a</sup>	Region (Scotland) <sup>b</sup>	Sex (f) <sup>c</sup>	Income (> £20,800) <sup>d</sup>	Smoker (ever) <sup>e</sup>	Experience of PI <sup>f</sup>
12-monthly PI from hygienist		0.013	-0.032	0.027	-0.025	-0.082*	0.130**
6-monthly PI from dentist		0.020	-0.001	0.103**	0.053	0.012	0.055
6-monthly PI from hygienist		0.001	-0.004	0.071*	0.069*	0.033	0.025
Bleeding gums							
Hardly ever		-0.022	0.045	0.069	0.098**	-0.018	0.033
Occasionally		-0.046	-0.075*	0.004	0.076**	-0.018	0.110
Fairly often		-0.036	0.094	-0.080	-0.059	0.057	-0.026
Very often		0.057	-0.024	-0.075	-0.204**	-0.083	-0.160
Teeth look and feel							
Very unclean		0.187	0.028	-0.334***	0.117	-0.004	-0.213
Unclean		-0.077	0.124*	-0.216***	-0.067	-0.003	-0.106
Moderately clean		-0.111***	-0.047	0.160***	-0.047	-0.004	0.104
Clean		0.070	-0.102*	0.177***	-0.014	-0.020	0.140*
Annual cost		0.002***	0.000	-0.002***	0.001***	-0.001**	0.002***
ASC (mean)		0.307***	-0.024	-0.064	0.081	-0.134*	0.320***
Likelihood ratio test							
Likelihood ratio	2176	2068	2177	2163	2132	2148	2025
$\chi^2  (df = 16)$		86.44	16.82	93.27	42.23	20.02	81.19
<i>p</i> -value		0.0000	0.3976	0.0000	0.0004	0.2195	0.0000

<sup>\*</sup>p < 0.1, \*\*p < 0.05, \*\*\*p < 0.01.

df, degrees of freedom; f, female.

a Experience of hygienist (1 = ever visited a hygienist; -1 = never visited).

b Region (1 = Scotland; -1 = rest of the UK).

c Sex (1 = female; -1 = male).

d Income (1 = moderate/high income reported,  $\geq$  £20,800 per year; -1 = low income <£20,800 per year).

e Smoker (1 = current or previous smoker; -1 = never smoked).

f Experience of PI (1 = ever had a PI; -1 = never had a PI).

NIHR Journals Library www.journalslibrary.nihr.ac.uk

OHA, WTP (f) **Routine OHA Personalised OHA** No PI 12-monthly PI 6-monthly PI **Total** No PI 12-monthly PI 6-monthly PI Total Region Mean Mean SD Mean SD Mean Mean Mean SD Mean SD UK (N = 290)(N = 866)(N = 338)(N = 289)(N = 287)(N = 334)(N = 335)(N = 1007)Constant 98.89 0.00 98.89 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 OHA -30.38 0.00 -30.38-30.38 0.00  $-30.38 \quad 0.00$ 19.80 9.66 19.99 9.56 19.88 9.62 19.89 0.00 9.60 PΙ -82.9270.59 -56.5560.80 -27.43 76.87 -55.60 73.29 -93.7363.83 -51.77 61.96 -21.6070.98 -55.65 71.97 Bleeding 30.02 45.86 29.98 47.63 31.74 45.15 33.86 45.78 42.39 35.25 34.19 35.28 41.60 33.48 44.17 44.10 Look and feel 63.12 61.06 67.05 59.84 75.61 58.32 68.60 59.96 69.52 61.83 66.69 62.00 75.24 58.34 70.47 60.84 Total WTP 79.09 128.46 97.68 114.25 95.04 146.46 104.91 113.30 103.01 104.80 167.28 95.92 207.65 116.31 167.87 110.72 Mean (95% CI) benefit -67.65-30.75N/A -17.7219.70 61.67 N/A difference vs. routine (-86.50 to -48.81) (-48.65 to -12.85) (-39.37 to 3.93) (-1.64 to 41.04) (40.19 to 83.14) OHA with 6-monthly PI Scotland (N = 196)(N = 195)(N = 195)(N = 586)(N = 252)(N = 255)(N = 252)(N = 759)Constant 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 98.89 0.00 OHA 0.00 -30.38 0.00 23.14 7.14 -£30.38 -30.380.00  $-30.38 \quad 0.00$ 23.16 7.13 23.03 7.28 23.25 7.03 PΙ -69.16 74.79 61.52 -9.42 72.91 -41.47 74.08 -89.21 58.52 -8.77 -46.80 71.16 -45.7164.70 -42.4666.05 Bleedina 24.97 45.56 31.52 43.20 26.13 49.98 27.52 46.37 34.40 45.06 31.86 41.91 35.07 45.42 33.77 44.13 Look and feel 62.85 £59.80 67.31 60.76 69.88 75.74 56.53 70.64 60.67 65.45 73.63 63.25 66.36 61.97 60.76 Total WTP 87.75 101.92 119.81 99.26 158.86 106.56 122.15 106.57 137.12 105.57 177.67 93.15 224.17 112.50 179.65 109.82 Mean (95% CI) benefit -71.63-39.09N/A -22.5814.31 64.09 N/A difference vs. routine (-94.39 to -48.86) (-61.03 to -17.14) (-45.90 to 0.74) (-8.62 to 37.24) (40.95 to 87.22) OHA with 6-monthly PI

	OHA, WTP (£)																
	Routine (	Routine OHA									Personalised OHA						
	No PI	No PI		12-monthly PI		6-monthly PI			No PI		12-monthly PI		6-monthly PI		Total		
Region	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	
England	(N = 93)		(N = 92)		(N = 95)	)	(N = 28	0)	(N = 82)		(N = 83)		(N = 83)		(N = 24	8)	
Constant	98.89	0.00	98.89	0.00	98.89	0.00	98.89	0.00	98.89	0.00	98.89	0.00	98.89	0.00	98.89	0.00	
OHA	-30.38	0.00	-30.38	0.00	-30.38	0.00	-30.38	0.00	9.45	9.07	10.68	9.75	9.65	9.20	9.93	9.32	
PI	-111.93	49.83	-79.52	52.57	-64.42	71.67	-85.16	62.04	-107.61	59.31	-80.38	63.77	-60.55	71.58	-82.75	67.67	
Bleeding	40.66	44.84	43.15	36.97	37.89	41.45	40.54	41.17	32.17	48.04	38.46	43.62	35.79	40.34	35.49	44.05	
Look and feel	63.67	62.04	70.44	56.33	79.71	55.07	71.31	58.18	68.40	57.29	67.71	62.17	73.72	63.75	69.96	61.13	
Total WTP	60.92	85.69	102.59	84.67	120.74	96.88	94.81	92.56	101.50	97.95	135.36	97.61	157.50	113.78	131.70	105.67	
Mean (95% CI) benefit difference vs. routine OHA with 6-monthly PI	–59.82 (–89.86 to	-29.78)	-15.06 (-44.61 to 1	4.48)			N/A		–12.45 (–51.68 to 2	26.79)	25.39 (–13.44 to	64.21)	45.46 (6.08 to	84.86)	N/A		

N/A, not applicable.

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Socience Park, Southampton SO16 7NS, UK.

mean WTPs for PI across all groups are due to the relatively few PIs provided across all groups, and the use of effects coding in which WTP values are estimated relative to the mean of the attribute. In terms of interpretation, the focus should therefore be on differences in WTP for PI across groups. Similarly, for example, 6-monthly PI with personalised OHA had a higher WTP for the OHA component (£19.88) than did 6-monthly PI with routine OHA (–£30.38), an additional WTP of £50.26 for the OHA provided in the personalised group. There were some differences in benefits across regions, driven by the provider of care (more treatments assumed to be provided by hygienists in England than in Scotland). The differences across groups appear to be driven mainly by the services provided rather than the bleeding or aesthetic outcomes. This observation matches the findings reported in the clinical effectiveness chapters.

## Cost-benefit analysis

#### Base-case analysis: NHS perspective

The CBA reports incremental net benefits (incremental benefits – incremental costs) for each policy, compared with standard care (routine OHA, with 6-monthly PI). *Table 29* shows results from the base-case NHS dental health-care cost perspective. The average incremental net benefit ranged from –£69 to £48 across the groups. For the base-case analysis, personalised OHA with 6-monthly PI is associated with the highest positive incremental net benefit compared with standard care and the difference is statistically significant. Therefore, this is the preferred option from a welfare-maximising perspective. The only other policy that is associated with positive incremental net benefits is personalised OHA with 12-monthly PI, although it should be noted that the difference in net benefits between this policy and routine care is not statistically significant.

Confidence ellipses on the cost-effectiveness plane for each comparison are reported in *Appendix 2* and show that personalised OHA with 6-monthly PI has a high probability of generating a positive incremental net benefit (i.e. with plots to the right of the diagonal line on the cost-benefit plane). Routine OHA with either no PI or 12-monthly PI is associated with negative incremental net benefits compared with standard care, and the differences in net benefits are statistically significant. *Figure 19, Appendix 2* shows that there is more uncertainty surrounding the incremental net benefits associated with personalised OHA (no PI or 12-monthly PI) relative to routine OHA (6-monthly PI), with confidence ellipses crossing the diagonal.

The main conclusion is consistent across regions, although the difference in positive incremental net benefits for personalised OHA and 6-monthly PI is not statistically significant for England. This may be caused by the relatively smaller sample in England.

### Wider perspective cost-benefit analysis

Table 30 presents the CBA results using the NHS and participant perspectives for costs. The preferred policy option is the same as in the base-case analysis, namely personalised OHA with 6-monthly Pl. The incremental net benefit is higher than in the base-case analysis (£68 vs. £48). Note that the CI is much wider, reflecting the relatively large variation in participant costs. The preferred policy varies across regions when using the wider perspective. For England, personalised OHA with no Pl generates the largest incremental net benefit. This result is driven by the lower cost of this policy compared with standard care. As noted in *Total participant costs*, this appears to be driven mainly by the lower costs of interdental brushes. The extent to which this can be attributable directly to the intervention is questionable. Furthermore, conclusions about participant-reported costs should be interpreted in the light of concerns about double-counting of private/NHS Pl.

This result may have also caused the second preferred policy for the UK as a whole to be different from the base-case analysis. The second preferred policy for the base-case analysis was personalised OHA with 12-monthly PI (incremental net benefit of £3) while the second preferred policy when taking the wider perspective was personalised OHA with no PI (incremental net benefit of £58).

TABLE 29 Cost-benefit analysis results: base case - NHS dental perspective costs (£)<sup>a</sup>

Region	Mean (SD) costs	Mean difference in costs <sup>b</sup> (95% CI) vs. routine OHA (6-monthly PI)	Mean (SD) benefits	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI)	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI)
UK					
Routine OHA					
6-monthly PI ( $n = 289$ )	74 (106)	_	146 (105)	_	_
No PI (n = 288)	75 (122)	2 (-19 to 22)	79 (98)	-67 (-86 to -48)	-69 (-97 to -41)
12-monthly PI ( $n = 285$ )	81 (140)	8 (-12 to 29)	114 (95)	−30 (−48 to −12)	−38 (−65 to −12)
Personalised OHA					
No PI ( <i>n</i> = 333)	62 (116)	-6 (-28 to 16)	128 (105)	-17 (-38 to 4)	-11 (-37 to 15)
12-monthly PI ( $n = 338$ )	86 (133)	18 (-4 to 40)	167 (96)	21 (0 to 42)	3 (-23 to 29)
6-monthly PI ( $n = 335$ )	83 (134)	15 (-8 to 37)	208 (116)	63 (42 to 84)	48 (22 to 74)
Scotland					
Routine OHA					
6-monthly PI ( $n = 195$ )	72 (93)	_	159 (107)	_	_
No PI ( <i>n</i> = 195)	64 (87)	-9 (-28 to 11)	88 (102)	-71 (-94 to -48)	-63 (-92 to -34)
12-monthly PI ( $n = 193$ )	67 (83)	-4 (-23 to 16)	120 (99)	−39 (−61 to −17)	−35 (−64 to −6)
Personalised OHA					
No PI $(n = 252)$	61 (103)	-5 (-28 to 18)	137 (106)	-23 (-49 to 3)	-18 (-46 to 10)
12-monthly PI ( $n = 255$ )	80 (121)	12 (–11 to 35)	178 (93)	14 (-11 to 40)	2 (-25 to 29)
6-monthly PI ( $n = 252$ )	79 (120)	12 (–11 to 35)	224 (112)	64 (38 to 89)	52 (24 to 79)
England					
Routine OHA					
6-monthly PI $(n = 94)$	79 (129)	_	121 (97)	_	_
No PI (n = 93)	99 (172)	25 (-25 to 74)	61 (86)	-59 (-90 to -29)	-84 (-143 to -25)
12-monthly PI $(n = 92)$	111 (214)	32 (–17 to 81)	103 (85)	-15 (-44 to 15)	-47 (-104 to 11)
Personalised OHA					
No PI (n = 81)	65 (149)	-13 (-68 to 43)	102 (98)	-10 (-49 to 29)	3 (-57 to 62)
12-monthly PI ( $n = 83$ )	107 (161)	31 (–25 to 87)	135 (98)	28 (-11 to 66)	-3 (-62 to 56)
6-monthly PI ( $n = 83$ )	95 (169)	19 (-37 to 74)	157 (114)	48 (9 to 87)	29 (-30 to 89)

a Costs and benefits rounded to the nearest whole pound.

b Note that mean difference in costs presented here are based on imputed data, accounting for correlation in costs and benefits, and, therefore, cannot be directly compared with *Table 19*.

TABLE 30 Cost-benefit analysis results: NHS and participant perspective costs (£)<sup>a</sup>

			<u> </u>		
Region	Mean (SD) costs	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI)	Mean (SD) benefits	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI)	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI)
UK					
Routine OHA					
6-monthly PI ( $n = 289$ )	420 (323)	_	146 (105)	_	_
No PI (n = 288)	378 (312)	-41 (-93 to 11)	79 (98)	-68 (-86 to -49)	-26 (-81 to 28)
12-monthly PI ( $n = 285$ )	398 (302)	-17 (-69 to 36)	114 (95)	−31 (−49 to −13)	-14 (-70 to 42)
Personalised OHA					
No PI (n = 333)	327 (279)	–75 (–126 to –25)	128 (105)	-17 (-39 to £4)	58 (7 to 110)
12-monthly PI ( $n = 338$ )	384 (302)	-21 (-73 to 30)	167 (96)	20 (-1 to 41)	41 (-11 to 94)
6-monthly PI ( $n = 335$ )	396 (295)	-6 (-57 to 46)	208 (116)	62 (41 to 83)	68 (15 to 120)
Scotland					
Routine OHA					
6-monthly PI ( $n = 195$ )	371 (257)	_	159 (107)	_	_
No PI (n = 195)	323 (264)	-49 (-106 to 8)	88 (102)	-72 (-94 to -49)	-23 (-83 to 37)
12-monthly PI ( $n = 195$ )	347 (255)	-25 (-83 to 33)	120 (99)	−39 (−61 to −17)	-14 (-75 to 47)
Personalised OHA					
No PI (n = 252)	320 (266)	-41 (-96 to 14)	137 (106)	-22 (-47 to 2)	18 (-38 to 74)
12-monthly PI ( $n = 255$ )	356 (291)	-13 (-68 to 42)	178 (93)	14 (-10 to 38)	27 (-29 to 84)
6-monthly PI ( $n = 252$ )	377 (284)	15 (-40 to 70)	224 (112)	64 (40 to 88)	49 (-7 to 105)
England					
Routine OHA					
6-monthly PI $(n = 94)$	520 (411)	_	121 (97)	_	_
No PI (n = 93)	494 (371)	-20 (-125 to 86)	61 (86)	-60 (-90 to -29)	-40 (-148 to 68)
12-monthly PI ( $n = 92$ )	507 (360)	6 (-102 to 115)	103 (85)	-15 (-45 to 15)	-21 (-131 to 89)
Personalised OHA					
No PI (n = 81)	346 (319)	-154 (-265 to -43)	102 (98)	-12 (-52 to 28)	142 (31 to 253)
12-monthly PI (n = 83)	469 (320)	-14 (-129 to 100)	135 (98)	26 (-14 to 66)	40 (-73 to 153)
6-monthly PI (n = 83)	453 (322)	-40 (-153 to 72)	157 (114)	46 (5 to 86)	86 (–27 to 200)
a Costs and benefits rounded	to the nearest	whole f.			

## Sensitivity analysis results

Table 31 reports the incremental net benefit results for a range of sensitivity analyses. All sensitivity analyses were conducted using NHS-perspective dental care costs. Further details of mean and incremental costs and benefits are presented for each analysis in *Appendix 2, Section 6: detailed results of further health economics analyses*, at a regional and UK level. Overall, the conclusions of our analyses remain robust to the range of sensitivity analyses undertaken. In all cases, a policy of personalised OHA with 6-monthly PI was associated with the greatest incremental net benefit relative to standard care. This was statistically significant for all scenarios except those for which an additional charge was included for personalised OHA. This analysis was conducted to reflect the likely opportunity cost of time spent delivering personalised OHA if it were to be rolled out as a policy. However, under current reimbursement structures, such a charge is unlikely to be directly incurred by the NHS. In all cases, routine OHA with no PI was associated with the lowest incremental net benefit relative to standard care. It should be noted that, across all sensitivity analyses, the results were driven by high estimates of WTP for PI in particular. Therefore, our results and conclusions are most applicable to decision-makers wishing to maximise societal welfare.

**TABLE 31** Main sensitivity analysis results

		Incremental net	cremental net benefits (95% CI) (£) vs. routine OHA (6-monthly PI)						
Po	licy	Base-case analysis	Including baseline and final visits	Discounting at 0%	Discounting at 6%	Assuming all PI and OHA delivered by dentist	Assuming all PI and OHA delivered by hygienist		
Roi	utine OHA								
	6-monthly PI $(n = 289)$	_	-	-	-	-	-		
	No PI (n = 288)	-69 (-97 to -41)	-68 (-103 to -33)	-70 (-99 to -42)	-67 (-93 to -40)	-70 (-98 to -43)	-61 (-88 to -34)		
	12-monthly PI ( <i>n</i> = 285)	−38 (−65 to −12)	-33 (-67 to 1)	-40 (-68 to -12)	−38 (−65 to −11)	−37 (−65 to −10)	-37 (-64 to -10)		
Per	rsonalised OH	4							
	No PI (n = 333)	-11 (-37 to 15)	9 (-24 to 42)	-13 (-40 to 15)	-10 (-36 to 15)	-8 (-34 to 19)	-18 (-44 to 8)		
	12-monthly PI ( <i>n</i> = 338)	3 (-23 to 29)	25 (-8 to 57)	1 (-26 to 28)	4 (-22 to 29)	8 (-18 to 35)	-13 (-39 to 13)		
	6-monthly PI $(n = 335)$	48 (22 to 74)	69 (36 to 102)	49 (21 to 76)	47 (22 to 73)	55 (28 to 82)	29 (2 to 55)		
		Incremental net	t benefits (95% C	I) (£) vs. routine	OHA (6-monthly	PI)			
Po	licy	Base-case analysis	Additional cost for personalised OHA	Decreasing the unit price of a UDA (-20%)	Increasing the unit price of a UDA (+20%)	Alternative map	ping of ne from the DCE		
Roi	utine OHA								
	6-monthly PI $(n = 289)$	_	-	-	-	-			
	No PI (n = 288)	-69 (-97 to -41)	-68 (-101 to -36)	-66 (-91 to -40)	-68 (-97 to -39)	−64 (−89 to −39)			
	12-monthly PI ( <i>n</i> = 285)	−38 (−65 to −12)	−39 (−70 to −7)	-36 (-61 to -10)	-40 (-69 to -10)	−37 (−61 to −12)			
Per	rsonalised OH	4							
	No PI (n = 333)	-11 (-37 to 15)	-54 (-104 to -4)	-12 (-37 to 13)	-9 (-38 to 19)	-5 (-29 to 18)			
	12-monthly PI ( <i>n</i> = 338)	3 (–23 to 29)	-48 (-100 to 5)	3 (-21 to 28)	2 (-27 to 30)	7 (-16 to 31)			
	6-monthly PI $(n = 335)$	48 (22 to 74)	28 (-15 to 70)	49 (23 to 74)	48 (20 to 77)	44 (20 to 68)			

### Cost-consequences analysis

The balance sheet of costs and outcomes outlined in *Table 32* narratively summarises the advantages and disadvantages of the different policy considerations within a cost–consequences framework. The analysis makes no attempt to conduct any further quantitative synthesis of economic findings.

