# Inhibition of WEE1 Is Effective in TP53and RAS-Mutant Metastatic Colorectal Cancer: A Randomized Trial (FOCUS4-C) Comparing Adavosertib (AZD1775) With Active Monitoring

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PURPOSE Outcomes in RAS-mutant metastatic colorectal cancer (mCRC) remain poor and patients have limited therapeutic options. Adavosertib is the first small-molecule inhibitor of WEE1 kinase. We hypothesized that aberrations in DNA replication seen in mCRC with both RAS and TP53 mutations would sensitize tumors to WEE1 inhibition.

METHODS Patients with newly diagnosed mCRC were registered into FOCUS4 and tested for TP53 and RAS mutations. Those with both mutations who were stable or responding after 16 weeks of chemotherapy were randomly assigned 2:1 between adavosertib and active monitoring (AM). Adavosertib (250 mg or 300 mg) was taken orally once on days 1-5 and days 8-12 of a 3-week cycle. The primary outcome was progression-free survival (PFS), with a target hazard ratio (HR) of 0.5 and 80% power with a one-sided 0.025 significance level.

RESULTS FOCUS4-C was conducted between April 2017 and Mar 2020 during which time 718 patients were registered; 247 (34%) were RAS/TP53-mutant. Sixty-nine patients were randomly assigned from 25 UK hospitals (adavosertib = 44; AM = 25). Adavosertib was associated with a PFS improvement over AM (median  $3.61 \text{ v} \cdot 1.87 \text{ months}$ ; HR = 0.35; 95% CI, 0.18 to 0.68; P = .0022). Overall survival (OS) was not improved with adavosertib versus AM (median 14.0 v 12.8 months; HR = 0.92; 95% CI, 0.44 to 1.94; P = .93). In prespecified subgroup analysis, adavosertib activity was greater in left-sided tumors (HR = 0.24; 95% CI, 0.11 to 0.51), versus right-sided (HR = 1.02; 95% CI, 0.41 to 2.56; interaction P = .043). Adavosertib was well-tolerated; grade 3 toxicities were diarrhea (9%), nausea (5%), and neutropenia (7%).

**CONCLUSION** In this phase II randomized trial, adavosertib improved PFS compared with AM and demonstrates potential as a well-tolerated therapy for RAS/TP53-mutant mCRC. Further testing is required in this sizable population of unmet need.

J Clin Oncol 39:3705-3715. © 2021 by American Society of Clinical Oncology

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## **ASSOCIATED** CONTENT Appendix

## **Data Supplement Protocol**

Author affiliations and support information (if applicable) appear at the end of this article.

Accepted on August 16, 2021 and nublished at ascopubs.org/journal/ jco on September 18, 2021: DOI https://doi. org/10.1200/JC0.21. 01435

### INTRODUCTION

Targeting the cellular DNA damage response (DDR) has been an effective therapeutic strategy in several tumor sites, including ovarian and pancreatic cancer.<sup>1,2</sup> These agents can be used as monotherapy in cancers with defective DDR, where we might anticipate a synthetic lethality interaction: two pathways together perform an essential function, and the loss of one pathway (eg, because of mutation) is tolerated but the loss of both pathways leads to cell death.<sup>3</sup>

WEE1 is a nuclear tyrosine kinase that has a central role in cell cycle regulation, including being the key regulator of the G2/M checkpoint through actions on CDK1,4 optimizing DNA-histone stoichiometry before mitotic

entry<sup>4</sup> and modulation of CDK1/2 during the intra-S phase to block replication initiation.<sup>5</sup> Inhibition of WEE1 causes unscheduled entry into mitosis, aberrant firing of replication origins leading to dNTP (Dithiobis [5-nitropyridine]) shortage and replication stress,4 and accumulation of DNA damage during S phase, leading to increased reliance on the G1/S checkpoint.4 Adavosertib (AZD1775) is the first small-molecule inhibitor of WEE1 kinase and has been tested in combination with chemotherapy and radiotherapy<sup>6,7</sup> but more recently as monotherapy to generate synthetic lethality in tumors with DDR defects.6

There has been limited investigation into agents targeting the DDR in metastatic colorectal cancer



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#### CONTEXT

#### **Key Objective**

To test if adavosertib, which is a small-molecule inhibitor of the WEE1 kinase, is effective as monotherapy in patients with *RAS/TP53*-mutant metastatic colorectal cancer (mCRC) as maintenance therapy following induction chemotherapy.

## **Knowledge Generated**

In this phase II randomized trial, adavosertib was well-tolerated and improved progression-free survival in *RAS/TP53*-mutant mCRC compared with active monitoring. Treatment effect may be affected by primary tumor location and *KRAS* subtype, with greater benefit seen in left-sided cancers and those with *KRAS* codon 12/13 mutations. *RAS/TP53* subgroup is a distinct moderately poor prognostic population.

#### Relevance

Adavosertib is a promising therapeutic agent in patients with *RAS/P53*-mutant mCRC, a poor prognostic population of unmet need, and was well-tolerated. This study demonstrates the potential of targeting the DNA damage response pathway in mCRC, which should be a research priority. Future studies of adavosertib should stratify patient outcomes according to primary tumor location and *RAS* subtype.