TABLE 32 Cost-consequences analysis: narrative summary

Policy	Analysis	Advantages/positives	Disadvantages/negatives
Routine OHA (6-monthly PI): SC	DCE	6-monthly PI is preferred by the general population	Personalised is preferable to routine OHA
	Clinical effectiveness	-	No evidence of clinical benefit
	Incremental costs (f) (vs. SC)	N/A	N/A
	Incremental benefits (£) (vs. SC)	N/A	N/A
	INB (£) (vs. SC)	N/A	N/A
Routine OHA (no PI)	DCE	-	Least preferred policy by the general population
	Clinical effectiveness	A policy of full disinvestment in PI would not adversely affect bleeding gums	-
	Incremental costs (£)	NHS -3 (95% CI -18 to 12)	N/A
	(vs. SC)	Participant -40 (95% CI -84 to 4)	
	Incremental benefits (£) (vs. SC)	N/A	-68 (95% CI -87 to -49)
	INB (£) (vs. SC)	N/A	–69 (95% CI –97 to –41)
Routine OHA (12-monthly PI)	DCE	-	12-monthly PI preferred to none but 6-monthly OHA would be more highly valued
	Clinical effectiveness	-	No evidence of clinical benefit of PI
	Incremental costs (£) (vs. SC)	Participant –23 (95% CI –70 to 23)	NHS 0 (95% CI –15 to 16)
	Incremental benefits (£) (vs. SC)	N/A	−31 (95% CI −49 to −13)
	INB (£) (vs. SC)	N/A	−38 (95% CI −65 to −12)
Personalised OHA (no Pl)	DCE	Personalised OHA preferred to routine	Policies with delivery of 12-monthly or 6-monthly Pl would be preferred
	Clinical effectiveness	-	No evidence of clinical benefit
	Incremental costs (£)	NHS –15 (95% CI –34 to 4)	N/A
	(vs. SC)	Participant –64 (95% CI –112 to –16)	
	Incremental benefits (£) (vs. SC)	N/A	–18 (95% CI –39 to 4)
	INB (£) (vs. SC)	N/A	–11 (95% CI –37 to 15)
Personalised OHA (12-monthly PI)	DCE	Personalised OHA preferred to routine; reduction in PI could, in part, be compensated for by personalised OHA	-
	Clinical effectiveness	-	No evidence of clinical benefit
	Incremental costs (£) (vs. SC)	Participant –17 (95% CI –68 to 35)	NHS 6 (95% CI –16 to 28)

TABLE 32 Cost-consequences analysis: narrative summary (continued)

Policy	Analysis	Advantages/positives	Disadvantages/negatives
	Incremental benefits (£) (vs. SC)	20 (95% CI –2 to 41)	N/A
	INB (£) (vs. SC)	3 (95% CI –23 to 29)	N/A
Personalised OHA (6-monthly PI)	DCE	Most preferred policy by the general population	-
	Clinical effectiveness	-	No evidence of clinical benefit
	Incremental costs (£) (vs. SC)	Participant –30 (95% CI –82 to 21)	NHS 12 (95% CI –11 to 34)
	Incremental benefits (£) (vs. SC)	62 (95% CI 40 to 83)	N/A
	INB (£) (vs. SC)	48 (95% CI 22 to 74)	N/A

The data presented in the balance sheet indicate the trade-offs decision-makers need to consider when determining the most efficient policy of PI and OHA to implement. It shows that, although personalised OHA with 6-monthly PI is the most cost-beneficial policy, owing to the high value placed on the service by the general population, there is no clear evidence of clinical benefit, or changes to provider beliefs. The preferred policy depends on the viewpoint taken. If the aim is to maximise welfare from a fixed NHS budget, then the CBA analysis shows that personalised OHA with 6-monthly PI is the preferred policy. If the aim is to maximise health from a fixed NHS budget, then it could be argued that no PI is the preferred policy given the lack of evidence of clinical benefit and given that it was associated with the lowest average costs (although it should be noted that differences were not significant).

## **Chapter 6** Cohort

#### **Introduction**

The pragmatic design of the IQuaD trial (reported in *Chapter 2*) outlined recruitment of healthy individuals or those with mild to moderate forms of periodontal disease. As one of the intervention groups received no PI, the IQuaD Periodontal Advisory Group recommended that potential participants exhibiting more severe forms of disease, as defined by BPE scores of 4 or \*, should not be recruited into the experimental trial. The most advanced form of periodontal disease affects only a relatively small percentage of the population<sup>11</sup> and there is a paucity of long-term data for these individuals in the primary care setting. A prospective longitudinal cohort study was therefore conducted in conjunction with the IQuaD trial, involving those individuals who were found to be ineligible for the IQuaD trial but who consented to be followed up for 3 years.

#### **Methods**

The IQuaD cohort study was conducted in conjunction with the IQuaD trial and the recruitment protocol of dental practices and participants is summarised in *Chapter 2*.

## **Ethics approval**

Favourable ethics approval for the IQuaD trial was confirmed by the East of Scotland Research Ethics Service on 24 March 2011 (REC reference number 10/S0501/65).

#### **Recruitment and consent**

The IQuaD trial and IQuaD cohort recruitment were conducted in 63 IQuaD trial dental practices in Scotland and north-east England from February 2012 to July 2013. All participants in the cohort study had previously consented to participate in the IQuaD trial. Following baseline clinical screening and outcome measurement, potential participants were advised that they were not eligible to be included in the trial if they were found to have a BPE score of 4 or \* or had an uncontrolled medical condition. They were then invited to participate in the IQuaD cohort study and signed consent to take part in this prospective longitudinal study. The IQuaD cohort study participants subsequently received a routine dental examination from their general dental practitioners, who were advised that the patient was ineligible for the IQuaD trial. The IQuaD cohort study participants received usual care and were not part of the experimental design.

#### **Outcomes**

The clinical and patient-centred outcomes were the same as the IQuaD trial outcomes (outlined in methods). The IQuaD cohort participants completed the same baseline questionnaire as the IQuaD trial participants and were also sent an annual questionnaire for 3 years.

#### **Statistical analyses**

No statistical analyses were planned and only descriptive data of the longitudinal cohort were reported.

#### **Results**

#### Baseline characteristics

Of the 2341 potential participants who attended for a baseline clinical outcome assessment, 160 were found to have a BPE score of 4 and/or \*. These potential participants were invited to participate in the IQuaD cohort study, with 16 (10%) declining to participate (reasons outlined in *Table 6*), resulting in 144 (90%) participants being recruited to the longitudinal study.

#### Clinical outcomes

Clinical outcome assessment, measurement and outcome mean calculations followed the same protocol as the IQuaD trial (outlined in *Chapter 2, Collection of clinical outcome measures*). At baseline, the 144 cohort participants had a mean number of 22 (SD 4.7) teeth present, with gingival inflammation/bleeding recorded in 35.1% (mean) (SD 23.4%) of sites. Supragingival calculus was recorded on 40.6% (mean) (SD 30.2%) of teeth and the mean clinical probing depth of the cohort patients was 2.2 mm (SD 0.4 mm).

Similar to the IQuaD trial, a number of recruited participants were not available for follow-up clinical assessment; 85 (59%) IQuaD cohort participants attended for follow-up clinical assessment at approximately 3 years following recruitment. When considering only the 85 participants who attended for both clinical assessment appointments, the baseline clinical characteristics were as follows: a mean number of 21.6 (SD 5) teeth present, 37.0% (mean) (SD 24%) of sites with recorded gingival inflammation, mean supragingival calculus of 36.0% (SD 27%) and mean clinical probing depth of 2.2 mm (SD 0 mm). A total of 97% (n = 82) of these participants had a BPE score of 4 or \* at baseline; the remaining three participants, who were ineligible due to medical history reasons, had a BPE score of 3 (1%, n = 1) or 2 (2%, n = 2).

#### Patient-centred outcomes

The collection, assessment and mean score calculations of the participant-centred outcomes are the same as in the IQuaD trial outlined in *Chapter 2*.

At baseline, the cohort participants' self-efficacy, assessed as confidence in one's ability to perform a behaviour and measured using a 7-point scale scored from 1 (not at all confident) to 7 (extremely confident), was, on average, 5.0 points (SD 1.2 points; completed the questionnaire, n = 131).

#### **Cognitions**

Perceived behaviour control score, assessed in terms of perceived ease or difficulty of performing the behaviour and measured using a 7-point scale varying from 1 (strongly agree) to 7 (strongly disagree), was, on average, 4.1 points (SD 1.4 points; completed the questionnaire, n = 131).

Attitude towards oral health behaviour, measured using a scale varying from 1 (strongly agree) to 7 (strongly disagree), was very positive, averaging 5.6 points (SD 1.2 points; completed the questionnaire, n = 131).

Subjective norm, assessed as attitude towards the behaviour and perceptions of social pressure to perform the behaviour and measured using a scale of 1 (strongly agree) to 7 (strongly disagree) scored, on average, 4.9 points (SD 1.3 points; completed the questionnaire, n = 131).

The score of mean intention to perform good oral hygiene practice was 5.5 points (SD 1.8 points; completed the questionnaire, n = 114), which was measured using questions to which responses were scored from 0 to 3 (the best possible intention), with 9 being the best possible score.

#### Behaviour

On the self-reported oral hygiene behaviours scale, which ranges from 0 to 9, with 9 being the best possible behaviour, the cohort group had an average score of 4.9 points (SD 1.7 points; completed the questionnaire, n = 130). Like the IQuaD trial participants, the cohort group showed room for improvement on this scale.

#### Quality of life

The self-reported QoL was measured using the OHIP-14, a 14-item oral health-specific patient-centred outcome referring to symptoms in the past 12 months. Each item is scored from to 0 to 4 (very often) and the scores are summed to produce a summary score ranging from 0 to 56, with 56 being the worst outcome. The cohort participants' mean score at baseline was 8.5 points (SD 9.5 points; completed the questionnaire, n = 126).

## **Follow-up characteristics**

#### **Clinical**

Follow-up examinations were conducted from May 2015 to July 2016. At the 3-year clinical follow-up, the 85 participants had an average of 21.2 (SD 5.4) teeth with mean gingival inflammation/bleeding recorded at 41% (SD 25.5%) of sites. More teeth (mean 43.5%, SD 33.1%) exhibited supragingival calculus and the mean clinical probing depths had increased minimally by 0.1 mm to 2.3 mm (SD 0.5 mm).

#### Patient-centred outcomes

At the 3-year follow-up, the cohort participant patient-centred outcomes were similar to the baseline scores. The average self-efficacy score was 5.1 points (SD 1.1 points; completed the questionnaire, n = 101) on a scale of 1–7.

### Cognitions

The average PBC, attitude and subjective norm scores were 4.4 points (SD 1.3 points; completed the questionnaire, n = 87), 5.2 points (SD 1.6 points; completed the questionnaire, n = 87) and 4.5 points (SD 1.4 points; completed the questionnaire, n = 86), respectively, with each measured on a scale of 1–7. The 3-year follow-up average intention score was 5.4 points (SD 1.8 points; completed the questionnaire, n = 83) on a scale from 0 to 9, with 9 being the best possible score.

#### **Behaviour**

On the self-reported oral hygiene behaviours scale, which ranges from 0 to 9, with 9 being the best possible behaviour, the cohort group had a mean score of 5.4 points (SD 1.8 points; completed the questionnaire, n = 87).

#### Quality of life

The mean score on the OHIP-14, which measures self-reported QoL, was 7.8 points (SD 8.0 points; completed the questionnaire, n = 83).

## **Discussion**

The IQuaD cohort study provided a unique opportunity to monitor NHS primary care patients with severe periodontal disease over a 3-year time period. The end-point outcome of periodontal disease is tooth loss, which is thought to be concentrated in a relatively small number of individuals with severe periodontal disease. There is a general paucity of long-term data on individuals with severe periodontal disease in primary care receiving routine care (which may include specialist periodontist referral). Among the 85 participants with baseline and follow-up data, there were only minimal changes in the mean scores of present teeth (–0.4 teeth), gingival inflammation (+4%), supragingival calculus (+7.5%) and mean clinical probing depths (+0.1 mm). Similarly, there were only minimal changes in participant-reported outcomes: self-efficacy (+0.1 points; 1–7 scale), PBC (+0.2 points; 1–7 scale), attitude (–0.3 points; 1–7 scale) and intention (no change). Interestingly, the patient-centred outcome scores (including QoL) were largely similar to those of the IQuaD trial participants.

The IQuaD cohort study has a number of limitations, including the high attrition rate, with only 59% of participants being available for 3-year clinical follow-up. The recruitment strategy of the IQuaD trial meant that only those individuals who had a BPE score of  $\leq 3$  at their most recent examination appointment were invited to the trial recruitment sessions. Therefore, it is unclear how representative this relatively self-selecting group of participants was, considering that the participants were all regular attenders at their primary care dental practice who had previously shown no signs of severe periodontal disease (as defined by a BPE score of 4).

It would be inappropriate to draw definitive conclusions from the outcomes, but it appears that periodontal condition and participant-centred outcomes did not markedly deteriorate or improve in the 85 IQuaD cohort participants over the course of this 3-year study.

## **Chapter 7** Discussion/conclusions

The IQuaD trial involving regular adult NHS dental attenders (with no or early signs of periodontitis) has shown that, over a 3-year period, scheduling 6-monthly or 12-monthly PIs has no additional beneficial effect (over not providing this treatment unless desired or recommended) on primary clinical (gingival inflammation/bleeding) and patient-centred (self-reported) outcomes and that there is no difference between personalised OHA and routine OHA (current practice). However, patients value, and are willing to pay for, both interventions, with greater financial value placed on PI than on OHA.

The IQuaD trial is the first pragmatic cluster RCT to evaluate PI and OHA in NHS dental practices. These interventions are the most frequently performed treatments in dentistry, with considerable cost to the NHS and society. The aim of this RCT in primary care dental practice was to provide evidence for the benefit or not of these routine treatments on the periodontal health of adults with no or early signs of periodontal disease.

The primary clinical outcome, gingival inflammation/bleeding, is a measure of gingivitis, a recognised precursor of periodontitis, caused by plaque retention, and is reversible with effective plaque removal. The PI in the IQuaD trial involved professional mechanical removal of plaque and calculus. Effective oral hygiene (self-care) involves tooth brushing with appropriate interdental cleaning, which, if undertaken effectively, will be sufficient to prevent or resolve gingivitis.

At baseline, most participants (98%) experienced gingival inflammation/bleeding, with, on average, 33% of sites affected. The prevalence of gingival inflammation/bleeding measured at baseline was similar to that reported in other studies<sup>25</sup> and confirmed that, if effective, the trial interventions had the opportunity/potential to improve periodontal health.

At the 3-year follow-up, there was no evidence of a difference in gingival inflammation/bleeding between those randomised to PI on a 6-monthly basis and those who were randomised to receive no PI (difference 0.87%, 95% CI -1.6% to 3.3%; p = 0.481). Similarly, there was no evidence of a difference between scheduled 6-monthly PI and scheduled 12-monthly PI (difference 0.11%, 95% CI -2.3% to 2.5%; p = 0.929). We are confident in the finding of no clinical benefit for 6-monthly PI over other frequencies of PI because the 95% CIs were small enough to exclude the prespecified clinically important difference in bleeding of 7.5%. Of the secondary clinical outcomes, a difference between groups was found only for calculus, with those participants in the no-PI group having 8% (95% CI 5.4% to 10.7%; p = 0.001) more sites with calculus than the 6-monthly PI group after 3 years. Although there was a statistically significant difference, the clinical significance or relevance of this result has to be questioned because there was no associated improvement in the primary clinical outcome of gingival inflammation/bleeding.

All participants received PI at baseline and were then randomised to be offered this treatment 6-monthly, 12-monthly or not at all for the 3-year duration of the trial. Participants' dental practices were asked to recall their participants for PI in keeping with their randomised allocation; however, a PI could be provided if either the participant requested it or the clinician thought it necessary. The pragmatic nature of the trial meant that scheduling of dental appointments and the intervals between PIs varied, as they do in routine practice. Prior to and since the publication of NICE dental recall guidelines, 66 common practice is to provide a PI at the same time as a dental recall visit and for this to be planned every 6 months. The intention was that participants randomised to the 6-monthly PI group would receive this frequency of care.

The claims recorded in the routine data collected for the IQuaD trial confirmed that the number and interval between PIs varied within groups and demonstrated clear separation in the mean number of PIs between groups. At the 3-year follow-up, there was a threefold difference in the mean number of PIs (1.1, 2.0 and 3.0 for the no-PI, 12-monthly and 6-monthly groups, respectively). National routine data<sup>67</sup> suggest that, in the NHS, the most frequent interval between visits with a PI is 9 months and this reflects the experience of participants in the 6-monthly PI group. The finding that 46% of participants randomised to

the no-PI group did not request, and were not thought to need, this treatment (PI) indicates that almost half of participants were not clinically compromised for the duration of the trial and would not have experienced any benefit had this treatment been provided.

Although participants with moderate to severe periodontal disease (BPE score of 4 or \*) were excluded from the trial, adults with a spectrum of periodontal disease and a wide age range were included. Using the baseline data, these groups were included in subgroup analyses along with dental practice/cluster employment of a hygienist, smoking habit and a post hoc criterion for moderate periodontitis evidenced by four or more pockets with a clinical probing depth of  $\geq 4$  mm. The IQuaD trial found no evidence of statistically significant differences at the 1% level for the prespecified and post hoc subgroup analyses and the primary clinical outcome of gingival inflammation/bleeding. However, the results suggest that there may be some clinical benefit of 6-monthly PI compared with other frequencies for participants with BPE scores of 3 at baseline (difference -4.7% 99% CI -10.5% to 1.1%; p=0.038). Current clinical guidance from the British Society of Periodontology<sup>16,68</sup> recommends providing PI for early signs of periodontitis (BPE score of 3 with bleeding or other risk factors) and these results suggest a possible benefit to such a targeted approach.

The IQuaD trial found no evidence of a difference between participants randomised to personalised OHA and those randomised to routine advice (current practice) for any outcome. Although the 95% CI for the difference in gingival inflammation/bleeding (-2.5%, 95% CI -8.3% to 3.3%; p=0.393) does not exclude the prespecified clinically important difference of 7.5%, it is unlikely that the personalised OHA has any impact > 8%. The results were robust to other adjusted/unadjusted models. Similarly to PI frequency, there was no evidence that any of the subgroup analyses were statistically significant at the 1% level. Although we found no evidence of a difference between the two OHA interventions, we have evidence of intervention fidelity from participant-reported responses in their follow-up questionnaires about desired oral health behaviours such as spit but not rinse, which leads us to believe that the personalised intervention was indeed delivered as intended. Participants having a plan for brushing or flossing better was another variable assessed. The randomised groups (personalised vs. routine) had similar levels of participants reporting to have a plan. This might reflect the fact that the question was asked too late (it would have been around 1 year between the participants making a plan with their clinicians and receiving their first participant annual questionnaire). Regardless, there is no good evidence that clinicians in the personalised arm delivered the intervention in a way that promoted planning of oral hygiene behaviour.

The personalised OHA was designed as a brief intervention (under 10 minutes) appropriate to be delivered in primary care dental practice. OHA is considered integral within a NHS recall/check-up appointment and only rarely attracts an additional fee. The trial intervention was designed 10 years ago using psychological behaviour change models; since that time national clinical guidance documents that provide similar advice have been disseminated in both England<sup>16</sup> and Scotland.<sup>69</sup> Dentists randomised to deliver routine OHA were asked to conform to their current practice. Therefore, one possible explanation of the outcome is that the personalised OHA was similar to current practice and therefore did not result in any additional benefit compared with current practice.

The confidence (self-efficacy) of participants to perform effective oral hygiene (i.e. to clean their teeth and gums regularly and well) was chosen to be the primary patient-centred outcome because gingivitis can be prevented with effective oral hygiene.<sup>34</sup> The IQuaD trial found participants to be moderately confident in performing good oral hygiene and the frequency of PI or type of OHA did not make a difference. The statistically significant difference found between those randomised to receive 6-monthly and 12-monthly PI, favouring 6-monthly PI (difference -0.097, 95% CI -0.188 to 0.006; p = 0.037), is most likely due to the very precise CI. The difference considered clinically significant was one-quarter of a SD around 0.275 in the self-efficacy scale and the CI excludes that value.

For all other clinical and patient-centred outcomes, the IQuaD trial found no evidence to suggest that the frequency of PI or whether or not OHA was personalised made a difference.

The health economics analysis assessed the costs and benefits of alternative policies of OHA with or without PI in terms of costs and benefits (WTP). The analysis focused on whether moving from standard care to an alternative policy increases or decreases welfare. The results showed that there were no statistically significant differences in total NHS costs for any of the policies compared with standard care (6-monthly PI and routine OHA). In terms of point estimates of total mean NHS costs, the policy of no PI and personalised OHA was associated with the lowest NHS costs (£62.42), followed by the policy of no PI and routine OHA (£72.23).

In Scotland, the fee-for-service structure allowed an assessment solely of the costs of periodontal treatments. Relative to standard care, and given the structure of current contracts, a policy of no PI (with routine OHA) would significantly reduce NHS spending on periodontal treatments by £7.47 per person over 2.25 years. A policy of offering routine OHA with 12-monthly PI would save £5.11 per person on periodontal treatments over the same time frame. However, we cannot definitively conclude cost savings to the overall NHS dental budget. To make this statement would require the assumption that the lack of statistical significance in total NHS costs is due to a lack of statistical power and the relatively large SDs around high-cost treatments, rather than any causal effects of disinvesting from PI. Given the lack of evidence of any clinical differences across policies, it could be argued that such an assumption may be valid, and that cost savings could be achieved, at least in the Scottish system, without having an adverse impact on the primary clinical outcome (gingival inflammation/bleeding).

When taking a wider perspective (NHS and participant), the policy of personalised OHA and no PI was less costly than standard care. However, this difference was mainly driven by the lower costs of interdental brushes in England and it is questionable whether or not it is attributable to the policy. The total cost differences between all other policies and standard care were not statistically significant.

The results of the DCE show strong evidence that the general population valued both PI and personalised OHA even when controlling for frequency of bleeding gums and aesthetics (the look and feel of teeth). Owing to the relatively high value placed by the general population on PI and personalised OHA, the policy of 6-monthly PI and personalised OHA was associated with the highest incremental benefit compared with standard care (£61.67). The policy of 12-monthly PI and personalised OHA was also associated with a positive but lower incremental benefit (£19.70) but the difference was not statistically significant.

The results of the CBA showed that the policy of 6-monthly PI with personalised OHA was associated with the largest incremental net benefit compared with standard care [£48 when taking a NHS cost perspective and £68 when taking a wider perspective (NHS and participant)]. The most efficient approach depends on the perspective taken. From a welfare maximisation perspective, the analysis shows that personalised OHA with 6-monthly PI is cost-beneficial, with the greatest incremental net benefit relative to standard care. It should be noted that this finding was driven primarily by differences in benefits rather than NHS-incurred costs.

From a health maximisation perspective, the no-PI approach generates the lowest average costs (though differences are not statistically significant), without detrimentally affecting clinical outcomes. The overall health economic conclusions were robust to a range of sensitivity analyses undertaken to explore the impact of uncertainty in key assumptions on results.

## **Comparisons with other randomised clinical trials**

The Cochrane systematic review<sup>24</sup> of routine scale and polish (PI) for periodontal health in adults (updated in 2013) reported that there was insufficient evidence to determine the effects of routine PI treatment.

Three studies  $^{25-27}$  were identified, one of which was conducted in a similar setting to the IQuaD trial (three NHS general dental practices in England, involving 369 participants). The trial inclusion criteria were less pragmatic than those in the IQuaD trial, with individuals with BPE scores of > 2, diabetes mellitus and those aged > 60 years excluded. Jones *et al.*  $^{25}$  compared PI intervals of 6-monthly PI, 12-monthly PI or 24-monthly PI (no PI throughout the trial). There was no evidence of any clinical differences between these different PI intervals provided for gingival health (gingival inflammation/bleeding). The Cochrane review is currently being updated by two authors of the monograph and will include the results of the IQuaD trial. The updated search of 5 December 2016 identified the IQuaD trial as the only new eligible study to be included in the updated review. When synthesising the data of both trials, the treatment effect is mean difference 0.00 (95% CI -0.03 to 0.03; p = 0.87) in gingival health, as shown in *Figure 17*.

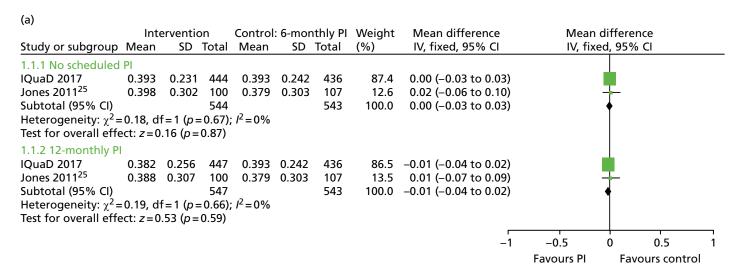
A recent systematic review<sup>23</sup> of psychological approaches to behaviour change for improved plaque control in periodontal management reported benefits of using goal-setting, self-monitoring and planning for improving oral hygiene. A number of the included trials were of short duration, of a non-experimental design and were rated as having a high risk of bias. A meta-analysis was not possible and the results and recommendations should be interpreted with caution. An ongoing Cochrane systematic review of delivering one-to-one oral hygiene in a variety of dental settings has identified a number of trials investigating enhanced oral hygiene techniques (Soldani FA, Young L, Jones K, Walsh T, Clarkson JE. Bradford District Care NHS Foundation Trust. 2017). The intensity of the interventions varied greatly and for a number of trials the risk of bias is rated as unclear or high. We have presented a diagrammatic display of the studies' effects to demonstrate the spread of effect size, direction and lack of CI overlap, highlighting the uncertainty of treatment clinical effectiveness (*Figure 18*).

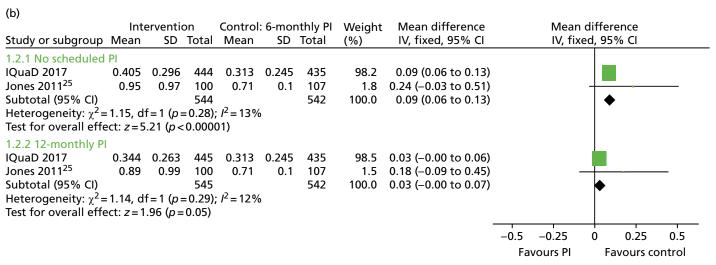
## **Strengths**

The IQuaD trial was a pragmatic trial in primary care dental practice designed to provide evidence for the clinical effectiveness and cost-effectiveness of the most frequently provided treatment in dentistry, PI, and the impact of personalised OHA. Despite the length of time that has elapsed since the trial began, uncertainty still exists for providers of dental care worldwide regarding the benefit of these common components of routine dental care.

Surprisingly, there have been few studies comparing different frequencies of PI, and none that have combined it with different approaches to providing OHA in a setting such as the NHS. Reviews evaluating PI frequency have included studies that involve few providers and recruit only participants with no or very early signs of periodontal disease (BPE scores of 0–2). One review concluded that there was little value in providing PI without OHA; this was largely influenced by the results of a study with an intensive advice programme involving several 45-minute appointments, which does not reflect routine NHS practice.<sup>74</sup>

The strengths of the IQuaD trial include the recruitment and retention of a large number of centres (n=63), with wide representation of geographical, clinical and dental practice characteristics and operating in two different contractual systems in the UK. Recruitment and retention in trials is a frequent challenge, but an achievement for the IQuaD trial was the recruitment of 87% of the potentially eligible patients and the retention at 3 years of 1450 participants with 1327 (71%) providing clinical data. The randomised groups were balanced/similar at baseline and the reasons for loss to follow-up were related to the inability of practices to contact their patients; therefore, we were confident in the robustness of the results regardless of the missing data. The dental behaviour and clinical characteristics of the participants mean that the findings of the IQuaD trial are generalisable to regular dental attenders in the NHS and, therefore, similar third-party funding systems. The cohort study provided long-term data on NHS primary care patients with moderate to severe periodontal disease, which has previously been lacking.





HEALTH TECHNOLOGY ASSESSMENT 2018 VOL. 22 NO. 38

FIGURE 17 Routine PI for periodontal health in adults. (a) Forest plot of comparison 1: PI vs. no PI (control), outcome 1.1 gingivitis at 24 months; and (b) forest plot of comparison 1: PI vs. no PI (control), outcome 1.2 calculus at 24 months. df, degrees of freedom; IV, inverse variance.

FIGURE 18 One-to-one OHA provided in a dental setting for oral health. Forest plot of comparison 1: personalised OHA vs. routine OHA, outcome 1.1 gingivitis. IV, inverse variance.

The pragmatic design of the trial did not prevent fidelity to the interventions with evidence of separation in the frequency of Pls between groups and of the delivery of personalised OHA. Conducting the trial in a primary care NHS setting with a design that involved minimal requirements for the dental practices, blind outcome assessment and access to routine data are strengths. The interventions delivered were not independently observed; however, we believe that they were delivered as in current dental practice and the fidelity information supports this.

The health economic evaluation has a number of strengths. To our knowledge, this trial provides the only evidence worldwide regarding the efficient allocation of NHS resources to the delivery of PI and personalised OHA. Uniquely among clinical trials of dental care interventions, we implement a CBA, using a DCE to capture the value placed on both the processes and outcomes of care by the general population. The values of the general population are most relevant as they provide the funding for health care in a tax-based system such as the NHS. The approach is novel within dentistry and has a number of distinct advantages over traditional cost-effectiveness and cost-utility analyses as it incorporates the simultaneous valuation of multiple processes and outcomes which are relevant to dental care recipients.

In terms of the within-trial analysis, we use best practice methodology, incorporating the most advanced recommendations for analysis of cluster randomised trials together with the appropriate use of missing data models to minimise the potential for bias.

## **Limitations**

As with all evaluations, it is important to acknowledge the limitations of the trial.

A weakness was not collecting detailed information about the reasons for additional Pls. A possible weakness is that gingival inflammation, measured as bleeding on probing, is a measure for which calibration is not possible. The difference in bleeding measures between baseline and follow-up probably reflects assessor's measurement error, but they were equally distributed across practice and participant randomised allocations, avoiding bias between randomised groups. The ICC for the primary clinical measure, bleeding on probing, was higher (0.24) than originally planned for (0.10), and most likely the observed difference was influenced by the 'assessor effect'. The trial was still powered to detect the clinically important difference due to statistical adjustment for other factors such as the baseline measures of bleeding and the minimisation variables.

The different contracts across Scotland and England provided several challenges to the health economic evaluation. First, the differences in costs across the policies were likely to vary across the regions as Pl attracts a separate payment in Scotland while it is included in band 1 treatments in England (which also include clinical check-ups). Second, the different contracts may provide different incentives for providing care. Some evidence suggests that health-care providers have a tendency to supply more services under fee-for-service than under other reimbursement contracts, but, in the IQuaD trial, we found no evidence of this, as average costs were actually higher in England than in Scotland. One could also make an argument that different contracts may have an impact on the private—public mix of services, for example the referral rate for private Pl. These issues potentially limit the direct comparability of Scottish and English cost data. However, this difference did not bias average UK results, as the ratio of English to Scottish practices was similar across the different arms of the trial. Furthermore, although subgroup analyses exploring regional-level resource use, cost and incremental net benefits show substantial variation across regions, the overall recommendations from the CBA remain unchanged.

Although the DCE was a useful method for obtaining WTP estimates for PI, personalised OHA and trial outcomes, it did not provide insights into why individuals value these attributes. Although source of value is not relevant from a CBA perspective, policy-makers may want some insights into this given the current climate of severe financial constraints. The results showed that the public value PI and personalised OHA

even when controlling for self-reported bleeding and aesthetics. There are a number of potential explanations for this finding:

- Respondents may have assumed that PI and personalised OHA were associated with clinical benefits other than bleeding and aesthetics.
- PI and personalised OHA represent another interaction with a dentist or hygienist, which provides an opportunity to identify any dental health issues and to provide reassurance.
- Respondents may simply value PI because that is what they always have (habitual behaviours) and/or their dentist has always recommended that they have a PI.
- Respondents might value having the option of PI and personalised OHA in a dental care package (option value).
- Respondents might want to minimise the regret that they may feel if their dental health gets worse and they did not have PIs or personalised OHA.
- It is possible that the general population value the additional PI service received as part of attending for a band 1 treatment in England. It is possible that withdrawal of PI may have an impact on the attendance for routine check-ups; however, that consequence was not observed in the trial participants when all groups continued to attend for dental check-ups at a similar rate.

Further research is required to explore the reasons behind the positive value attached to PI.

The economic analysis is based on the price paid by the NHS for dental treatments under the current contracts and structures in place in England and Scotland. Thus, our analysis is the most accurate reflection of the current impact on NHS budgets of delivering alternative policies. However, there are limitations associated with this. First, the price may not be an accurate reflection of the cost of time and materials to the dentists of delivering the respective interventions. Owing to contract structures, it is likely that there are substantial differences between price and cost. Second, if the way in which dental practices are paid were to change substantially under new contracts, the results of our costing analyses would need to be considered in the light of such changes. Third, we have not explicitly estimated the opportunity costs of alternative policies (i.e. the benefits forgone from not having those resources available to expand or implement other policies). This is an important consideration, given a fixed NHS budget; however, the issue is rarely explicitly incorporated within economic evaluation frameworks. It should be noted that, for this study, incremental net benefit results were driven by differences in benefits rather than costs, so the notion of opportunity costs is less relevant in this scenario, as NHS costs were similar across policies.

The majority of economic evaluations of health-care interventions tend to extrapolate trial outcomes over a longer time horizon using decision-modelling studies. For this study, we have conducted a CBA alongside the RCT. We have not attempted to extrapolate trial results as there is a paucity of evidence to directly predict the probability of final clinical end points (such as tooth loss or long-term caries), or to determine when such events may happen. This is a limitation of our analysis. However, to attempt such a modelling exercise, given the paucity of available data, could generate misleading conclusions for policy-makers. Furthermore, given that there was no evidence of short-term clinical benefit, one would not expect any differences in long-term tooth loss or caries across different policy options in this population with relatively healthy dentitions. Compared with other areas of health economic evaluation, there is a lack of evidence generally to inform the population of economic decision models in dentistry and further research is required to bridge this gap.

Despite attempting to collect information on the wider NHS perspective, it was found that rates of secondary care consultations exceeded what might be expected in the general population. After a validation exercise, the data were excluded. Reasons for the poor validity may include a participant's difficulty in determining what was meant by the questionnaire text asking to report 'admissions for problems related to your teeth' or issues of recall bias over a full year of recall, or the question may have been misinterpreted, with participants providing details of all contacts with secondary care, rather than dental care-specific attendances. Future studies should explore direct linkage to Scottish Morbidity Reports data in Scotland and Hospital Episode Statistics data in England to obtain more accurate estimates at a

participant level. The impact of omitting these data on the results is likely to be minimal, given that resource use was evenly distributed across groups.

## Patient and public involvement and engagement

Prior to the start of the IQuaD trial, patients were involved with the trial design and provided invaluable feedback on trial recruitment and communication strategies. Patients also contributed to the content and layout of the trial invitation, trial newsletters and the design of patient participant questionnaires. This ensured that trial participants could understand and easily complete these materials. Members of the public also contributed to trial oversight through membership of the TSC, including helping to interpret the trial findings and preparation of the monograph. We found that the most effective feedback was provided in face-to-face meetings with trial staff and patient and public involvement (PPI) personnel. This provided an opportunity for clarification and more comprehensive and constructive feedback. In preparation for oversight committee meetings and the final PMG meeting, the PPI representative found that the opportunity for a pre-meeting session to clarify issues made their participation of these meetings easier. The results were discussed with the Periodontal Advisory Group and Chief Dental Officers of the UK.

## **Generalisability**

The IQuaD trial was designed pragmatically to investigate OHA and PI delivery to NHS patients in dental primary care that did not exhibit moderate or severe periodontitis.

The 63 recruiting dental practices were situated across Scotland and the north-east of England, with 72% of practices employing dental hygienists and 75% having three or more dentists working in the practice.

In the year to 31 March 2014, 84.4% of the population of Scotland was registered with a NHS dentist, 76.4% of whom had attended the dentist in the past 2 years. The ADHS 2009<sup>11</sup> reported that over half of the UK population had attended a dental practice within the last 3 years. The IQuaD trial recruited a total of 1877 participants at their routine dental examination, with 1348 (72%) of these participants recruited across Scotland, including 121 (6%) recruited from the Scottish islands. Over 90% of participants reported having attended a dental practice at least once over the previous 2 years.

At baseline, around 60% of participants reported having received OHA and a PI during their last dental appointment. The mean percentage of sites with gingival inflammation/bleeding was 33%, and 35% of teeth had calculus present. The mean clinical probing depth was 1.8 mm. Two-thirds of participants had BPE scores of  $\leq$  2, and between 10% and 12% of participants in each group had four or more pockets with a clinical probing depth of  $\geq$  4 mm.

We are confident that the practices and participants recruited to the IQuaD trials are a true representation of those adults who are periodontally healthy or have very early periodontal disease and attend NHS dental practices across Scotland and England.

#### **Recommendations for research**

- Research is needed to assess the clinical effectiveness and cost-effectiveness of providing multifaceted periodontal care packages (e.g. OHA, oral care products, PI) in primary dental care for those with periodontitis.
- Research is required to better understand the source of WTP values and the extent to which this is influenced by perceptions and current practice.
- Further research is required to explore the relative value of different data sources for estimating resource use in dentistry including routine data, patient-reported data and practice records.

## **Acknowledgements**

The authors wish to thank Mark Forrest and the programming team at CHaRT; Cynthia Fraser, our information specialist, for assistance with referencing; Moira Swan, who was the dental research nurse and part of the OA team in Newcastle upon Tyne; Louise Campbell for secretarial support and data management; our original statistician in the group, Andy Elders; senior IT manager Gladys Macpherson; senior trial administrator at the TCOD Marilyn Laird; Luke Vale for his involvement with the design of the health economic analysis at the inception of the trial; Maria Dimitrova, who assisted the health economists in the collection of unit costs; staff of the Scottish Primary Care Research Network, who assisted with screening eligible patients at dental practices; staff of the North East Commissioning Support Unit who assisted with research payments to dental practices in the north-east; members of the TMC and Periodontal Advisory Group for their ongoing advice and support of the trial; the independent members of the TSC and DMC; and the staff at recruitment sites who facilitated recruitment, treatment and follow-up of trial participants.

The Health Services Research Unit and the Health Economics Research Unit is core funded by the Chief Scientist Office of the Scottish Government Health and Social Care Directorate.

## **Trial Management Committee**

Paul Averley, Graham Chadwick, Andy Hall, Penny Hodge, Isobel Madden, Wendy McCombes, Nigel Pitts, David Ricketts, Margaret Ross, Martin Tickle and Helen Worthington.

## **Independent members of the Trial Steering Committee**

Tina Halford-McGuff, Dione Macmorran, the late Eleanor Grey, James McCaul and Elizabeth Treasure (chairperson).

#### **Independent members of the Data Monitoring Committee**

Pollyanna Hardy, Peter Robinson and Damien Walmsley (chairperson).

#### **Members of the Periodontal Advisory Committee**

Mary Cullinan, Mark Ide, Ian Needleman, Tim Newton and Greg Seymour.

#### **Recruitment sites**

We would like to thank the staff and participants of the following dental practices:

Ardmillan Dental Practice, Baillieston Dental Care, Bank Street Dental Surgery (formally known as Peacock Dental Surgery), Birch Valley Dental Clinic, Bridge of Don Dental Practice, Bute Dental Surgery, Care Dental – Crieff, Chong Kwan Dental Centre, City Health Clinic, Clark and Watson Family Dental Practice, Clyde Dental Practice, David Gilchrist Dental Surgery, Dental Care Perth Ltd, Dental Plus, Discovery Dental Care, Drumbrae Dental Surgery, Dunbar Dental Practice, Duns Dental Practice, East Neuk Dental Practice, Framwellgate Dental Surgery, Consett, Grange Dental Centre (formally known as Gosforth Dental), Hetton Dental Practice, Invergowrie Dental Practice, JL Barrack Dental Practice, Kenton Lane Dental Practice, King Street Dental Practice (Orkney), Laurencekirk Dental Practice, L C Milton Dental Practice

(Barrhead Dental practice), Lochboisdale Dental Clinic (South Uist), Long and Gilmour Dental Care, Lux Dental (formally known as K H Stirling), Mastrick Dental Centre, Montgomery Street Dental Care, Mr S B Pabary and Associates, Mullins Dental Practice, Mydentist – Dean Road, Mydentist – Front Street (formally known as Front Street Dental Practice), Nepali Dental Practice, Number One Dental Practice, Osborne Dental Practice, Park View Family Dental (formally known as Mr A I Robson and Associates Dental Practice), P A Penney Dental Practice, Perfect Smile Blaydon (formally known as Blaydon Dental Practice), Pickering Dental Care Ltd, Princes Street Dental Practice, Riverview Dental Practice, S. Rankin, Selkirk Dental Practice, Shotley Bridge Dental Care, Silver Dental Practice, Smith and Smith Dental Practice, St Leonards Dental Practice, Stirling Dental Care, Stobswell Dental Practice (formally known as Karolak and Iwanowicz), Stuart Eaborn Dental Practice, Sunderland Road Dental Practice, The Hollies Dental Practice, West End Dental Practice, Whickham Village Dental Practice, Whitburn Cosmetic Dental Clinic (formally known as Whitburn Dental Practice), Wishaw Cross Dental Care and Woodside Dental Practice.

#### Contributions of authors

**Craig R Ramsay** (Professor and Triallist) contributed to the conception and design of the trial, the conduct of the trial, the statistical analysis, the interpretation of the results and the writing/editing of the report.

**Jan E Clarkson** (Professor and Chief Investigator) contributed to the conception and design of the trial, the conduct of the trial, the interpretation of the results and the writing/editing of the report.

**Anne Duncan** (Trials Manager) was responsible for the day-to-day management of the trial, contributed to the interpretation of data and made significant contributions to drafting the monograph.

**Thomas J Lamont** (Clinical Research Fellow) was responsible for the conduct of the trial in Dundee, contributed to recruitment of participants and clinical data collection, contributed to the interpretation of data and made significant contributions to drafting the monograph.

**Peter A Heasman** (Professor and Triallist) contributed to the design of the trial, was responsible for the conduct of the trial in Newcastle upon Tyne, contributed to recruitment of dentists, contributed to the interpretation of data and made significant contributions to writing and editing the report.

**Dwayne Boyers** (Research Fellow) undertook the trial-based health economics analysis and drafted the chapter on health economic methods, *Chapter 3*, and results of the economic analysis, *Chapter 5*, under the supervision of Marjon van der Pol.

**Beatriz Goulão** (Research Fellow and Statistician) conducted the statistical analyses, contributed to drafting the statistical methods section in *Chapter 2* and drafted the clinical effectiveness results in *Chapter 4* under the supervision of Craig R Ramsay.

**Debbie Bonetti** (Health Psychologist) led on the conception and design of the patient-reported self-efficacy questionnaire, the CBQ and the development of the personalised OHA intervention, contributed to analysis and interpretation of behavioural data and CBQ, and contributed to reviewing and approving the final report.

**Rebecca Bruce** (Trials Data Co-ordinator) was responsible for the day-to-day data management of the trial, contributed to interpreting the data and reviewing and approving the final report.

**Jill Gouick** (Research Dental Nurse and OA) was responsible for recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**Lynne Heasman** (Research Hygienist and OA) was responsible for recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**Laura A Lovelock-Hempleman** (Research Hygienist and OA) was responsible for recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**Lorna E Macpherson** (Trials Administrator) was responsible for the day-to-day management of the trial, and contributed to the interpretation of data and to drafting and approving the monograph.

**Giles I McCracken** (Clinical Senior Lecturer and Triallist) contributed to the design of the trial, was responsible for the conduct of the trial in Newcastle upon Tyne, contributed to recruitment of dentists, contributed to the interpretation of data and reviewed and approved the final report.

**Alison M McDonald** (Senior Trials Manager) was responsible for overseeing management of the trial, and contributed to the interpretation of data and to reviewing and approving the monograph.

**Fiona McLaren-Neil** (Trials Administrator) was responsible for overseeing management of the trial, and contributed to the interpretation of data and to reviewing and approving the monograph.

**Fiona E Mitchell** (Research Dental Nurse and OA) was responsible for recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**John DT Norrie** (Professor and Triallist) contributed to the design of the trial, the interpretation of the data and reviewing and approving the monograph.

**Marjon van der Pol** (Professor and Health Economist) contributed to the design of the trial, the interpretation of the data and contributed to the writing and editing of the final report.

**Kirsty Sim** (Research Hygienist and OA) was responsible for the recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**James G Steele** (Professor and Triallist) contributed to the design of the trial, the interpretation of the data and reviewing and approving the monograph.

**Alex Sharp** (Research Dental Nurse and OA) was responsible for the recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**Graeme Watt** (Research Hygienist and OA) was responsible for the recruitment of participants and collection of clinical data outcomes, and contributed to interpreting the data and reviewing and approving the final report.

**Helen V Worthington** (Professor and Triallist) contributed to the design of the trial, the interpretation of the data and reviewing and approving the monograph.

**Linda Young** (Research Manager and Economist) contributed to the design of the trial, the interpretation of the data and reviewing and approving the monograph.

## **Publications**

Heasman PA, Macpherson LE, Haining SA, Breckons M. Clinical research in primary dental care. *BDJ* 2015;**219**:159–163.

## **Data sharing statement**

All available data can be obtained by contacting the corresponding author. See the HTA programme website for further project information.

#### **Patient data**

This work uses data provided by patients and collected by the NHS as part of their care and support. Using patient data is vital to improve health and care for everyone. There is huge potential to make better use of information from people's patient records, to understand more about disease, develop new treatments, monitor safety and plan NHS services. Patient data should be kept safe and secure, to protect everyone's privacy, and it's important that there are safeguards to make sure that it is stored and used responsibly. Everyone should be able to find out about how patient data are used. #datasaveslives You can find out more about the background to this citation here: https://understandingpatientdata.org.uk/data-citation.

## References

- Clarkson JE, Ramsay CR, Averley P, Bonetti D, Boyers D, Campbell L, et al. IQuaD dental trial; improving the quality of dentistry: a multicentre randomised controlled trial comparing oral hygiene advice and periodontal instrumentation for the prevention and management of periodontal disease in dentate adults attending dental primary care. BMC Oral Health 2013;13:58. https://doi.org/ 10.1186/1472-6831-13-58
- 2. Papapanou PN. Epidemiology of periodontal diseases: an update. J Int Acad Periodontol 1999;1:110–16.
- 3. Petersen PE, Ogawa H. The global burden of periodontal disease: towards integration with chronic disease prevention and control. *Periodontol 2000* 2012;**60**:15–39. https://doi.org/10.1111/j.1600-0757.2011.00425.x
- Kassebaum NJ, Bernabé E, Dahiya M, Bhandari B, Murray CJ, Marcenes W. Global burden of severe periodontitis in 1990–2010: a systematic review and meta-regression. *J Dent Res* 2014;93:1045–53. https://doi.org/10.1177/0022034514552491
- 5. Neely AL, Holford TR, Löe H, Anerud A, Boysen H. The natural history of periodontal disease in man. Risk factors for progression of attachment loss in individuals receiving no oral health care. *J Periodontol* 2001;**72**:1006–15. https://doi.org/10.1902/jop.2001.72.8.1006
- 6. Kinane DF, Attström R, European Workshop in Periodontology group B. Advances in the pathogenesis of periodontitis. Group B consensus report of the fifth European Workshop in Periodontology. *J Clin Periodontol* 2005;**32**(Suppl. 6):130–1. https://doi.org/10.1111/j.1600-051X.2005.00823.x
- 7. Chapple IL, Genco R, working group 2 of the joint EFP/AAP workshop. Diabetes and periodontal diseases: consensus report of the Joint EFP/AAP Workshop on Periodontitis and Systemic Diseases. *J Periodontol* 2013;**40**:S106–S12. https://doi.org/10.1111/jcpe.12077
- 8. Linden GJ, Lyons A, Scannapieco FA. Periodontal systemic associations: review of the evidence. *J Clin Periodontol* 2013;**40**(Suppl. 14):S8–S19. https://doi.org/10.1111/jcpe.12064
- 9. Schenkein HA, Loos BG. Inflammatory mechanisms linking periodontal diseases to cardiovascular diseases. *J Periodontol* 2013;**40**(Suppl. 14):S51–S69. https://doi.org/10.1111/jcpe.12060
- 10. Tonetti MS, Van Dyke TE. Periodontitis and atherosclerotic cardiovascular disease: consensus report of the Joint EFP/AAP Workshop on Periodontitis and Systemic Diseases. *J Clin Periodontol* 2013;**40**(Suppl. 14):S24–S9. https://doi.org/10.1111/jcpe.12089
- Steele J, O'Sullivan I. Executive Summary: Adult Dental Health Survey 2009. NHS Digital; 2011.
   URL: http://content.digital.nhs.uk/catalogue/PUB01086/adul-dent-heal-surv-summ-them-exec-2009-rep2.pdf (accessed 1 February 2017).
- Chapple IL, Van der Weijden F, Doerfer C, Herrera D, Shapira L, Polak D, et al. Primary prevention of periodontitis: managing gingivitis. J Clin Periodontol 2015;42(Suppl. 16):S71–6. https://doi.org/ 10.1111/jcpe.12366.
- Tonetti MS, Chapple IL, Jepsen S, Sanz M. Primary and secondary prevention of periodontal and peri-implant diseases: introduction to, and objectives of the 11th European Workshop on Periodontology consensus conference. *J Clin Periodontol* 2015;42(Suppl. 16):S1–S4. https://doi.org/ 10.1111/jcpe.12382
- 14. Tonetti MS, Eickholz P, Loos BG, Papapanou P, van der Velden U, Armitage G, et al. Principles in prevention of periodontal diseases: consensus report of group 1 of the 11th European Workshop on Periodontology on effective prevention of periodontal and peri-implant diseases. *J Clin Periodontol* 2015;**42**(Suppl. 16):S5–11. https://doi.org/10.1111/jcpe.12368

- 15. Key Note. *Market Update Toiletries*. 2014. URL: www.keynote.co.uk/market-update/retail/toiletries (accessed 1 February 2017).
- British Society of Peiodontology (BSP). The Good Practitioner's Guide to Periodontology. Selby: BSP;
   2016. URL: www.bsperio.org.uk/publications/good\_practitioners\_guide\_2016.pdf (accessed
   1 February 2017).
- 17. NHS National Services Scotland. *Primary Care in Dentistry in Scotland, Annual Report 2014/15*. Edinburgh: NHS National Services Scotland; 2016. URL: www.isdscotland.org/Health-Topics/Dental-Care/Publications/2016-11-01/2016-11-01-PrimaryCareDentistry-Report.pdf (accessed 1 February 2017).
- 18. NHS. NHS Dental Statistics for England 2015–16, Annual Report. NHS; 2016. URL: www.content. digital.nhs.uk/catalogue/PUB21701 (accessed 1 February 2017).
- 19. Axelsson P, Nyström B, Lindhe J. The long-term effect of a plaque control program on tooth mortality, caries and periodontal disease in adults. Results after 30 years of maintenance. *J Clin Periodontol* 2004;**31**:749–57. https://doi.org/10.1111/j.1600-051X.2004.00563.x
- 20. Kay E, Locker D. A systematic review of the effectiveness of health promotion aimed at improving oral health. *Community Dent Health* 1998;**15**:132–44.
- 21. Renz A, Ide M, Newton T, Robinson PG, Smith D. Psychological interventions to improve adherence to oral hygiene instructions in adults with periodontal diseases. *Cochrane Database Syst Rev* 2007;**2**:CD005097. https://doi.org/10.1002/14651858.CD005097.pub2
- 22. Sprod AJ, Anderson R, Treasure ET. *Effective Oral Health Promotion: Literature Review. Technical Report 20*. Cardiff: Health Promotion Wales; 1996.
- 23. Newton JT, Asimakopoulou K. Managing oral hygiene as a risk factor for periodontal disease: a systematic review of psychological approaches to behaviour change for improved plaque control in periodontal management. *J Clin Periodontol* 2015;**42**:S36–S46. https://doi.org/10.1111/jcpe.12356
- 24. Worthington HV, Clarkson JE, Bryan G, Beirne PV. Routine scale and polish for periodontal health in adults. *Cochrane Database Syst Rev* 2013;**11**:CD004625. https://doi.org/10.1002/14651858. CD004625.pub4.
- 25. Jones CL, Milsom KM, Ratcliffe P, Wyllie A, Macfarlane TV, Tickle M. Clinical outcomes of single-visit oral prophylaxis: a practice-based randomised controlled trial. *BMC Oral Health* 2011;**11**:35. https://doi.org/10.1186/1472-6831-11-35
- 26. Lightner LM, O'Leary JT, Drake RB, Crump PP, Allen MF. Preventive periodontic treatment procedures: results over 46 months. *J Periodontol* 1971;**42**:555–61. https://doi.org/10.1902/jop.1971.42.9.555
- 27. Listgarten MA, Schifter CC, Laster L. 3-year longitudinal study of the periodontal status of an adult population with gingivitis. *J Clin Periodontol* 1985;**12**:225–38. https://doi.org/10.1111/j.1600-051X.1985.tb00920.x
- 28. Gaunt F, Devine M, Pennington M, Vernazza C, Gwynnett E, Steen N, Heasman P. The cost-effectiveness of supportive periodontal care for patients with chronic periodontitis. *J Clin Periodontol* 2008;**35**(Suppl. 8):67–82. https://doi.org/10.1111/j.1600–051X.2008.01261.x
- 29. Hefti AF, Preshaw PM. Examiner alignment and assessment in clinical periodontal research. *Periodontol 2000* 2012;**59**:41–60. https://doi.org/10.1111/j.1600-0757.2011.00436.x
- 30. Bandura A. Health promotion from the perspective of social cognitive theory. *Psychol Health* 1998;**13**:623–49. https://doi.org/10.1080/08870449808407422
- 31. Gollwitzer PM. Implementation intentions: strong effects of simple plans. *Am Psychol* 1999;**54**:493–503. https://doi.org/10.1037/0003-066X.54.7.493

- 32. Clarkson JE, Young L, Ramsay CR, Bonner BC, Bonetti D. How to influence patient oral hygiene behavior effectively. *J Dent Res* 2009;**88**:933–7. https://doi.org/10.1177/0022034509345627
- 33. Chapple IL, Hill K. Getting the message across to periodontitis patients: the role of personalised feedback. *Int Dent J* 2008;**58**:294–306. https://doi.org/10.1111/j.1875-595X.2008.tb00207.x
- 34. Löe H. The Gingival Index, the Plaque Index and the Retention Index Systems. *J Periodontol* 1967;**38**(Suppl.):610–16. https://doi.org/10.1902/jop.1967.38.6.610
- 35. Slade GD. Derivation and validation of a short-form oral health impact profile. *Community Dent Oral Epidemiol* 1997;**25**:284–90. https://doi.org/10.1111/j.1600-0528.1997.tb00941.x
- 36. Clarkson JE, Turner S, Grimshaw JM, Ramsay CR, Johnston M, Scott A, et al. Changing clinicians' behavior: a randomized controlled trial of fees and education. *J Dent Res* 2008;**87**:640–4. https://doi.org/10.1177/154405910808700701
- 37. NHS Digital. *NHS Payments to General Practice, England, 2014–15*. Leeds: NHS Digital; 2015. URL: http://content.digital.nhs.uk/catalogue/PUB18468 (accessed 1 February 2017).
- 38. NHS Digital. *A Guide to NHS Dental Publications*. Leeds; NHS Digital; 2015. URL: http://content.digital.nhs.uk/catalogue/PUB18129/nhs-dent-stat-eng-14-15-anx5-beg-gui.pdf (accessed 1 February 2017).
- 39. The Scottish Government. *Statement of Dental Remuneration Amendment No 130 Letter*. The Scottish Government; 2015. URL: www.scottishdental.org/library/statement-of-dental-remuneration-amendment-no-130-letter/ (accessed 1 February 2017).
- 40. Steele J. *NHS Dental Services in England: An Independent Review*. NHS; 2009. URL: www.sigwales. org/wp-content/uploads/dh\_101180.pdf (accessed 1 February 2017).
- 41. Lord J, Longworth L, Singh J, Onyimadu O, Fricke J, Bayliss S, Meads C. *Oral Health Guidance Economic Analysis of Oral Health Promotion Approaches for Dental Teams*. Birmingham & Brunel Consortium External Assessment Centre; 2015. URL: www.nice.org.uk/guidance/NG30/documents/oral-health-promotion-approaches-for-dental-teams-health-economic-analysis2 (accessed 1 February 2017).
- 42. National Institute for Health and Care Excellence (NICE). *Guide to the Methods of Technology Appraisal. PMG9*. London: NICE; 2013. URL: www.nice.org.uk/article/pmg9/resources/non-guidance-guide-to-the-methods-of-technology-appraisal-2013-pdf (accessed 1 February 2017).
- 43. Department of Health and Social Care. *NHS Reference Costs 2014 to 2015*. London: Department of Health and Social Care; 2015. URL: www.gov.uk/government/publications/nhs-reference-costs-2014-to-2015 (accessed 1 February 2017).
- 44. Information Services Division (ISD) Scotland. *Costs Book Tables 2015*. Edinburgh: ISD Scotland; 2015. URL: www.isdscotland.org/Health-Topics/Finance/Costs/File-Listings-2015.asp (accessed 1 February 2017).
- 45. Office for National Statistics (ONS). *Annual Survey of Hours and Earnings: 2015 Provisional Results*. London: ONS; 2015. URL: www.ons.gov.uk/employmentandlabourmarket/peopleinwork/earningsandworkinghours/bulletins/annualsurveyofhoursandearnings/2015provisionalresults (accessed 1 February 2017).
- 46. Boox. *The Boox Report: Analysis of the UK's Self-employed Workforce*. Boox; 2014. URL: www.boox.co.uk/wp-content/uploads/2014/01/The-Boox-Report-2014-Download.pdf (accessed 1 February 2017).
- 47. The Campbell and Cochrane Economics Methods Group (CCEMG) and the Evidence for Policy and Practice Information and Co-ordinating Centre (EPPI-Centre). *Cost Converter*. London: EPPI Centre, Social Science Research Unit, University College London; 2016. URL: http://eppi.ioe.ac.uk/costconversion/ (accessed 1 February 2017).

- 48. Office for National Statistics (ONS). *Self-employed Workers in the UK 2014*. London: ONS; 2014. URL: http://webarchive.nationalarchives.gov.uk/20160105160709/http://www.ons.gov.uk/ons/dcp171776\_374941.pdf (accessed 1 February 2017).
- 49. Her Majesty's Revenue and Customs (HMRC). *Expenses and Benefits: Business Travel Mileage for Employees' Own Vehicles*. London: HMRC; 2016. URL: www.gov.uk/expenses-and-benefits-business-travel-mileage/rules-for-tax (accessed 1 February 2017).
- 50. Department of Health and Social Care. *NHS Pay Review Body. Review for 2012*. London: Department of Health and Social Care; 2012. URL: www.gov.uk/government/uploads/system/uploads/attachment\_data/file/215410/dh\_130332.pdf (accessed 1 February 2017).
- 51. Transport Analysis Guidance (TAG). WebTAG: TAG Data Book, July 2016. UK Department of Transport; 2016. URL: www.gov.uk/government/publications/webtag-tag-data-book-july-2016 (accessed 1 February 2017).
- 52. Gomes M, Ng ES, Grieve R, Nixon R, Carpenter J, Thompson SG. Developing appropriate methods for cost-effectiveness analysis of cluster randomised trials. *Med Decis Making* 2012;**32**:350–61. https://doi.org/10.1177/0272989X11418372
- 53. Gomes M, Grieve R, Nixon R, Edmunds WJ. Statistical methods for cost-effectiveness analyses that use data from cluster randomized trials: a systematic review and checklist for critical appraisal. *Med Decis Making* 2012;**32**:209–20. https://doi.org/10.1177/0272989X11407341
- 54. Gomes M, Grieve R, Nixon R, Ng ES, Carpenter J, Thompson SG. Methods for covariate adjustment in cost-effectiveness analysis that use cluster randomised trials. *Health Econ* 2012;**21**:1101–18. https://doi.org/10.1002/hec.2812
- 55. Gomes M, Diaz-Ordaz K, Grieve R, Kenward MG. Multiple imputation methods for handling missing data in cost-effectiveness analyses that use data from hierarchical studies: an application to cluster randomized trials. *Med Decis Making* 2013;**33**:1051–63. https://doi.org/10.1177/0272989X13492203
- 56. Faria R, Gomes M, Epstein D, White IR. A guide to handling missing data in cost-effectiveness analysis conducted within randomised controlled trials. *PharmacoEconomics* 2014;**32**:1157–70. https://doi.org/10.1007/s40273-014-0193-3
- 57. Rubin DB. Multiple Imputation for Nonresponse in Surveys. New York, NY: John Wiley & Sons; 2004.
- 58. Johnson FR, Lancsar E, Marshall D, Kilambi V, Muhlbacher A, Regier DA, et al. Constructing experimental designs for discrete-choice experiments: report of the ISPOR conjoint analysis experimental design good research practices task force. Value Health 2013;16:3–13. https://doi.org/10.1016/j.jval.2012.08.2223
- 59. Bridges JFP, Hauber AB, Marshall D, Lloyd A, Prosser LA, Regier DA, et al. Conjoint analysis applications in health a checklist: a report of the ISPOR Good Research Practices for Conjoint Analysis Task Force. Value Health 2011;14:403–13. https://doi.org/10.1016/j.jval.2010.11.013
- 60. McFadden D. Conditional Logit Analysis of Qualitative Choice Behaviour. In Zarembka P, editor. *Frontiers in Econometrics*. New York, NY: Academic Press; 1974.
- 61. McIntosh E. Using discrete choice experiments within a cost-benefit analysis framework: some considerations. *PharmacoEconomics* 2006;**24**:855–68. https://doi.org/10.2165/00019053-200624090-00004
- 62. NHS Digital. *Hospital Outpatient Activity 2014-15*. London: NHS Digital; 2015. URL: https://digital. nhs.uk/catalogue/PUB19608 (accessed 1 February 2017).
- 63. NHS Digital. *Hospital Episode Statistics, Admitted Patient Care England, 2014–15.* London: NHS Digital; 2015. URL: https://digital.nhs.uk/catalogue/PUB19124 (accessed 1 February 2016).

- 64. Office for National Statistics, National Records of Scotland, Northern Ireland Statistics & Research Agency. 2011 Census Aggregate Data. UK Data Service Census Support; 2016. URL: https://discover.ukdataservice.ac.uk/doi/2011-census-aggregate (accessed 1 February 2017).
- 65. Clark MD, Determann D, Petrou S, Moro D, de Bekker-Grob EW. Discrete choice experiments in health economics: a review of the literature. *PharmacoEconomics* 2014;**32**:883–902. https://doi.org/10.1007/s40273-014-0170-x
- 66. National Institute for Health and Care Excellence (NICE). *Dental Checks: Intervals Between Oral Health Reviews. Clinical Guideline* [CG19]. London: NICE; 2014. URL: www.nice.org.uk/guidance/cg19 (accessed 1 February 2017).
- 67. Tilley CJ, Chalkley MJ. Measuring access to health services: General Dental Services in Scotland. Br Dent J 2005;**199**:599–601. https://doi.org/10.1038/sj.bdj.4812905
- 68. British Society of Periodontology (BSP). *Basic Periodontal Examination (BPE)*. Selby: BSP; 2016. URL: www.bsperio.org.uk/publications/downloads/94\_154250\_bpe-2016-po-v5-final-002.pdf (accessed 1 February 2017).
- 69. Scottish Dental Clinical Effectiveness Programme (SDCEP). Prevention and Treatment of Periodontal Diseases in Primary Care: Dental Clinical Guidance. Dundee: SDCEP; 2014. URL: www.sdcep.org.uk/wp-content/uploads/2015/01/SDCEP+Periodontal+Disease+Full+Guidance.pdf (accessed 1 February 2017).
- 70. Jönsson B, Lindberg P, Oscarson N, Ohrn K. Improved compliance and self-care in patients with periodontitis—a randomized control trial. *Int J Dent Hyg* 2006;**4**:77–83. https://doi.org/10.1111/j.1601-5037.2006.00175.x
- 71. Jönsson B, Ohrn K, Oscarson N, Lindberg P. The effectiveness of an individually tailored oral health educational programme on oral hygiene behaviour in patients with periodontal disease: a blinded randomized-controlled clinical trial (one-year follow-up). *J Clin Periodontol* 2009;**36**:1025–34. https://doi.org/10.1111/j.1600-051X.2009.01453.x
- 72. López-Jornet P, Fabio CA, Consuelo RA, Paz AM. Effectiveness of a motivational–behavioural skills protocol for oral hygiene among patients with hyposalivation. *Gerodontology* 2012;**31**:288–95. https://doi.org/10.1111/ger.12037
- 73. Soderholm G, Nobreus N, Attstrom R, Egelberg J. Teaching plaque control. I. A five-visit versus a two-visit program. *J Clin Periodontol* 1982;**9**:203–13. https://doi.org/10.1111/j.1600-051X.1982. tb02060.x
- 74. Needleman I, Nibali L, Di Iorio A. Professional mechanical plaque removal for prevention of periodontal diseases in adults systematic review update. *J Clin Periodontol* 2015;**42**(Suppl. 16):S12–35. https://doi.org/10.1111/jcpe.12341
- 75. NHS Business Services Authority. *FP17 Processing and Payments*. URL: www.nhsbsa.nhs.uk/fp17-processing-and-payments (accessed 1 February 2017).
- 76. Dyer TA, Owens J, Robinson PG. What matters to patients when their care is delegated to dental therapists? *Br Dent J* 2013;**214**:E17. https://doi.org/10.1038/sj.bdj.2013.275
- 77. Sonneveld RE, Brands WG, Bronkhorst EM, Welie JV, Truin GJ. Patients' priorities in assessing organisational aspects of a general dental practice. *Int Dent J* 2013;**63**:30–8. https://doi.org/10.1111/idj.12001
- 78. Sonneveld RE, Wensing M, Bronkhorst EM, Truin GJ, Brands WG. The estimation of patients' views on organisational aspects of a general dental practice by general dental practitioners: a survey study. *BMC Health Serv Res* 2011;**11**:263. https://doi.org/10.1186/1472-6963-11-263

- 79. Gussy M, Dickson-Swift V, Adams J. A scoping review of qualitative research in peer-reviewed dental publications. *Int J Dent Hyg* 2013;**11**:174–9. https://doi.org/10.1111/idh.12008
- 80. Guzeldemir E, Toygar HU, Cilasun U. Pain perception and anxiety during scaling in periodontally healthy subjects. *J Periodontol* 2008;**79**:2247–55. https://doi.org/10.1902/jop.2008.080152
- 81. Stenman J, Hallberg U, Wennstrom JL, Abrahamsson KH. Patients' attitudes towards oral health and experiences of periodontal treatment: a qualitative interview study. *Oral Health Prev Dent* 2009;**7**:393–401.
- 82. Mussard J, Ashley FA, Newton JT, Kendall N, Crayford TJ. What do you think of your dentist? A dental practice assessment questionnaire. *J Eval Clin Pract* 2008;**14**:181–4. https://doi.org/10.1111/j.1365-2753.2006.00826.x
- 83. Ohrn K, Hakeberg M, Abrahamsson KH. Dental beliefs, patients' specific attitudes towards dentists and dental hygienists: a comparative study. *Int J Dent Hyg* 2008;**6**:205–13. https://doi.org/10.1111/j.1601-5037.2008.00300.x

# **Appendix 1** Clinical effectiveness outcomes

## **Section 1: methods for computing patient-reported outcomes**

Table 33 describes the calculations used to produce the outcomes in the CBQ at baseline.

Table 34 describes the calculations used to produce the outcomes in the CBQ at follow-up.

*Table 35* describes the calculations used to produce the patient-reported outcomes. All outcomes were calculated for each year of the patient annual questionnaire (baseline and years 1, 2 and 3).

TABLE 33 Calculations for outcomes in the CBQ at baseline

Outcome	Questionnaire item(s)	Scoring
Self-efficacy	Q1 (a) to (g)	Mean of all Q1
Attitude	Q2 (a) to (d)	Mean of: $2(a) + 2(b) + 2(c) + 2(d) + 4(a) + 4(b) + 4(c)R^* + 3(a) + 3(b) + 3(c) + 3($
	Q3 (a) to (d)	3(d) + 5(a) + 5(b) + 5(c)R*
	Q4 (a) to (c)	
	Q5 (a) to (c)	
PBC	Q4 (d), Q5 (d)	Mean of: $4(d) + 9(a) + 9(b) + 9(c) + 9(d) + 9(e) + 9(f) + 9(g) + 5(d)$
	Q9 (a) to (f)	
Intention	Q7 (a) to (c)	Mean of: 7(a) + 7(b)
Subjective norm	Q11 (a) to (c)	Mean of: subjective norm advice + subjective norm treat
	Q12 (a) to (c)	Subjective norm advice = mean of $11(a) \times 13(a) + 11(b) \times 13(b) + 11(c) \times 13(c)$
	Q13 (a) to (c)	Subjective norm treat = mean of $12(a) \times 13(a) + 12(b) \times 13(b) + 12(c) \times 13(c)$
Has a plan to give OHA	Q8	8(a)
Has a plan to give Pl		8(b)
R*, reverse score.		

TABLE 34 Calculations for outcomes in the CBQ at follow-up

Outcome	Questionnaire item(s)	Scoring
Self-efficacy	Follow-up CBQ	Mean of all Q1 items
	Q1 (a) to (g)	
Attitude	Follow-up CBQ	Mean of: $2(a) + 2(b) + 2(c) + 2(d) + 4(a) + 4(b) + 4(c)R^* + 3(a) + 3(b) + 3(c) + 3($
	Q2 (a) to (d)	3(d) + 5(a) + 5(b) + 5(c)R*
	Q3 (a) to (d)	
	Q4 (a) to (c)	
	Q5 (a) to (c)	

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

TABLE 34 Calculations for outcomes in the CBQ at follow-up (continued)

Outcome	Questionnaire item(s)	Scoring
PBC	Follow-up CBQ	Mean of: $4(d) + 9(a) + 9(b) + 9(c) + 9(d) + 9(e) + 9(f) + 9(g) + 5(d) + 10(a) + 40(b) + 40(b$
	Q4 (d)	10(b) + 10(c) + 10(d) + 10(e) + 10(f) + 10(g)
	Q5 (d)	
	Q6 (d)	
	Q9 (a) to (g)	
	Q10 (a) to (g)	
Intention	Follow-up CBQ	Mean of: $7(a) + 7(b)$
	Q7 (a) to (b)	
Subjective norm	Follow-up CBQ	Mean of: subjective norm advice + subjective norm treat
	Q11 (a) to (c)	Subjective norm advice = mean of 11(a) $\times$ 14(a) + 11(b) $\times$ 14(b) + 11(c) $\times$ 14(c)
	Q12 (a) to (c)	Subjective norm treat = mean of $12(a) \times 14(a) + 12(b) \times 14(b) + 12(c) \times 14(c)$
	Q14 (a) to (c)	
Has a plan to give OHA	Follow-up CBQ	8(a)
Has a plan to give Pl	Q8	8(b)
R*, reverse score.		

Outcome	Questionnaire item(s)	Scoring
Self-efficacy	Section 3	Mean of all Q1 items
	Q1 (a) to (g)	
PBC	Section 3	Mean of: $2(a)R^* + 2(b) + 2(c) + 2(d) + 2(e) + 3(a) + 3(b) + 3(c)$
	Q2 (a) to (e)	
	Q5 (a) to (c)	
Subjective norm	Section 3	Mean of: $4(a)R^* + 4(b) + 4(f)$
	Q4 (a), (b) and (f)	
Attitude	Section 3	Mean of: $3(a)R^* + 3(b) + 3(c)R^* + 3(d)R^* + 3(e)R^* + 3(f)R^* + 3(g)R^* + 5(d)R^*$
	Q3 (a) to (f)	+ 5(e)R*
	Q5 (d) and (e)	
Behaviour	Section 2	Q1 + Q2 + Q8 or $Q12$ (the highest of the two)
	Q1, Q2, Q8 and Q12	
Intention	Section 2	Q4 + Q5 + Q10
	Q4, Q5 and Q10	
OHIP	Section 4	According to instrument instructions
	Q1-14	

## Section 2: participant dental characteristics (years 1 and 2)

Section 2 shows the participant dental characteristics at follow-up at year 1 (Table 36) and at year 2 (Table 37).

TABLE 36 Participant dental characteristics at year 1

	Randomised gr	oup, <i>n</i> (%)			
	PI			ОНА	
Participant dental characteristics	No ( <i>N</i> = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 860)
Date of last visit to the dental practice (yea	ars ago)				
< 1	449 (72)	445 (71)	458 (73)	715 (71)	637 (74)
1–2	23 (4)	23 (4)	4 (1)	31 (3)	19 (2)
> 2	_	1 (0)	1 (0)	_	2 (0)
Missing	151 (24)	156 (25)	163 (26)	262 (26)	208 (24)
Do you think of yourself as					
A regular attendee	452 (73)	446 (71)	457 (73)	718 (71)	637 (74)
Someone who sees a dentist when in pain or trouble	20 (3)	22 (4)	6 (1)	28 (3)	20 (2)
Missing	151 (24)	157 (25)	163 (26)	262 (26)	209 (24)
In the last 12 months, did you receive OH	Δ?				
Yes	343 (55)	378 (60)	406 (65)	635 (63)	492 (57)
No	124 (20)	88 (14)	51 (8)	103 (10)	160 (18)
Missing	156 (25)	159 (25)	169 (27)	270 (27)	214 (25)
By whom?					
Dentist	241 (39)	233 (37)	209 (33)	395 (39)	288 (33)
Hygienist	61 (10)	89 (14)	110 (18)	132 (13)	128 (15)
Both	43 (7)	59 (9)	89 (14)	116 (12)	75 (9)
Missing	278 (45)	244 (39)	218 (35)	365 (36)	375 (43)
In the last 12 months, did you receive a sc	ale and polish?				
Yes	269 (43)	353 (56)	434 (69)	575 (57)	481 (56)
No	201 (32)	115 (18)	27 (4)	172 (17)	171 (20)
Missing	153 (25)	157 (25)	165 (26)	261 (26)	214 (25)
By whom?					
Dentist	180 (29)	219 (35)	253 (40)	354 (35)	298 (34)
Hygienist	79 (13)	120 (19)	154 (25)	189 (19)	164 (19)
Both	6 (1)	12 (2)	22 (4)	24 (2)	16 (2)
Missing	358 (57)	274 (44)	197 (31)	441 (44)	388 (45)
Smoked in the last 12 months?					
Yes	66 (11)	77 (12)	70 (11)	109 (11)	104 (12)
No	407 (65)	394 (63)	392 (63)	642 (64)	551 (64)
Missing	150 (24)	154 (25)	164 (26)	257 (25)	211 (24)

TABLE 36 Participant dental characteristics at year 1 (continued)

	Randomised group, <i>n</i> (%)						
	PI	PI			ОНА		
Participant dental characteristics	No ( <i>N</i> = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 860)		
What type of toothbrush do you normally use?							
Manual	283 (45)	280 (45)	266 (42)	443 (44)	386 (45)		
Electric	168 (27)	166 (27)	167 (27)	271 (27)	230 (27)		
Do not use brush	21 (3)	26 (4)	28 (4)	38 (4)	37 (4)		
Missing	151 (24)	153 (24)	165 (26)	256 (25)	213 (25)		
How often do you prefer to have a scale and polish?							
Never	17 (3)	14 (2)	7 (1)	22 (2)	16 (2)		
Once every 2 years	19 (3)	15 (2)	7 (1)	23 (2)	18 (2)		
Once a year	109 (17)	127 (20)	70 (11)	149 (15)	157 (18)		
Twice a year	233 (37)	213 (34)	266 (42)	398 (39)	314 (36)		
Three times a year	39 (6)	55 (9)	46 (7)	74 (7)	66 (8)		
Four times a year	42 (7)	37 (6)	50 (8)	68 (7)	61 (7)		
More often	10 (2)	8 (1)	11 (2)	13 (1)	16 (2)		
Missing	154 (25)	156 (25)	169 (27)	261 (26)	218 (25)		

TABLE 37 Participant dental characteristics at year 2

	Randomised group, n (%)						
	PI			ОНА			
Participant dental characteristics	No (N = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 860)		
Date of last visit to the dental practice (years ago)							
< 1	400 (64)	408 (65)	400 (64)	645 (64)	563 (65)		
1–2	22 (4)	20 (3)	3 (0)	23 (2)	22 (3)		
> 2	5 (1)	-	2 (0)	4 (0)	3 (0)		
Missing	196 (31)	197 (32)	221 (35)	336 (33)	278 (32)		
Do you think of yourself as							
A regular attendee	411 (66)	403 (64)	399 (64)	642 (64)	571 (66)		
Someone who sees a dentist when in pain or trouble	17 (3)	25 (4)	5 (1)	28 (3)	19 (2)		
Missing	195 (31)	197 (32)	222 (35)	338 (34)	276 (32)		
In the last 12 months, did you receive OHA?							
Yes	302 (48)	346 (55)	343 (55)	550 (55)	441 (51)		
No	123 (20)	79 (13)	64 (10)	121 (12)	145 (17)		
Missing	198 (32)	200 (32)	219 (35)	337 (33)	280 (32)		

TABLE 37 Participant dental characteristics at year 2 (continued)

	Randomised group, n (%)					
	PI			ОНА		
Participant dental characteristics	No ( <i>N</i> = 623)	12-monthly ( <i>N</i> = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 860)	
By whom?						
Dentist	226 (36)	221 (35)	189 (30)	368 (37)	268 (31)	
Hygienist	37 (6)	73 (12)	90 (14)	92 (9)	108 (12)	
Both	32 (5)	48 (8)	61 (10)	87 (9)	54 (6)	
Missing	328 (53)	283 (45)	286 (46)	461 (46)	436 (50)	
In the last 12 months, did you receive a	scale and polish?					
Yes	219 (35)	342 (55)	370 (59)	494 (49)	437 (50)	
No	204 (33)	83 (13)	37 (6)	172 (17)	152 (18)	
Missing	200 (32)	200 (32)	219 (35)	342 (34)	277 (32)	
By whom?						
Dentist	153 (25)	213 (34)	216 (35)	317 (31)	265 (31)	
Hygienist	51 (8)	119 (19)	139 (22)	155 (15)	154 (18)	
Both	6 (1)	8 (1)	12 (2)	16 (2)	10 (1)	
Missing	413 (66)	285 (46)	259 (41)	520 (52)	437 (50)	
Smoked in the last 12 months?						
Yes	56 (9)	47 (8)	50 (8)	78 (8)	75 (9)	
No	370 (59)	380 (61)	355 (57)	592 (59)	513 (59)	
Missing	197 (32)	198 (32)	221 (35)	338 (34)	278 (32)	
What type of toothbrush do you norma	illy use?					
Manual	255 (41)	233 (37)	218 (35)	369 (37)	337 (39)	
Electric	148 (24)	165 (26)	166 (27)	265 (26)	214 (25)	
Do not use brush	24 (4)	29 (5)	22 (4)	36 (4)	39 (5)	
Missing	196 (31)	198 (32)	220 (35)	338 (34)	276 (32)	
How often do you prefer to have a scale	e and polish?					
Never	14 (2)	12 (2)	10 (2)	20 (2)	16 (2)	
Once every 2 years	24 (4)	7 (1)	7 (1)	22 (2)	16 (2)	
Once a year	98 (16)	120 (19)	59 (9)	137 (14)	140 (16)	
Twice a year	210 (34)	204 (33)	225 (36)	351 (35)	288 (33)	
Three times a year	30 (5)	40 (6)	48 (8)	68 (7)	50 (6)	
Four times a year	39 (6)	34 (5)	38 (6)	50 (5)	61 (7)	
More often	6 (1)	5 (1)	13 (2)	14 (1)	10 (1)	
Missing	202 (32)	203 (32)	226 (36)	346 (34)	285 (33)	

## Section 3: clinical and patient-reported outcomes by year of follow-up

Section 3 shows descriptive data for the clinical (*Tables 38* and *39*) and patient-reported (*Tables 40* and *41*) outcomes.

**TABLE 38** Attendance rates by randomised allocation

	Randomised group, n (%)						
	PI		ОНА				
Attended 3-year examination	No ( <i>N</i> = 623)	12-monthly ( <i>N</i> = 625)	6-monthly ( <i>N</i> = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)		
	444 (71)	447 (72)	436 (70)	712 (71)	615 (71)		

TABLE 39 Clinical outcomes for participants who attended clinical follow-up only

		Randomised group, mean (SD), <i>n</i>						
	Time	PI		ОНА				
Clinical outcomes	point	No	12-monthly	6-monthly	Personalised	Routine		
Present teeth	Baseline	23.4 (4.7), 444	23.5 (4.5), 447	23.8 (4.4), 436	23.4 (4.7), 712	23.7 (4.3), 615		
	Year 3	23.6 (4.8), 444	23.6 (4.6), 447	23.9 (4.5), 436	23.5 (4.8), 712	23.8 (4.3), 615		
% of sites bleeding	Baseline	32.8 (22.8), 444	31.5 (23.7), 447	32.8 (22.9), 436	33.7 (23.1), 712	30.8 (23.1), 615		
	Year 3	39.3 (23.1), 444	38.2 (25.6), 447	39.3 (24.2), 436	39.2 (23.8), 712	38.7 (24.9), 615		
% of teeth with calculus present	Baseline	35.0 (26.0), 444	35.9 (27.4), 447	33.3 (25.3), 436	33.3 (25.1), 712	36.4 (27.4), 615		
	Year 3	40.5 (29.6), 444	34.4 (26.3), 445	31.3 (24.5), 435	35.7 (27.1), 711	35.1 (27.2), 613		
Mean clinical pocket depth (mm)	Baseline	1.8 (0.3), 444	1.8 (0.3), 447	1.8 (0.3), 436	1.8 (0.3), 712	1.8 (0.3), 615		
	Year 3	1.9 (0.3), 444	1.9 (0.3), 446	1.9 (0.3), 436	1.9 (0.3), 712	1.9 (0.3), 614		
Mean clinical pocket depth	Baseline	47 (11)	58 (13)	47 (11)	89 (13)	63 (10)		
(four sites with clinical pocket depth of $\geq 4$ mm), $n$ (%)	Year 3	63 (14)	62 (14)	59 (14)	98 (14)	86 (14)		

TABLE 40 Questionnaire response rates by randomised group

	Randomised group	Randomised group, n (%)									
	PI			ОНА							
Year	No ( <i>N</i> = 623)	12-monthly (N = 625)	6-monthly ( <i>N</i> = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)						
1	476 (76)	472 (76)	463 (74)	753 (75)	658 (76)						
2	430 (69)	428 (68)	408 (65)	674 (67)	592 (68)						
3	494 (79)	486 (78)	472 (75)	780 (77)	672 (78)						

TABLE 41 Patient-reported outcomes of those who replied to the follow-up questionnaire at 3 years

		Randomised group, mean (SD), <i>n</i>						
		PI			ОНА			
Outcomes	Time point	No	12-monthly	6-monthly	Personalised	Routine		
Self-efficacy	Baseline	5.2 (1.1), 490	5.2 (1.2), 477	5.2 (1.2), 463	5.2 (1.2), 766	5.2 (1.2), 664		
	Year 1	5.2 (1.1), 418	5.1 (1.1), 417	5.2 (1.1), 407	5.2 (1.1), 665	5.2 (1.1), 577		
	Year 2	5.3 (1.1), 398	5.2 (1.2), 394	5.3 (1.1), 386	5.2 (1.1), 632	5.3 (1.2), 546		
	Year 3	5.3 (1.1), 493	5.2 (1.1), 485	5.3 (1.1), 472	5.3 (1.1), 780	5.2 (1.1), 670		
PBC	Baseline	4.4 (1.2), 486	4.4 (1.2), 474	4.5 (1.2), 462	4.5 (1.2), 764	4.4 (1.2), 658		
	Year 1	4.4 (1.2), 418	4.3 (1.2), 417	4.4 (1.2), 409	4.4 (1.2), 666	4.3 (1.2), 578		
	Year 2	4.4 (1.2), 395	4.5 (1.3), 394	4.3 (1.3), 386	4.4 (1.3), 630	4.4 (1.2), 545		
	Year 3	4.4 (1.3), 412	4.4 (1.2), 411	4.4 (1.2), 397	4.4 (1.3), 660	4.4 (1.2), 560		
Attitude	Baseline	5.8 (1.2), 489	5.9 (1.1), 475	5.7 (1.3), 463	5.8 (1.2), 767	5.8 (1.2), 660		
	Year 1	5.4 (1.6), 418	5.5 (1.5), 417	5.7 (1.4), 409	5.5 (1.5), 666	5.5 (1.5), 578		
	Year 2	5.6 (1.4), 397	5.6 (1.5), 394	5.6 (1.4), 386	5.5 (1.5), 632	5.6 (1.4), 545		
	Year 3	5.5 (1.6), 414	5.6 (1.4), 410	5.6 (1.5), 397	5.5 (1.5), 659	5.5 (1.5), 562		
Subjective norm	Baseline	5.3 (1.1), 479	5.3 (1.2), 469	5.3 (1.1), 453	5.3 (1.1), 749	5.2 (1.1), 652		
	Year 1	4.9 (1.2), 412	4.9 (1.2), 416	5.0 (1.2), 408	4.9 (1.2), 663	4.9 (1.2), 573		
	Year 2	4.9 (1.1), 392	4.9 (1.1), 390	4.9 (1.1), 384	4.9 (1.1), 624	4.9 (1.1), 542		
	Year 3	4.9 (1.2), 409	4.9 (1.2), 409	4.9 (1.1), 396	4.9 (1.2), 654	4.9 (1.1), 560		
Behaviour score	Baseline	4.7 (1.7), 485	4.7 (1.8), 476	4.8 (1.7), 463	4.6 (1.7), 762	4.9 (1.8), 662		
	Year 1	5.0 (1.7), 418	5.1 (1.7), 416	5.2 (1.7), 406	5.0 (1.7), 665	5.2 (1.7), 575		
	Year 2	5.1 (1.6), 398	5.1 (1.7), 396	5.3 (1.6), 383	5.1 (1.6), 632	5.2 (1.6), 545		
	Year 3	5.1 (1.7), 412	5.1 (1.7), 411	5.4 (1.6), 395	5.1 (1.7), 658	5.2 (1.6), 560		
Intention score	Baseline	5.5 (1.8), 445	5.4 (1.8), 451	5.5 (1.8), 418	5.3 (1.8), 707	5.6 (1.7), 607		
	Year 1	5.7 (1.7), 400	5.6 (1.7), 401	5.7 (1.7), 393	5.7 (1.7), 642	5.6 (1.7), 552		
	Year 2	5.6 (1.7), 387	5.6 (1.7), 377	5.7 (1.7), 373	5.6 (1.7), 614	5.7 (1.7), 523		
	Year 3	5.6 (1.7), 403	5.6 (1.7), 390	5.8 (1.7), 384	5.7 (1.7), 636	5.7 (1.7), 541		
OHIP	Baseline	5.6 (7.0), 476	6.0 (7.3), 466	6.1 (7.5), 456	5.8 (7.4), 752	6.1 (7.2), 646		
	Year 1	4.9 (6.5), 415	5.6 (7.0), 407	5.0 (6.6), 395	4.9 (6.5), 652	5.4 (7.0), 565		
	Year 2	4.7 (6.5), 389	5.4 (7.0), 387	4.4 (6.0), 381	4.8 (6.5), 619	5.0 (6.6), 538		
	Year 3	5.2 (6.8), 408	5.3 (6.7), 399	4.9 (6.7), 387	4.8 (6.5), 641	5.5 (6.9), 553		
Has a plan to either	Baseline	0.3 (0.4), 494	0.2 (0.4), 486	0.3 (0.4), 472	0.3 (0.4), 780	0.2 (0.4), 672		
brush or floss better	Year 1	0.2 (0.4), 494	0.2 (0.4), 486	0.2 (0.4), 472	0.2 (0.4), 780	0.2 (0.4), 672		
	Year 2	0.2 (0.4), 494	0.2 (0.4), 486	0.2 (0.4), 472	0.2 (0.4), 780	0.2 (0.4), 672		
	Year 3	0.2 (0.4), 494	0.2 (0.4), 486	0.2 (0.4), 472	0.2 (0.4), 780	0.2 (0.4), 672		
Have you had bleeding from your gums when brushing your teeth?	Year 3	2.2 (1.0), 484	2.2 (1.0), 473	2.2 (1.0), 459	2.2 (1.0), 777	2.2 (1.0), 639		
Received private PI, n (%)	Year 3	75 (15)	103 (21)	106 (22)	149 (19)	135 (20)		
Missing, n (%)		228 (46)	221 (45)	227 (48)	372 (48)	304 (45)		

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

# **Section 4: tertiary outcomes (at baseline and 3 years)**

Section 4 describes the tertiary measures at baseline (Table 42) and 3-year follow-up (Table 43).

**TABLE 42** Tertiary outcomes at baseline

	Randomised group							
	PI			ОНА				
Tertiary outcomes at baseline	No (N = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)			
How clean do your teeth fee	, mean (SD), <i>n</i>							
After you brush	5.8 (0.9), 605	5.8 (0.9), 608	5.9 (0.9), 605	5.8 (0.9), 975	5.8 (0.9), 843			
After a scale and polish	6.6 (0.8), 574	6.5 (0.9), 589	6.6 (0.8), 574	6.6 (0.9), 927	6.6 (0.8), 810			
How clean do your teeth loo	k, mean (SD), <i>n</i>							
After you brush	5.3 (1.2), 608	5.3 (1.3), 603	5.3 (1.2), 601	5.3 (1.2), 970	5.2 (1.2), 842			
After a scale and polish	6.2 (1.1), 575	6.1 (1.2), 585	6.1 (1.1), 567	6.2 (1.1), 928	6.1 (1.1), 799			
How pleasant do your teeth	feel, mean (SD), <i>n</i>							
After you brush	5.8 (1.1), 607	5.7 (1.1), 604	5.8 (1.1), 600	5.8 (1.1), 971	5.7 (1.1), 840			
After a scale and polish	6.3 (1.0), 574	6.3 (1.0), 581	6.3 (1.0), 565	6.3 (1.0), 924	6.3 (1.1), 796			
How pleasant do your teeth	look, mean (SD), <i>n</i>							
After you brush	5.0 (1.4), 608	5.0 (1.4), 603	5.1 (1.4), 598	5.1 (1.4), 968	5.0 (1.4), 841			
After a scale and polish	5.8 (1.4), 579	5.7 (1.3), 580	5.7 (1.4), 566	5.7 (1.4), 923	5.7 (1.4), 802			
Do you experience sensitivity	in your teeth? n (%	(o)						
Yes	304 (49)	293 (47)	282 (45)	445 (44)	434 (50)			
No	304 (49)	309 (49)	323 (52)	527 (52)	409 (47)			
Missing	15 (2)	23 (4)	21 (3)	36 (4)	23 (3)			
How sensitive are your teeth	? n (%)							
Rarely sensitive	59 (9)	44 (7)	41 (7)	82 (8)	62 (7)			
Sometimes sensitive, rarely interfering with what I eat or drink	166 (27)	152 (24)	135 (22)	221 (22)	232 (27)			
Sometimes sensitive, occasionally interfering with what I eat or drink	79 (13)	90 (14)	98 (16)	135 (13)	132 (15)			
Always sensitive	13 (2)	17 (3)	10 (2)	21 (2)	19 (2)			
Missing	306 (49)	322 (52)	342 (55)	549 (54)	421 (49)			
How sensitive are your teeth?	2.1 (0.8), 317	2.3 (0.8), 303	2.3 (0.7), 284	2.2 (0.8), 459	2.2 (0.7), 445			
On contact with hot and cold, how often do you get pain?	3.1 (1.4), 321	3.1 (1.4), 321	3.0 (1.4), 295	3.1 (1.4), 480	3.1 (1.3), 457			
On contact with hot and cold, how severe is the pain?	2.9 (1.3), 319	2.8 (1.3), 321	2.9 (1.2), 292	2.8 (1.3), 477	2.9 (1.2), 455			

**TABLE 43** Tertiary outcomes at 3 years

	Randomised group							
	PI			ОНА				
Tertiary outcomes at Byears	No ( <i>N</i> = 623)	12-monthly (N = 625)	6-monthly (N = 626)	Personalised (N = 1008)	Routine ( <i>N</i> = 866)			
How clean do your teeth feel	, mean (SD), <i>n</i>							
After you brush	6.0 (0.8), 408	6.0 (0.8), 408	6.1 (0.8), 395	6.0 (0.8), 656	6.0 (0.8), 555			
After a scale and polish	6.6 (0.6), 397	6.6 (0.8), 407	6.7 (0.7), 386	6.7 (0.7), 643	6.6 (0.7), 547			
How clean do your teeth look	k, mean (SD), <i>n</i>							
After you brush	5.5 (1.2), 406	5.4 (1.2), 405	5.6 (1.2), 394	5.5 (1.2), 652	5.5 (1.2), 553			
After a scale and polish	6.3 (1.0), 391	6.1 (1.1), 404	6.2 (1.1), 383	6.2 (1.1), 639	6.2 (1.1), 539			
How pleasant do your teeth f	eel, mean (SD), <i>n</i>							
After you brush	5.9 (1.0), 408	5.9 (1.0), 408	6.0 (1.0), 392	6.0 (1.0), 652	5.9 (1.0), 556			
After a scale and polish	6.5 (0.9), 393	6.4 (1.0), 404	6.5 (0.9), 385	6.5 (0.9), 640	6.4 (1.0), 542			
How pleasant do your teeth l	ook, mean (SD), <i>n</i>							
After you brush	5.3 (1.3), 406	5.2 (1.3), 407	5.4 (1.3), 391	5.3 (1.3), 650	5.2 (1.4), 554			
After a scale and polish	5.9 (1.3), 396	5.8 (1.4), 404	5.9 (1.3), 382	5.9 (1.3), 638	5.9 (1.4), 544			
Do you experience sensitivity	in your teeth?, n (%	o)						
Yes	161 (26)	165 (26)	157 (25)	241 (24)	242 (28)			
No	235 (38)	224 (36)	224 (36)	385 (38)	298 (34)			
Missing	227 (36)	236 (38)	245 (39)	382 (38)	326 (38)			
How sensitive are your teeth?	n (%)							
Rarely sensitive	47 (8)	46 (7)	38 (6)	69 (7)	62 (7)			
Sometimes sensitive, rarely interfering with what I eat or drink	73 (12)	87 (14)	85 (14)	131 (13)	114 (13)			
Sometimes sensitive, occasionally interfere	50 (8)	51 (8)	48 (8)	73 (7)	76 (9)			
Always sensitive	9 (1)	3 (0)	1 (0)	4 (0)	9 (1)			
Missing	444 (71)	438 (70)	454 (73)	731 (73)	605 (70)			
How sensitive are your teeth?, mean (SD), n	2.1 (0.9), 179	2.1 (0.8), 187	2.1 (0.7), 172	2.0 (0.8), 277	2.1 (0.8), 261			
On contact with hot and cold, how often do you get pain?, mean (SD), n	2.8 (1.5), 199	2.8 (1.2), 196	2.8 (1.3), 183	2.7 (1.2), 299	3.0 (1.4), 279			
On contact with hot and cold, how severe is the pain?, mean (SD), n	2.5 (1.2), 199	2.7 (1.2), 195	2.7 (1.2), 183	2.6 (1.2), 298	2.6 (1.2), 279			

# **Appendix 2** Health economics

# Section 1: additional detailed methods for costing, the discrete choice experiment and mapping discrete choice experiment valuations to the trial outcomes

#### Calculations of unit costs

The cost of consumer products was based on a sample of commonly available products in the marketplace. Given that data on the expense of replacing products (e.g. toothbrushes, interdental brushes) were not collected, we have taken a sample of products to represent the variability and the range available. This is not intended to be an exhaustive list of products.

**TABLE 44** Cost of consumer dental care products

Item	Туре	Unit cost (£)	Reference
Interdental brushes	Superdrug Totalcare interdental brushes (6 pack)	2.49	Superdrug (www.superdrug.com; accessed 1 February 2017)
	TePe interdental brushes 0.4 mm (6 pack)	3.05	Superdrug (www.superdrug.com; accessed 1 February 2017)
	Boots Expert TePe interdental brushes 0.4 mm (6 pack)	3.25	Boots (www.boots.com; accessed 1 February 2017)
	TePe interdental brush blue 0.6 mm (6 pack)	5.70	British Corner Shop (www.britishcornershop.co.uk; accessed 1 February 2017)
	Average	3.62 (0.60 each)	
Electric toothbrush	Oral-B® (Proctor & Gamble Co., Cincinatti, OH, USA) Genius 9000 Black	140.00	Boots (www.boots.com; accessed 1 February 2017)
	Oral-B® Pro 600	49.99	Superdrug (www.superdrug.com; accessed 1 February 2017)
	Philips (Amsterdam, the Netherlands) Sonicare DiamondClean	249.99	Boots (www.boots.com; accessed 1 February 2017)
	Colgate® (Colgate-Palmolive, New York, NY, USA) ProClinical® C350	59.99	Boots (www.boots.com; accessed 1 February 2017)
	Oral-B® Vitality Precision	18.99	Tesco (www.tesco.com; accessed 1 February 2017)
	Average	103.79	
Manual toothbrush	Sensodyne® (GlaxoSmithKline plc, GSK House, Middlesex, UK) Precision Toothbrush	3.00	Tesco (www.tesco.com; accessed 1 February 2017)
	Aquafresh® (GSK) Complete Care Medium	2.50	Tesco (www.tesco.com; accessed 1 February 2017)
	Colgate® 360 Max White	4.00	Asda (www.asda.com; accessed 1 February 2017)
	Colgate® Slim Soft	1.22	Sainsbury's (www.sainsburys.co.uk; accessed 1 February 2017)
	Average	2.68	

© Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SC16 7NS, UK.

TABLE 44 Cost of consumer dental care products (continued)

Item	Туре	Unit cost (£)	Reference
electric heads (4 pack) toothbrush Philips Sonicare Pro R	Oral-B® Precision Clean brush heads (4 pack)	18.75 (4.69 each)	Superdrug (www.superdrug.com; accessed 1 February 2017)
	Philips Sonicare Pro Results Standard Sonic toothbrush heads (4 pack)	25.00 (6.25 each)	Philips (www.philips.co.uk; accessed 1 February 2017)
	Colgate® ProClinical Sensitive refill brush heads (4 pack)	18.99 (4.75 each)	Boots (www.boots.com; accessed 1 February 2017)
	Oral-B® Sensitive Clean brush heads (4 pack)	16.00 (4.00 each)	Sainsbury's (www.sainsburys.co.uk; accessed 1 February 2017)
	Average	19.69 (4.92 each)	

# Discrete choice experiment attributes and levels by segment of design

TABLE 45 Experimental design, by segmented group

	Dental Health	Dental Health								
	Good		Moderate		Poor					
Attribute	Levels considered for choice sets	Opt-out	Levels considered for choice sets	Opt-out	Levels considered for choice sets	Opt-out				
Detailed and personalised OHA	All	None	All	None	All	None				
Frequency of PI	All	None	All	None	All	None				
Bleeding gums	Never	Occasional	Hardly ever	Fairly	Occasional	Very				
	Hardly ever		Occasional	often	Fairly often	often				
	Occasional		Fairly often		Very often					
Look and feel	Very clean	Moderately	Clean	Unclean	Moderately clean	Very				
	Clean	clean	Moderately clean		Unclean	unclean				
	Moderately clean		Unclean		Very unclean					
Cost (£)	All	0	All	0	All	0				

#### Mapping discrete choice experiment valuations to trial outcomes

**TABLE 46** Mapping DCE WTP values to trial outcomes

DCE outcome	Trial outcome/variable	Mapping method/notes/assumptions
WTP value for routine or personalised OHA provided by dentist/hygienist	Type of OHA by randomisation  Provider of initial intervention	<ul> <li>Mean WTP value applied to individuals based on randomised group, according to provider of treatment to each individual trial participant – full compliance with type of OHA</li> <li>Total value assigned to year 1 as OHA given at baseline</li> </ul>
WTP value for different annual frequency of PI, provided by hygienist/dentist	Frequency of PI (routine data) Provider (participant questionnaire)	<ul> <li>WTP value assigned to number of Pls actually received</li> <li>Number of Pls based on routine data sources for Scotland (Pl item) and England (clinical data set from FP17 routine data forms<sup>75</sup>), excluding assumed baseline and final study visits</li> <li>Assumed stepwise linear function between 0, 3, 6 and more Pls over trial follow-up</li> <li>Provider of Pl (dentist/hygienist) sourced from baseline clinical examination forms and assumed consistent for duration of follow-up</li> <li>Value assigned to the year in which Pl was given</li> </ul>
WTP value for frequency of bleeding gums	Frequency of bleeding gums at the 3-year follow-up time point	<ul> <li>Direct map from DCE question onto clinical trial outcome, at an individual trial participant level at 3 years' follow-up (i.e. how frequently do your gums bleed when you brush your teeth?)</li> <li>WTP values attached to trial outcomes for complete cases. Values imputed for participants with missing trial outcomes using multiple imputation methods</li> <li>Total value assigned to year 3</li> </ul>
WTP value for aesthetic outcome (teeth look and feel clean and healthy)	How clean do your teeth look and feel 'right now'?	<ul> <li>Mapping for the base case based on an aggregate outcome from four different measures in the trial asking respondents how clean and healthy their teeth look and feel 'right now'</li> <li>Average score on a 7-point scale in the trial (rounded to nearest whole number) reduced to a 5-point scale collapsing extreme ends of the values and assigning WTP values from the DCE directly to each trial participant aggregate score</li> <li>Sensitivity analysis to explore a scenario in which all WTP values are mapped to a similar question asking respondents how clean teeth look and feel 'after brushing', making the alternative assumption that respondents to the DCE may relate how clean their teeth look and feel to when they brush</li> <li>WTP value attached to trial outcomes for complete cases. Values imputed for participants with missing trial outcomes using multiple imputation methods</li> <li>Value assigned to each year</li> </ul>
The annual cost to you	N/A	<ul> <li>Cost attribute used to calculate annual WTP for the outcomes measured above</li> </ul>
N/A, not applicable.		

#### Section 2: research conducted for questionnaire development

#### Literature review

The initial set of draft attributes and levels presented to our advisory group for feedback was informed by a structured review of the literature regarding patient preferences for the processes and outcomes of dental care. The review searched for qualitative studies and surveys from 2000 to 2014 of preferences, experiences, attitudes, beliefs or opinions of adult patients or the general population. Studies were included if they related to routine or preventative dental care. The literature review focused on studies which could inform the selection of attributes for the DCEs. In brief, 15 articles were included and assessed. There was clear qualitative evidence on processes and organisational aspects of care that patients value. Continuity of care, and being treated 'as a person, not as a patient' were viewed positively.<sup>76–78</sup> Dental anxiety and fear were significant barriers to attending the dentist.<sup>79,80</sup> The review reaffirmed the importance of costs,<sup>76,81,82</sup> and provider of care.<sup>76,83</sup>

There was less information on preferences for dental health outcomes, such as bleeding gums (the primary clinical end point for the trial). Therefore, it was crucial to include this as an attribute and to ensure that it was framed correctly within the DCE. There was no information from the literature about the trade-offs between aesthetic and clinical outcomes, particularly for people having PI. The lack of relevant literature motivated the decision to undertake primary FG research.

#### Focus groups

After development of an initial set of attributes and levels important for the trial, and based on the literature available, we ran FG discussions to see if these issues were important to the general population (potential preventative dental care patients) and to determine if we omitted anything relevant and important for our research questions.

Eighteen respondents from the general population in Aberdeen were recruited into four FG sessions. Each group consisted of four or five respondents. FG discussions were recorded, transcribed and analysed following a qualitative thematic framework. The FG protocol was approved by the University of Aberdeen College Ethics Review Board. The FGs provided a rich data set with varied demographic characteristics. Saturation was achieved after the second group.

Focus group results are discussed in the context of developing attributes and levels for the DCEs. The goods to be valued within the DCE were all found to be important to participants, namely frequency of visits, PI and OHA. Participants preferred to avoid dental pain, tooth loss and bad breath. Bleeding gums were important, but only after the facilitator encouraged discussion of this issue. Bleeding was associated with brushing. A description of how bleeding from a clinical point of view might translate into brushing is presented in the DCE to ground the attribute in reality. Dental pain was not directly included as an attribute as it would dominate all of the other attributes in the choice sets, limiting the ability to value any potential differences between groups in trial outcomes. As aesthetic improvement was a prominent reason in the FGs for having a PI, we included an attribute asking respondents to value how clean their teeth look and feel. This aesthetic attribute was intended to value wider, non-health, benefits of PI or OHA.

Focus group data were further used to frame the questions and explain the attributes in the DCEs. For example, many respondents were unaware of who a dental hygienist was, and their knowledge was clearly dependent on experience. A comprehensive description of the roles and responsibilities of a dental hygienist was thus included. There was a preference for personalised care; however, the meaning of 'personalised' is specific to each individual. The survey therefore included a detailed description of what we meant by 'personalised' OHA, in line with the intervention delivered in the trial. Furthermore, many FG participants tended to merge their preferences for PI with a routine visit to the dentist as the two often happen at the same visit. In order to obtain preferences for PI, as opposed to a regular check-up, we tailored the explanation of the PI attribute, emphasising the difference between PI and routine check-ups.

Focus group data identified important contextual information. For example, respondents who had frequent bleeding tended to be less bothered by it. Respondents with experience of a dental hygienist had a more positive view of their role. Fear, anxiety, continuity of care and trust were important drivers of preferences. These findings shaped the selection of a number of contextual questions for inclusion at the end of the DCE surveys.

#### Think-aloud interviews

Once the final design was decided, we recruited five members of the general population and five colleagues to work through the survey, providing comments on the content and framing of the questions. We asked respondents to focus on areas of the questionnaire that were difficult to understand or any problems that they experienced as they went along. Particular attention was paid to how the choice tasks were interpreted and if there was any evidence of decision heuristics or attribute non-attendance. In general, respondents reacted well to the questionnaire. A number of minor bugs in the process were identified and corrected and some minor changes were made to clarify the instructions for the choice tasks, in particular explaining what should and should not be considered when making the choices. No changes were made to the DCE design at this stage as all attributes and levels were found to be acceptable to respondents.

#### Pilot survey

We conducted a comprehensive pilot study with an online representative sample of the general population. A major concern with the external validity of DCE responses relates to the issue of hypothetical bias, in which the respondent's stated preferences are not as they would act in reality. A particularly problematic issue is that, in the presence of hypothetical bias, respondents often overstate the value they place on a good or service. The implication is biased estimates of WTP and, hence, an overvaluation of services in a CBA. Hypothetical bias may therefore contribute to incorrect policy recommendations. There are many potential causes of hypothetical bias and much literature on how best to address it.

We compared a number of suggested approaches, including a cheap talk script, a consequentiality script and asking respondents to sign an honesty oath. All approaches were compared with a standard practice approach of reminding respondents about the availability of close substitutes and asking them to consider their budget constraints when answering the choice tasks. The pilot questionnaire was administered to a nationally representative sample of the general population, with oversampling (n = 100) in Scotland. The Qualtrics survey platform was used to deliver the survey online.

To determine the presence or otherwise of hypothetical bias, we estimated marginal WTP and predicted uptake from the DCE. We compared results across groups. We found no clear evidence of differences across groups. While our sample was small, we were reasonably confident that the base-case approach was sufficient for the main study.

The pilot study was also used to assess the direction of effect of the included attributes and levels. The pilot data were analysed using both simple conditional logit model and mixed logit models, finding that all attributes and levels had good face validity, with the expected coefficient signs and we found that WTP estimates were within the extremes of the levels of the cost attribute presented (i.e. no attribute level was valued at a WTP value of > £200 per year). Therefore, we concluded that the pilot version without correction for hypothetical bias was appropriate for the main phase of data collection.

## **Section 3: detailed claims for Scottish routine data**

TABLE 47 Details of dental claims (Scotland)

	OHA, num	ber of claims							
	Routine	Routine				Personalised			
Routine data claim	No PI (n = 192)	12-monthly PI ( <i>n</i> = 193)	6-monthly PI ( <i>n</i> = 195)	Total (n = 580)	No PI (n = 250)	12-monthly PI (n = 255)	6-monthly PI (n = 252)	Total (n = 757)	Overall total, n (n = 1337)
Clinical	498	495	522	1515	608	629	673	1910	3425
Simple scaling and polishing	253	333	519	1105	238	447	644	1329	2434
Small film	160	149	162	471	232	229	208	669	1140
Composite or synthetic resin	95	112	94	301	117	134	122	373	674
Amalgam	103	102	106	311	129	109	122	360	671
Additional fee for each visit	28	39	27	94	35	43	31	109	203
Fee per course of treatment	28	39	27	94	35	43	31	109	203
Incisal acid etch	26	33	30	89	44	42	27	113	202
Glass ionomer, silicate or silico-phosphate	28	27	33	88	23	39	30	92	180
Additional fee for first inlay or crown in the same arch	17	11	18	46	29	33	31	93	139
Stoning and smoothing surface of a tooth	5	9	8	22	21	38	31	90	112
Treatment of sensitive cementum or dentine	6	13	19	38	14	25	15	54	92
Extensive clinical	13	11	11	35	23	14	18	55	90
Refixing or recementing a crown	15	13	12	40	15	18	14	47	87
Bonded full or jacket crown – non-precious metal	11	8	10	29	17	18	20	55	84
Treatment urgently required for acute conditions	14	16	11	41	10	16	12	38	79
Other treatment claims	158	131	133	422	192	253	216	661	1083
Total claims	1458	1541	1742	4741	1782	2130	2245	6157	10,898

If a code is not claimed, zero resource use is assumed.

DOI: 10.3310/hta22380

## Section 4: participant-reported contact with non-dental health services

TABLE 48 Descriptive statistics for participant-reported contact with non-dental health services

		OHA, n/N (%	)						
		Routine				Personalised			
NHS resource use item		No Pl (n = 289)	12-monthly PI (n = 287)	6-monthly PI (n = 290)	Total (n = 866)	No Pl (n = 334)	12-monthly PI (n = 338)	6-monthly PI (n = 335)	Total (n = 1007)
NHS 24	Year 1	9/204 (4)	7/203 (3)	4/200 (2)	20/607 (3)	4/242 (2)	6/233 (3)	6/240 (3)	16/715 (2)
	Year 2	5/196 (3)	5/186 (3)	6/179 (3)	16/561 (3)	6/207 (3)	3/214 (1)	2/209 (1)	11/630 (2)
	Year 3	1/186 (1)	2/177 (1)	3/170 (2)	6/533 (1)	4/207 (2)	2/215 (1)	1/209 (0)	7/631 (1)
GP	Year 1	26/204 (13)	24/205 (12)	20/200 (10)	70/609 (11)	25/244 (10)	28/237 (12)	17/240 (7)	70/721 (10)
	Year 2	21/196 (11)	20/188 (11)	18/179 (10)	59/563 (10)	24/208 (12)	19/213 (9)	19/209 (9)	62/630 (10)
	Year 3	23/187 (12)	9/179 (5)	7/168 (4)	39/534 (7)	20/207 (10)	17/217 (8)	13/209 (6)	50/633 (8)
A&E	Year 1	10/202 (5)	9/204 (4)	4/198 (2)	23/604 (4)	3/242 (1)	10/236 (4)	7/239 (3)	20/717 (3)
	Year 2	9/196 (5)	6/186 (3)	9/178 (5)	24/560 (4)	7/207 (3)	8/214 (4)	4/210 (2)	19/631 (3)
	Year 3	1/188 (1)	1/176 (1)	3/169 (2)	5/533 (1)	3/207 (1)	4/215 (2)	3/208 (1)	10/630 (2)
Outpatients	Year 1	13/204 (6)	12/203 (6)	18/202 (9)	43/609 (7)	14/243 (6)	12/237 (5)	10/239 (4)	36/719 (5)
	Year 2	13/196 (7)	12/188 (6)	12/178 (7)	37/562 (7)	16/209 (8)	8/214 (4)	10/210 (5)	34/633 (5)
	Year 3	13/188 (7)	7/179 (4)	8/171 (5)	28/538 (5)	9/207 (4)	9/216 (4)	8/209 (4)	26/632 (4)
Inpatients	Year 1	10/203 (5)	5/202 (2)	5/199 (3)	20/604 (3)	3/243 (1)	7/236 (3)	3/239 (1)	13/718 (2)
	Year 2	7/194 (4)	5/185 (3)	6/176 (3)	18/555 (3)	5/208 (2)	4/212 (2)	1/208 (0)	10/628 (2)
	Year 3	3/188 (2)	1/177 (1)	1/167 (1)	5/532 (1)	1/207 (0)	1/212 (0)	2/209 (1)	4/628 (1)

A&E, accident and emergency; GP, general practitioner.

# **Section 5: unit opportunity cost of time and travel to dental appointments**

TABLE 49 Calculation of unit opportunity cost of time and travel to dental appointments

	OHA, n, mean cost (SD)									
	Routine				Personalised	b				
Time and travel costs	No Pl ( <i>N</i> = 289)	12-monthly PI ( <i>N</i> = 287)	6-monthly PI ( <i>N</i> = 290)	Total ( <i>N</i> = 866)	No PI ( <i>N</i> = 334)	12-monthly PI ( <i>N</i> = 338)	6-monthly PI ( <i>N</i> = 335)	Total ( <i>N</i> = 1007)		
Participant time and travel costs										
Participant mode of transport, n (%)										
Data completeness	284 (98)	281 (98)	283 (98)	848 (98)	326 (98)	329 (97)	324 (97)	979 (97)		
Walked	57 (20)	64 (22)	66 (23)	187 (22)	77 (23)	76 (22)	72 (21)	225 (22)		
Cycled	7 (2)	5 (2)	5 (2)	17 (2)	8 (2)	8 (2)	4 (1)	20 (2)		
Private car	186 (64)	173 (60)	177 (61)	536 (62)	209 (63)	213 (63)	212 (63)	634 (63)		
Bus	27 (9)	35 (12)	32 (11)	94 (11)	25 (7)	24 (7)	31 (9)	80 (8)		
Taxi	1 (0)	3 (1)	0 (0)	4 (0)	5 (1)	2 (1)	2 (1)	9 (1)		
Other, non-specified	3 (1)	0 (0)	1 (0)	4 (0)	1 (0)	2 (1)	1 (0)	4 (0)		
Metro	1 (0)	0 (0)	1 (0)	2 (0)	0 (0)	2 (1)	1 (0)	3 (0)		
Boat	0 (0)	0 (0)	0 (0)	0 (0)	1 (0)	2 (1)	0 (0)	3 (0)		
Train	2 (1)	1 (0)	1 (0)	4 (0)	0 (0)	0 (0)	1 (0)	1 (0)		
Missing	5 (2)	6 (2)	7 (2)	18 (2)	8 (2)	9 (3)	11 (3)	28 (3)		
Transportation costs, $n$ , mean (£) (SD, £)	227, 3.46 (4.70)	238, 3.62 (6.32)	238, 3.47 (5.54)	267, 3.46 (6.35)	274, 3.60 (5.11)	703, 3.52 (5.56)	274, 3.63 (8.40)	815, 3.56 (6.75)		
Minutes travelling/waiting, n, mean (SD)	281, 31 (25.56)	276, 31.2 (27.81)	279, 31.36 (25.29)	318, 28.72 (23.66)	322, 29.88 (27.4)	836, 31.19 (26.21)	326, 29.91 (28.11)	966, 29.51 (26.46)		
Participant alternative activity, n (%)										
Data completeness	281 (97)	276 (96)	276 (95)	833 (96)	324 (97)	328 (97)	323 (96)	975 (97)		
Housework	59 (20)	57 (20)	55 (19)	171 (20)	58 (17)	60 (18)	75 (22)	193 (19)		

DOI: 10.3310/hta22380

	OHA, <i>n</i> , mear	n cost (SD)						
	Routine				Personalised	d		
Time and travel costs	No PI ( <i>N</i> = 289)	12-monthly PI ( <i>N</i> = 287)	6-monthly PI ( <i>N</i> = 290)	Total (N = 866)	No Pl ( <i>N</i> = 334)	12-monthly PI ( <i>N</i> = 338)	6-monthly PI ( <i>N</i> = 335)	Total (N = 1007)
Child care	24 (8)	17 (6)	17 (6)	58 (7)	19 (6)	31 (9)	20 (6)	70 (7)
Caring for a friend/relative	6 (2)	6 (2)	3 (1)	15 (2)	7 (2)	6 (2)	5 (1)	18 (2)
Unemployed	14 (5)	11 (4)	15 (5)	40 (5)	9 (3)	18 (5)	14 (4)	41 (4)
Paid work	114 (39)	118 (41)	114 (39)	346 (40)	138 (41)	140 (41)	138 (41)	416 (41)
Voluntary work	3 (1)	5 (2)	5 (2)	13 (2)	6 (2)	2 (1)	4 (1)	12 (1)
Leisure activities	35 (12)	47 (16)	46 (16)	128 (15)	66 (20)	47 (14)	48 (14)	161 (16)
Other non-specified	5 (2)	8 (3)	8 (3)	21 (2)	8 (2)	13 (4)	10 (3)	31 (3)
Studying	12 (4)	3 (1)	5 (2)	20 (2)	5 (1)	6 (2)	4 (1)	15 (1)
Retired	5 (2)	3 (1)	5 (2)	13 (2)	7 (2)	3 (1)	4 (1)	14 (1)
Self-employed	2 (1)	1 (0)	1 (0)	4 (0)	1 (0)	2 (1)	1 (0)	4 (0)
Long-term sick/disabled	2 (1)	0 (0)	2 (1)	4 (0)	0 (0)	0 (0)	0 (0)	0 (0)
Missing	8 (3)	11 (4)	14 (5)	33 (4)	10 (3)	10 (3)	12 (4)	32 (3)
Opportunity cost, alternative activities, $n$ , mean $(f)$ (SD, $f$ )	280, 7.16 (5.10)	274, 7.20 (5.80)	274, 7.23 (5.36)	320, 6.42 (4.54)	323, 6.99 (5.77)	828, 7.20 (5.42)	325, 6.90 (6.12)	968, 6.77 (5.52)
Participant unit cost for time and travel, $n$ , mean $(f)$ (SD, $f$ )	222, 10.53 (8.42)	234, 11.07 (10.70)	230, 10.83 (9.71)	261, 9.85 (9.59)	272, 10.74 (9.87)	686, 10.81 (9.66)	270, 10.61 (13.26)	803, 10.41 (11.04)
Companion time and travel costs								
Participant accompanied, n (%)								
Yes	52 (18)	36 (13)	38 (13)	126 (15)	55 (16)	50 (15)	41 (12)	146 (14)
No	220 (76)	231 (80)	233 (80)	684 (79)	265 (79)	268 (79)	274 (82)	807 (80)
Missing	17 (6)	20 (7)	19 (7)	56 (6)	14 (4)	20 (6)	20 (6)	54 (5)
Relationship of companion <sup>a</sup>								
Partner	21 (36)	16 (41)	15 (35)	52 (37)	29 (51)	25 (45)	14 (33)	68 (44)
Other relative	16 (28)	14 (36)	19 (44)	49 (35)	15 (26)	16 (29)	13 (30)	44 (28)
								continue

TABLE 49 Calculation of unit opportunity cost of time and travel to dental appointments (continued)

	OHA, n, mean o	ost (SD)						
	Routine				Personalise	d		
Time and travel costs	No PI ( <i>N</i> = 289)	12-monthly PI ( <i>N</i> = 287)	6-monthly PI ( <i>N</i> = 290)	Total ( <i>N</i> = 866)	No PI (N = 334)	12-monthly PI ( <i>N</i> = 338)	6-monthly PI ( <i>N</i> = 335)	Total (N = 1007)
Paid companion	0 (0)	0 (0)	1 (2)	1 (1)	0 (0)	0 (0)	0 (0)	0 (0)
Friend	2 (3)	1 (3)	0 (0)	3 (2)	0 (0)	1 (2)	0 (0)	1 (1)
Other	19 (33)	8 (21)	8 (19)	35 (25)	13 (23)	13 (24)	16 (37)	42 (27)
Companion travel costs								
Bus fare, $n$ , mean (£) (SD, £)	34, 0.16 (0.75)	28, 0.08 (0.30)	30, 0.34 (0.84)	92, 0.19 (0.68)	29, 0.21 (0.53)	27, 0.26 (0.77)	24, 0.12 (0.42)	80, 0.20 (0.59)
Total costs, $n$ , mean (£) (SD, £) <sup>b</sup>	263, 0.38 (2.55)	263, 0.09 (0.59)	265, 0.22 (1.60)	791, 0.23 (1.77)	309, 0.15 (0.85)	311, 0.20 (1.00)	308, 0.23 (1.34)	928, 0.19 (1.08)
Companion alternative activity, n (%)								
Housework	5 (9)	3 (8)	6 (15)	14 (11)	12 (23)	11 (21)	3 (8)	26 (18)
Child care	3 (5)	1 (3)	3 (8)	7 (5)	1 (2)	4 (8)	4 (11)	9 (6)
Caring for a relative/friend	3 (5)	1 (3)	3 (8)	7 (5)	1 (2)	3 (6)	1 (3)	5 (4)
Unemployed	0 (0)	1 (3)	3 (8)	4 (3)	1 (2)	0 (0)	0 (0)	1 (1)
Paid work	12 (21)	7 (19)	6 (15)	25 (19)	11 (21)	11 (21)	9 (24)	31 (22)
Voluntary work	0 (0)	1 (3)	1 (3)	2 (2)	2 (4)	0 (0)	0 (0)	2 (1)
Leisure activities	18 (32)	15 (42)	10 (25)	43 (33)	14 (27)	13 (25)	14 (37)	41 (29)
Other non-specified	4 (7)	3 (8)	7 (18)	14 (11)	7 (13)	5 (10)	1 (3)	13 (9)
Studying	9 (16)	3 (8)	1 (3)	13 (10)	1 (2)	3 (6)	4 (11)	8 (6)
Retired	2 (4)	1 (3)	0 (0)	3 (2)	1 (2)	2 (4)	2 (5)	5 (4)
Disabled	0 (0)	0 (0)	0 (0)	0 (0)	1 (2)	0 (0)	0 (0)	1 (1)

DOI: 10.3310/hta22380

	OHA, n, mean cost (SD)								
	Routine				Personalised	Personalised			
Time and travel costs	No PI	12-monthly	6-monthly	Total	No PI	12-monthly	6-monthly	Total	
	( <i>N</i> = 289)	PI ( <i>N</i> = 287)	PI ( <i>N</i> = 290)	(N = 866)	( <i>N</i> = 334)	PI ( <i>N</i> = 338)	PI ( <i>N</i> = 335)	(N = 1007)	
Companion unit cost (£) for time and travel	260, 1.35	260, 0.85	260, 1.03	780, 1.08	301, 0.89	306, 0.98	304; 0.96	911, 0.94	
	(4.57)	(2.94)	(4.07)	(3.92)	(3.28)	(3.16)	(3.72)	(3.40)	
Average number of visits to dentist	271, 3.57	273, 3.75	283, 3.9	827, 3.74	327, 3.21	331, 3.61	328, 3.74	986, 3.52	
	(1.99)	(2.17)	(2.37)	(2.19)	(1.87)	(1.96)	(1.87)	(1.91)	
Total cost (£) of time and travel:complete case	186, 44.08	201, 47.53	195, 47.16	582, 46.30	234, 35.16	249, 43.67	243, 44.38	726, 41.17	
	(51.69)	(57.18)	(48.61)	(52.61)	(36.66)	(58.72)	(50.50)	(49.72)	
Total cost (£) of time and travel: imputation of missing data	289, 43.55	287, 47.31	290, 46.44	866, 45.77	334, 34.57	338, 42.75	335, 44.36	1007, 0.57	
	(51.33)	(57.12)	(48.35)	(52.36)	(36.17)	(56.46)	(50.95)	(48.81)	

a Respondents were able to tick more than one box to indicate who accompanied them (e.g. partner and child 'other relative').

b Companion total travel costs based on 2 × one-way bus fare, if reported. Otherwise, companion travel costs assumed equal to individual participant costs; includes £0 for those who were not accompanied and, hence, paid nothing.

### Section 6: detailed results of further health economic analyses

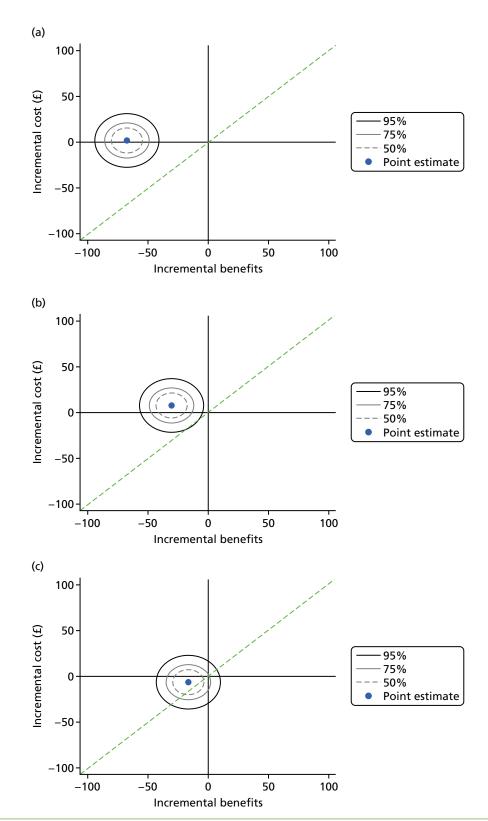
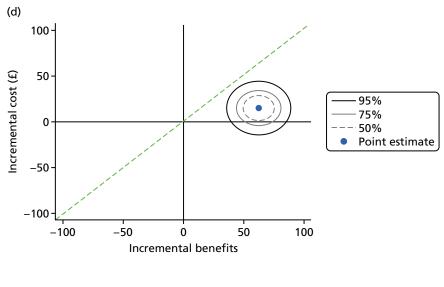


FIGURE 19 Confidence ellipses for base-case health economic results. NHS dental costs: (a) routine OHA, no PI vs. routine OHA, 6-monthly PI; (b) routine OHA, 12-monthly PI vs. routine OHA, 6-monthly PI; (c) personalised OHA, no PI vs. routine OHA, 6-monthly PI; (d) personalised OHA, 6-monthly PI vs. routine OHA, 6-monthly PI; and (e) personalised OHA, 12-monthly PI vs. routine OHA, 6-monthly PI. (continued)



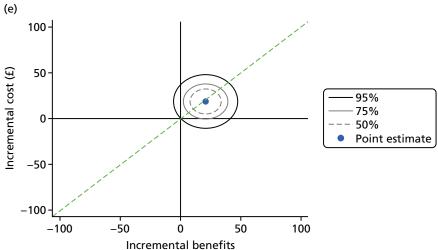


FIGURE 19 Confidence ellipses for base-case health economic results. NHS dental costs: (a) routine OHA, no PI vs. routine OHA, 6-monthly PI; (b) routine OHA, 12-monthly PI vs. routine OHA, 6-monthly PI; (c) personalised OHA, no PI vs. routine OHA, 6-monthly PI; (d) personalised OHA, 6-monthly PI vs. routine OHA, 6-monthly PI; and (e) personalised OHA, 12-monthly PI vs. routine OHA, 6-monthly PI.

TABLE 50 Sensitivity analysis: including baseline and final clinical visits

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	97 (123)	-	231 (139)	_	-
No PI (n = 288)	97 (134)	2 (–22 to 25)	164 (134)	-67 (-92 to -42)	-68 (-103 to -33)
12-monthly PI $(n = 285)$	105 (160)	10 (–13 to 33)	205 (133)	-23 (-48 to 1)	-33 (-67 to 1)
Personalised OHA					
No PI (n = 333)	87 (132)	-3 (-31 to 25)	236 (145)	6 (–23 to 35)	9 (–24 to 42)
12-monthly PI $(n = 338)$	110 (150)	21 (-8 to 49)	277 (134)	45 (17 to 74)	25 (–8 to 57)
6-monthly PI $(n = 335)$	109 (164)	19 (–9 to 47)	317 (150)	88 (59 to 117)	69 (36 to 102)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	101 (118)	_	251 (136)	-	-
No PI (n = 195)	88 (102)	-12 (-36 to 12)	192 (130)	–59 (–89 to –29)	–47 (–85 to –9)
12-monthly PI $(n = 195)$	97 (116)	–2 (–26 to 22)	226 (133)	-24 (-54 to 5)	–22 (–60 to 15)
Personalised OHA					
No PI (n = 252)	87 (114)	-5 (-34 to 23)	260 (141)	7 (–26 to 41)	13 (-23 to 49)
12-monthly PI $(n = 255)$	107 (142)	13 (–16 to 41)	302 (123)	45 (12 to 77)	32 (–3 to 67)
6-monthly PI $(n = 252)$	108 (152)	14 (-14 to 43)	345 (138)	92 (58 to 125)	77 (42 to 113)
England					
Routine OHA					
6-monthly PI $(n = 94)$	89 (133)	-	189 (137)	-	-
No PI (n = 93)	115 (183)	32 (-23 to 86)	105 (122)	-83 (-124 to -43)	-115 (-185 to -45)
12-monthly PI (n = 92)	122 (227)	33 (–20 to 87)	161 (123)	-24 (-65 to 17)	-58 (-126 to 11)
Personalised OHA					
No PI (n = 81)	85 (175)	-2 (-66 to 63)	161 (133)	-15 (-69 to 38)	-14 (-84 to 56)
12-monthly PI $(n = 83)$	119 (173)	35 (–30 to 101)	199 (138)	30 (–24 to 84)	-5 (-76 to 65)
6-monthly PI (n = 83)	111 (197)	27 (–38 to 92)	233 (157)	60 (5 to 116)	34 (-38 to 105)

TABLE 51 Sensitivity analysis: discounting 0% for costs and benefits

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	77 (109)	-	150 (110)	-	-
No PI (n = 288)	78 (126)	2 (–20 to 23)	81 (102)	-69 (-88 to -49)	-70 (-99 to -42)
12-monthly PI $(n = 285)$	85 (147)	9 (–12 to 30)	117 (99)	−31 (−50 to −12)	−40 (−68 to −12)
Personalised OHA					
No PI (n = 333)	65 (120)	-6 (-29 to 17)	130 (110)	-18 (-41 to 4)	-13 (-40 to 15)
12-monthly PI $(n = 338)$	90 (139)	19 (-4 to 43)	170 (100)	20 (0 to 41)	1 (-26 to 28)
6-monthly PI $(n = 335)$	86 (139)	15 (-8 to 38)	212 (120)	64 (42 to 86)	49 (21 to 76)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	75 (96)	-	163 (112)	-	-
No PI (n = 195)	66 (90)	-9 (-29 to 12)	90 (106)	–73 (–97 to –49)	−64 (−95 to −34)
12-monthly PI $(n = 195)$	70 (86)	-4 (-24 to 17)	122 (103)	−40 (−63 to −17)	−36 (−66 to −6)
Personalised OHA					
No PI (n = 252)	63 (106)	-5 (-29 to 19)	139 (112)	–25 (–53 to 2)	-20 (-50 to 9)
12-monthly PI $(n = 255)$	83 (127)	13 (–11 to 37)	181 (97)	14 (–12 to 39)	1 (-27 to 29)
6-monthly PI $(n = 252)$	82 (124)	13 (–11 to 36)	229 (117)	65 (38 to 92)	52 (23 to 81)
England					
Routine OHA					
6-monthly PI $(n = 94)$	82 (132)	-	123 (102)	-	-
No PI $(n = 93)$	102 (177)	25 (–27 to 76)	62 (89)	-60 (-92 to -29)	-85 (-146 to -24)
12-monthly PI $(n = 92)$	116 (224)	35 (–16 to 86)	105 (89)	-14 (-47 to 18)	-50 (-111 to 12)
Personalised OHA					
No PI $(n = 81)$	68 (156)	-12 (-69 to 45)	103 (100)	-11 (-52 to 30)	1 (-62 to 64)
12-monthly PI $(n = 83)$	113 (171)	35 (–24 to 93)	136 (102)	26 (–14 to 67)	-9 (-71 to 54)
6-monthly PI (n = 83)	98 (175)	20 (–38 to 78)	162 (116)	50 (9 to 90)	30 (–32 to 91)

TABLE 52 Sensitivity analysis: discounting 6% for costs and benefits

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI (n = 289)	72 (103)	-	144 (102)	-	-
No PI (n = 288)	72 (117)	1 (-19 to 21)	79 (95)	-65 (-84 to -47)	-67 (-93 to -40)
12-monthly PI (n = 285)	79 (139)	8 (-11 to 28)	113 (91)	−29 (−48 to −11)	−38 (−65 to −11)
Personalised OHA					
No PI (n = 333)	60 (112)	-6 (-27 to 16)	127 (102)	-16 (-36 to 5)	-10 (-36 to 15)
12-monthly PI (n = 338)	84 (129)	18 (–4 to 39)	166 (93)	21 (1 to 41)	4 (–22 to 29)
6-monthly PI (n = 335)	81 (131)	15 (–7 to 36)	205 (112)	62 (41 to 83)	47 (22 to 73)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	71 (91)	-	157 (103)	-	_
No PI (n = 195)	62 (84)	-9 (-28 to 10)	87 (99)	-70 (-93 to -47)	-62 (-90 to -33)
12-monthly PI $(n = 195)$	66 (80)	-4 (-23 to 15)	119 (95)	−38 (−62 to −15)	−34 (−64 to −5)
Personalised OHA					
No PI $(n = 252)$	60 (100)	-5 (-27 to 17)	136 (103)	-22 (-48 to 3)	-18 (-45 to 10)
12-monthly PI $(n = 255)$	78 (118)	12 (–11 to 34)	176 (91)	15 (–10 to 40)	3 (-24 to 30)
6-monthly PI $(n = 252)$	77 (117)	12 (–11 to 34)	221 (108)	63 (37 to 88)	51 (24 to 78)
England					
Routine OHA					
6-monthly PI $(n = 94)$	76 (125)	-	118 (94)	-	-
No PI ( <i>n</i> = 93)	95 (165)	23 (–26 to 72)	61 (84)	−56 (−85 to −27)	–79 (–136 to –22)
12-monthly PI $(n = 92)$	109 (213)	33 (–15 to 82)	101 (82)	-13 (-42 to 15)	-47 (-103 to 10)
Personalised OHA					
No PI (n = 81)	63 (144)	-12 (-65 to 42)	101 (94)	-8 (-46 to 30)	3 (-54 to 61)
12-monthly PI $(n = 83)$	104 (157)	31 (-23 to 86)	133 (94)	27 (–10 to 65)	-4 (-62 to 54)
6-monthly PI (n = 83)	93 (166)	20 (–34 to 73)	155 (109)	48 (9 to 88)	28 (-30 to 87)

TABLE 53 Sensitivity analysis: all PI and OHA delivered by a dentist

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	74 (106)	-	153 (107)	-	-
No PI (n = 288)	74 (119)	1 (-20 to 21)	83 (99)	−70 (−89 to −51)	-70 (-98 to -43)
12-monthly PI $(n = 285)$	81 (141)	9 (–12 to 29)	122 (96)	−29 (−48 to −10)	−37 (−65 to −10)
Personalised OHA					
No PI (n = 333)	62 (115)	-6 (-27 to 16)	137 (104)	-13 (-35 to 8)	-8 (-34 to 19)
12-monthly PI $(n = 338)$	87 (134)	18 (-3 to 40)	178 (97)	27 (6 to 47)	8 (–18 to 35)
6-monthly PI $(n = 335)$	83 (134)	15 (–7 to 37)	220 (113)	70 (48 to 91)	55 (28 to 82)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	72 (93)	-	161 (107)	-	-
No PI (n = 195)	64 (86)	-8 (-28 to 11)	89 (104)	-72 (-95 to -49)	-64 (-93 to -34)
12-monthly PI $(n = 195)$	67 (82)	-4 (-23 to 16)	122 (99)	−39 (−62 to −17)	−35 (−64 to −6)
Personalised OHA					
No PI (n = 252)	61 (103)	-5 (-27 to 18)	141 (106)	-20 (-46 to 6)	-15 (-44 to 13)
12-monthly PI $(n = 255)$	80 (121)	12 (–11 to 35)	184 (95)	19 (-6 to 44)	7 (–20 to 35)
6-monthly PI $(n = 252)$	79 (120)	12 (–10 to 35)	231 (110)	69 (43 to 95)	57 (29 to 85)
England					
Routine OHA					
6-monthly PI $(n = 94)$	78 (128)	-	135 (106)	-	_
No PI (n = 93)	95 (167)	21 (-28 to 71)	69 (87)	-65 (-97 to -34)	-86 (-145 to -28)
12-monthly PI $(n = 92)$	112 (216)	33 (–16 to 83)	122 (90)	-9 (-42 to 24)	–43 (–103 to 17)
Personalised OHA					
No PI (n = 81)	65 (148)	-13 (-68 to 42)	123 (97)	-3 (-43 to 36)	9 (-50 to 69)
12-monthly PI $(n = 83)$	108 (165)	33 (–23 to 88)	160 (101)	38 (–3 to 79)	6 (–56 to 67)
6-monthly PI ( <i>n</i> = 83)	95 (170)	19 (–36 to 74)	185 (116)	62 (20 to 103)	43 (–19 to 104)

TABLE 54 Sensitivity analysis: all PI and OHA delivered by a hygienist

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	74 (106)	-	126 (100)	-	-
No PI (n = 288)	74 (119)	1 (-20 to 21)	65 (89)	-60 (-78 to -43)	-61 (-88 to -34)
12-monthly PI $(n = 285)$	81 (141)	9 (–12 to 29)	95 (89)	−29 (−46 to −11)	−37 (−64 to −10)
Personalised OHA					
No PI (n = 333)	62 (115)	-6 (-27 to 15)	100 (94)	-24 (-43 to -4)	-18 (-44 to 8)
12-monthly PI (n = 338)	87 (134)	18 (-3 to 39)	130 (88)	5 (–14 to 24)	-13 (-39 to 13)
6-monthly PI $(n = 335)$	83 (134)	15 (-7 to 36)	167 (105)	43 (23 to 63)	29 (2 to 55)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	72 (93)	-	131 (102)	-	-
No PI (n = 195)	64 (86)	-8 (-28 to 11)	68 (92)	-63 (-84 to -41)	-54 (-83 to -25)
12-monthly PI $(n = 195)$	67 (82)	-4 (-23 to 16)	94 (92)	−37 (−58 to −16)	−33 (−61 to −4)
Personalised OHA					
No PI $(n = 252)$	61 (103)	-5 (-27 to 17)	103 (96)	-28 (-53 to -4)	-24 (-51 to 4)
12-monthly PI $(n = 255)$	80 (121)	12 (–10 to 34)	133 (87)	-1 (-24 to 22)	-13 (-40 to 14)
6-monthly PI $(n = 252)$	79 (120)	12 (-10 to 34)	175 (103)	44 (19 to 68)	31 (4 to 59)
England					
Routine OHA					
6-monthly PI $(n = 94)$	78 (128)	-	115 (97)	-	-
No PI (n = 93)	95 (167)	21 (-28 to 71)	58 (82)	−56 (−86 to −27)	–78 (–135 to –20)
12-monthly PI $(n = 92)$	112 (216)	33 (–16 to 83)	98 (81)	-13 (-45 to 18)	-47 (-106 to 13)
Personalised OHA					
No PI (n = 81)	65 (148)	-14 (-67 to 39)	89 (88)	-19 (-54 to 17)	-5 (-64 to 54)
12-monthly PI $(n = 83)$	108 (165)	32 (–22 to 85)	118 (91)	15 (–22 to 51)	-17 (-78 to 43)
6-monthly PI (n = 83)	95 (170)	18 (–35 to 71)	141 (106)	35 (–2 to 72)	17 (–43 to 77)

TABLE 55 Sensitivity analysis: adding an additional cost for personalised OHA to all respondents in this cluster

	Mean (SD)	Mean difference in costs (95% CI) vs. routine OHA	Mean (SD)	Mean difference in benefits (95% CI) vs. routine OHA	Incremental net benefits (95% CI) vs. routine OHA
Region	costs, £ª	(6-monthly PI), £	benefits, £	(6-monthly PI), £	(6-monthly PI), £
UK OUA					
Routine OHA	74 (106)		1.47 (10.4)		
6-monthly PI $(n = 289)$	74 (106)	_	147 (104)	_	_
No PI (n = 288)	75 (121)	1 (-26 to 28)	79 (98)	-67 (-85 to -49)	-68 (-101 to -36)
12-monthly PI $(n = 285)$	81 (140)	8 (–18 to 35)	115 (95)	−30 (−48 to −12)	−39 (−70 to −7)
Personalised OHA					
No PI ( <i>n</i> = 333)	109 (215)	37 (–13 to 87)	129 (104)	-17 (-37 to 3)	−54 (−104 to −4)
12-monthly PI (n = 338)	135 (190)	66 (16 to 116)	166 (96)	19 (–2 to 39)	-48 (-100 to 5)
6-monthly PI $(n = 335)$	105 (177)	34 (–5 to 72)	207 (117)	61 (40 to 82)	28 (–15 to 70)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	72 (93)	-	159 (105)	-	-
No PI ( <i>n</i> = 195)	64 (86)	-9 (-40 to 23)	88 (102)	−72 (−94 to −50)	-64 (-102 to -26)
12-monthly PI ( <i>n</i> = 195)	67 (82)	-4 (-36 to 27)	120 (98)	−39 (−62 to −17)	-35 (-74 to 3)
Personalised OHA					
No PI $(n = 252)$	122 (231)	52 (-12 to 116)	138 (105)	-23 (-46 to 0)	−75 (−141 to −10)
12-monthly PI $(n = 255)$	143 (196)	75 (11 to 140)	177 (93)	12 (–11 to 35)	-63 (-130 to 4)
6-monthly PI $(n = 252)$	106 (177)	36 (-12 to 84)	223 (113)	62 (39 to 86)	26 (-27 to 80)
England					
Routine OHA					
6-monthly PI $(n = 94)$	79 (130)	-	120 (96)	_	-
No PI ( <i>n</i> = 93)	98 (170)	23 (–28 to 73)	62 (86)	−57 (−87 to −27)	−80 (−140 to −21)
12-monthly PI $(n = 92)$	112 (214)	33 (–17 to 83)	103 (85)	-13 (-43 to 17)	-46 (-104 to 12)
Personalised OHA					
No PI ( <i>n</i> = 81)	69 (152)	-9 (-65 to 47)	103 (97)	-8 (-47 to 31)	1 (-59 to 61)
12-monthly PI (n = 83)	111 (164)	35 (–22 to 91)	132 (98)	26 (–13 to 65)	-9 (-70 to 52)
6-monthly PI (n = 83)	102 (177)	25 (–32 to 82)	158 (113)	49 (11 to 87)	24 (-37 to 85)

<sup>©</sup> Queen's Printer and Controller of HMSO 2018. This work was produced by Ramsay et al. under the terms of a commissioning contract issued by the Secretary of State for Health and Social Care. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

TABLE 56 Sensitivity analysis: decreasing value of UDA (-20%)

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	69 (96)	-	146 (105)	-	-
No PI (n = 288)	68 (104)	0 (-19 to 18)	79 (98)	-66 (-85 to -48)	-66 (-91 to -40)
12-monthly PI $(n = 285)$	74 (119)	7 (–12 to 25)	115 (95)	–29 (–48 to –11)	-36 (-61 to -10)
Personalised OHA					
No PI (n = 333)	59 (107)	-5 (-25 to 16)	128 (105)	-17 (-38 to 5)	-12 (-37 to 13)
12-monthly PI $(n = 338)$	81 (124)	18 (–3 to 38)	166 (97)	21 (0 to 42)	3 (–21 to 28)
6-monthly PI $(n = 335)$	78 (124)	15 (–6 to 35)	207 (115)	63 (41 to 85)	49 (23 to 74)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	72 (93)	-	159 (107)	-	-
No PI ( <i>n</i> = 195)	64 (86)	-8 (-28 to 11)	88 (102)	-71 (-94 to -49)	−63 (−92 to −34)
12-monthly PI $(n = 195)$	67 (82)	-4 (-23 to 16)	120 (99)	−39 (−61 to −17)	−35 (−64 to −6)
Personalised OHA					
No PI (n = 252)	61 (103)	-5 (-28 to 18)	137 (106)	-23 (-49 to 3)	-18 (-47 to 10)
12-monthly PI $(n = 255)$	80 (121)	12 (–11 to 35)	178 (94)	14 (-11 to 39)	2 (–25 to 29)
6-monthly PI $(n = 252)$	79 (120)	12 (–11 to 35)	225 (110)	64 (38 to 90)	52 (24 to 80)
England					
Routine OHA					
6-monthly PI $(n = 94)$	63 (103)	-	118 (97)	-	-
No PI (n = 93)	76 (133)	17 (–22 to 57)	61 (85)	−56 (−87 to −26)	−73 (−123 to −24)
12-monthly PI $(n = 92)$	89 (173)	27 (–13 to 67)	104 (85)	-10 (-42 to 22)	-37 (-89 to 14)
Personalised OHA					
No PI (n = 81)	52 (119)	-9 (-55 to 36)	101 (97)	-8 (-47 to 30)	1 (-50 to 52)
12-monthly PI $(n = 83)$	86 (132)	27 (–19 to 73)	132 (98)	28 (–12 to 67)	0 (–53 to 53)
6-monthly PI (n = 83)	76 (136)	16 (–30 to 62)	154 (113)	48 (8 to 88)	32 (–21 to 84)

TABLE 57 Sensitivity analysis: increasing value of UDA (+20%)

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	79 (117)	-	146 (105)	-	-
No PI (n = 288)	80 (136)	2 (-21 to 25)	79 (98)	-66 (-85 to -48)	−68 (−97 to −39)
12-monthly PI $(n = 285)$	89 (164)	11 (–12 to 34)	115 (95)	–29 (–48 to –11)	−40 (−69 to −10)
Personalised OHA					
No PI (n = 333)	65 (125)	-7 (-31 to 17)	128 (105)	-16 (-38 to 5)	-9 (-38 to 19)
12-monthly PI $(n = 338)$	92 (145)	19 (-5 to 43)	166 (97)	21 (0 to 41)	2 (–27 to 30)
6-monthly PI $(n = 335)$	87 (146)	15 (–9 to 39)	207 (115)	63 (42 to 85)	48 (20 to 77)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	72 (93)	-	159 (107)	-	-
No PI ( <i>n</i> = 195)	64 (86)	-8 (-28 to 11)	88 (102)	-71 (-94 to -49)	-63 (-92 to -34)
12-monthly PI $(n = 195)$	67 (82)	-4 (-23 to 16)	120 (99)	−39 (−61 to −17)	−35 (−64 to −6)
Personalised OHA					
No PI (n = 252)	61 (103)	-5 (-28 to 18)	137 (106)	-23 (-49 to 3)	-18 (-47 to 10)
12-monthly PI $(n = 255)$	80 (121)	12 (–11 to 35)	178 (94)	14 (-11 to 39)	2 (-25 to 29)
6-monthly PI $(n = 252)$	79 (120)	12 (–11 to 35)	225 (110)	64 (38 to 90)	52 (24 to 80)
England					
Routine OHA					
6-monthly PI $(n = 94)$	94 (154)	-	118 (97)	-	_
No PI (n = 93)	114 (200)	26 (-34 to 85)	61 (85)	−57 (−87 to −26)	−82 (−148 to −16)
12-monthly PI $(n = 92)$	134 (259)	40 (–19 to 100)	104 (85)	-10 (-42 to 22)	–51 (–119 to 17)
Personalised OHA					
No PI (n = 81)	78 (178)	-16 (-80 to 49)	101 (97)	-9 (-49 to 31)	7 (-61 to 74)
12-monthly PI ( <i>n</i> = 83)	130 (198)	39 (–27 to 105)	132 (98)	27 (–14 to 69)	-12 (-81 to 58)
6-monthly PI (n = 83)	114 (204)	22 (–43 to 88)	154 (113)	47 (6 to 89)	25 (–44 to 94)

TABLE 58 Alternative method to map from DCE to trial for aesthetic outcome (mapping to 'after brushing')

Region	Mean (SD) costs, £	Mean difference in costs (95% CI) vs. routine OHA (6-monthly PI), £	Mean (SD) benefits, £	Mean difference in benefits (95% CI) vs. routine OHA (6-monthly PI), £	Incremental net benefits (95% CI) vs. routine OHA (6-monthly PI), £
UK					
Routine OHA					
6-monthly PI $(n = 289)$	74 (105)	-	177 (92)	-	-
No PI (n = 288)	74 (118)	1 (-20 to 21)	113 (85)	-64 (-79 to -49)	-64 (-89 to -39)
12-monthly PI $(n = 285)$	81 (141)	9 (–12 to 29)	148 (78)	−28 (−43 to −13)	−37 (−61 to −12)
Personalised OHA					
No PI (n = 333)	62 (116)	-6 (-28 to 17)	166 (84)	-11 (-29 to 7)	-5 (-29 to 18)
12-monthly PI (n = 338)	86 (133)	18 (-4 to 40)	203 (78)	26 (8 to 44)	7 (–16 to 31)
6-monthly PI $(n = 335)$	83 (134)	15 (–8 to 37)	235 (100)	59 (40 to 77)	44 (20 to 68)
Scotland					
Routine OHA					
6-monthly PI $(n = 195)$	72 (93)	-	190 (92)	-	-
No PI (n = 195)	63 (86)	-9 (-28 to 11)	123 (89)	-67 (-86 to -49)	−59 (−85 to −33)
12-monthly PI $(n = 195)$	67 (82)	-4 (-23 to 16)	153 (82)	−38 (−56 to −20)	−34 (−59 to −8)
Personalised OHA					
No PI (n = 252)	61 (103)	-5 (-27 to 17)	173 (86)	-18 (-39 to 3)	-13 (-37 to 11)
12-monthly PI $(n = 255)$	79 (121)	12 (-11 to 34)	212 (75)	18 (-3 to 40)	7 (–18 to 31)
6-monthly PI $(n = 252)$	79 (120)	12 (-10 to 34)	253 (95)	62 (41 to 84)	50 (26 to 75)
England					
Routine OHA					
6-monthly PI $(n = 94)$	78 (127)	-	150 (88)	-	-
No PI (n = 93)	95 (165)	21 (-28 to 70)	93 (72)	−57 (−81 to −33)	−79 (−134 to −23)
12-monthly PI $(n = 92)$	112 (215)	34 (-15 to 83)	137 (68)	-10 (-34 to 13)	-45 (-99 to 10)
Personalised OHA					
No PI (n = 81)	66 (150)	-12 (-68 to 44)	142 (73)	-1 (-36 to 35)	11 (-45 to 68)
12-monthly PI $(n = 83)$	108 (162)	32 (–24 to 89)	176 (81)	36 (0 to 71)	3 (-53 to 60)
6-monthly PI (n = 83)	95 (170)	19 (–37 to 75)	180 (94)	38 (3 to 74)	19 (–37 to 76)

# **Appendix 3** Recruitment

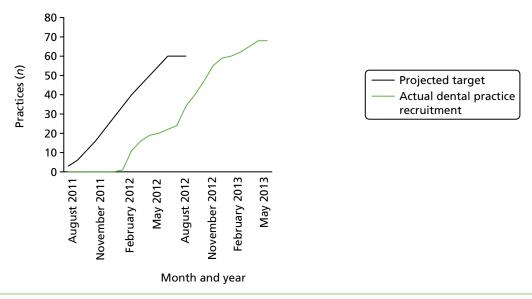


FIGURE 20 Target vs. actual practice accrual.

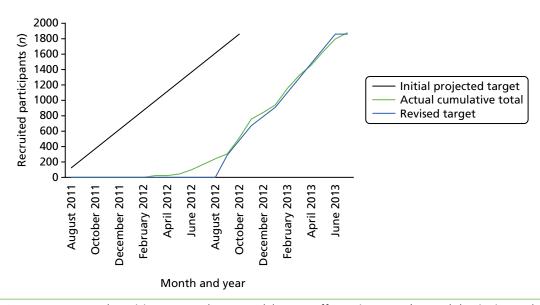


FIGURE 21 Target vs. actual participant accrual. Note: a delay to staff appointments (research hygienists and dental research nurses) resulted in a delay to the start of recruitment of dental practices and participants against the initial projected target accrual.

**TABLE 59** Recruitment by centre

	ОНА		Randomisation date	
Centre	Personalised	Routine	First participant	Last participant
Scotland				
Dental Care Glenrothes	_	31	2 March 2012	18 May 2012
Discovery Dental Care	33	_	17 July 2012	28 September 2012
Mullins Dental Practice	35	_	11 June 2012	15 June 2012
Laurencekirk Dental Practice	_	13	17 May 2012	30 May 2012
Woodside Dental Practice	_	32	19 July 2012	16 August 2012
Baillieston Dental Care	31	_	19 June 2012	24 October 2012
P A Penney Dental Practice	_	14	26 July 2012	20 September 2012
Wishaw Cross Dental Care	31	_	6 August 2012	18 October 2012
Clyde Dental	_	23	18 July 2012	23 October 2012
S. Rankin	31	-	1 October 2012	5 December 2012
Barrhead Dental practice	29	_	13 August 2012	10 September 2012
The Hollies Dental Practice	_	39	29 October 2012	1 November 2012
Chong Kwan and Associates	32	-	5 November 2012	9 November 2012
Duns Dental Practice	_	35	9 October 2012	2 November 2012
Clark and Watson	_	33	26 September 2012	14 November 2012
Dental Care Perth	_	18	23 October 2012	25 February 2013
Dental Plus	31	_	3 October 2012	20 December 2012
Stirling Dental Care	31	-	5 November 2012	25 March 2013
West End Dental Practice	32	_	15 November 2012	13 December 2012
JL Barrack Dental Practice	_	31	31 October 2012	24 January 2013
Selkirk	34	-	22 October 2012	9 November 2012
Bridge of Don Dental Practice	_	38	19 November 2012	10 December 2012
Ardmillan Dental Practice	_	34	11 December 2012	22 January 2013
Drumbrae Dental Surgery	16	_	13 November 2012	11 January 2013
Pickering Dental Care Ltd	_	25	5 December 2012	6 May 2013
Karolak and Iwanowicz	_	22	14 January 2013	15 April 2013
City Health Clinic	_	34	25 January 2013	23 May 2013
Care Dental (Crieff)	31	-	7 January 2013	3 June 2013
Dunbar Dental Practice	43	-	4 February 2013	8 February 2013
Long and Gilmour Dental Care	22	-	11 March 2013	25 June 2013
Riverview Dental Practice	_	30	10 April 2013	25 June 2013
Montgomery Street Dental	32	-	29 March 2013	22 May 2013
Invergowrie Dental	32	_	17 January 2013	15 March 2013
East Neuk Dental	_	30	25 January 2013	12 April 2013
Mastrick Dental	32	_	12 February 2013	3 April 2013
Birch Valley Dental Clinic	34	-	19 February 2013	21 February 2013

TABLE 59 Recruitment by centre (continued)

	ОНА		Randomisation date	
Centre	Personalised	Routine	First participant	Last participant
Whitburn Dental Practice	_	34	27 March 2013	19 June 2013
K H Stirling	29	-	21 March 2013	16 May 2013
Bute Dental Practice	_	34	27 May 2013	31 May 2013
Peacocks Dental Surgery	22	_	20 May 2013	24 May 2013
St Leonards	32	-	1 May 2013	3 May 2013
Lochboisdale, South Uist	41	_	1 July 2013	5 July 2013
King Street, Orkney	46	_	10 June 2013	14 June 2013
D A Gilchrist	_	36	26 June 2013	3 July 2013
England				
Mr A I Robson and Associates	31	_	6 September 2012	18 October 2012
Mr S B Pabary and Associates	31	_	13 November 2012	30 April 2013
Sunderland Road Dental Practice	_	31	12 September 2012	6 November 2012
Framwellgate Dental Practice	24	_	18 September 2012	5 February 2013
Osborne Dental Practice	26	-	12 June 2012	3 December 2012
Hetton Dental	_	24	15 November 2012	24 April 2013
Shotley Bridge	31	-	29 November 2012	11 December 2012
Number One	_	14	18 February 2013	4 July 2013
Stuart Eaborn Dentist	31	-	5 February 2013	23 April 2013
Dean Road Dental Surgery	33	-	10 January 2013	11 March 2013
Smith and Smith	_	33	28 February 2013	14 March 2013
Kenton Lane	_	31	15 April 2013	28 May 2013
Whickham Village	34	-	13 February 2013	28 March 2013
Blaydon Dental	_	32	5 March 2013	27 March 2013
Gosforth Dental	_	31	15 May 2013	5 June 2013
Nepali Dental	_	30	15 May 2013	4 July 2013
Princes Street	_	32	28 May 2013	27 June 2013
Front Street, Consett	7	-	5 June 2013	5 June 2013
Silver Dental Practice	_	23	20 June 2013	4 July 2013

# EME HS&DR HTA PGfAR PHR

Part of the NIHR Journals Library www.journalslibrary.nihr.ac.uk

This report presents independent research funded by the National Institute for Health Research (NIHR). The views expressed are those of the author(s) and not necessarily those of the NHS, the NIHR or the Department of Health