(mCRC), mainly because of the lack of systematic identification of alterations in DDR genes.8 Here, we test adavosertib in RAS- and TP53-mutant (RAS/TP53-mut) mCRC, which we hypothesize would be sensitive to WEE1 inhibition. TP53 is a key regulator of the G1/S checkpoint<sup>9</sup>; loss of function leads to dependence on the intra-S and G2/M checkpoints to detect DNA damage and initiate repair. 10 In preclinical studies, AZD1775 possessed preferential killing effect in TP53-deficient compared with TP53 wild-type tumors. 11 Mutant RAS, as well as recognized actions through downstream mitogenactivated protein kinase B (MAPK-AKT) pathway signaling, also drives cell cycle progression leading to replication stress during S phase. 12 In preclinical studies, mutant RAS drives cells into S phase through regulation of the CDK4 or CDK6 complex and provides sustained mitogenic signals through sustained CDK2 activity. These effects activate the replication stress response including checkpoint activation. 13 Theoretically, RAS/TP53-mut tumors will be highly vulnerable to adavosertib, with G1 checkpoint failure, evidence of replication stress, and reliance on the intra-S phase and G2/M checkpoints.

The FOCUS4 trial program was an adaptive molecularly stratified umbrella platform trial that evaluated the safety and efficacy of novel treatments in targeted biomarker subgroups within a phase II/III trial setting in the interval after 16 weeks of first-line therapy of mCRC. The design has been published separately, <sup>14</sup> and the trial schema, registration, and biomarker methods are provided in the Data Supplement (online only). Here, we report the findings of FOCUS4-C, which tested the safety and efficacy of adavosertib in patients with *RAS/TP53*-mut mCRC compared with active monitoring (AM) and has achieved disease stability following induction chemotherapy.

#### **METHODS**

## Trial Approvals, Patient Eligibility, and Recruitment

The trial and subsequent amendments were approved by the UK National Ethics Committee Oxford—Panel C (reference 13/SC/0111) and by the relevant regulatory body MHRA (CTA No. 20363/0400/001 and EudraCT No. 2012-005111-12).

Patients age more than 18 years with newly diagnosed mCRC were registered into the FOCUS4 trial program, while undergoing induction chemotherapy, from a total of 88 UK hospitals. Following registration, a tumor sample was tested using next generation sequencing platform for stratification into molecular subtypes including *BRAF*, *PIK3CA*, *TP53*, and *RAS* mutations (Fig 1 and Data Supplement). Patients were required to provide written informed consent for both tissue testing and entry into any of the randomized subtrials including FOCUS4-C.

Patients were randomly assigned into the FOCUS4-C trial in a subset of 25 hospitals between July 2017 and March 2020. Patients were eligible if their tumor had both *RAS* and *TP53* mutations and they had disease stability or response as assessed by computed tomography (CT) scan at the end of 16 weeks of induction chemotherapy, at which point the chemotherapy ceased and the patient was randomly assigned. Patients required a baseline CT scan 4 weeks before random assignment, a minimum 3-week washout period between the last dose of chemotherapy or biologic therapy and the first dose of adavosertib, adequate renal (creatinine clearance > 50 mL/min) and liver function, a WHO performance status of 0-1, and no evidence of prolonged QT interval on ECG.

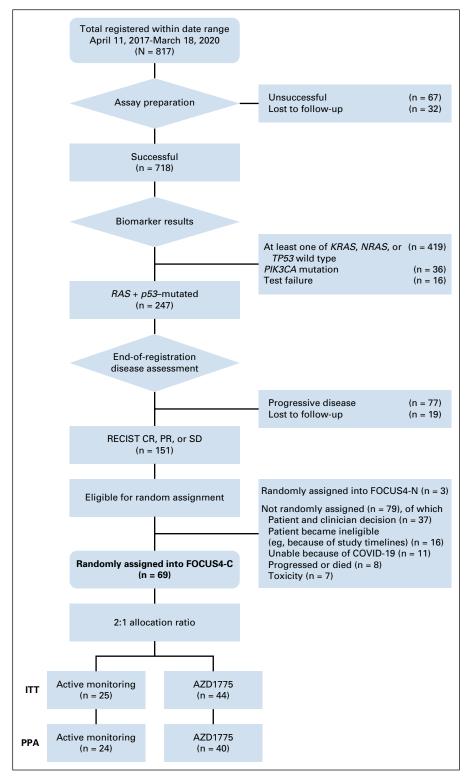


FIG 1. Flowchart of patients through the trial. CR, complete response; ITT, intention-to-treat; PPA, perprotocol analysis; PR, partial response; SD, stable disease.

#### **Trial Procedures**

Adavosertib was supplied by AstraZeneca Ltd (Cambridge, UK); packaging, labeling, and distribution were undertaken by Fisher Services (Horsham, UK). Patients randomly

assigned to adavosertib continued the drug until disease progression, death, or intolerable toxicity. The first 21 patients received adavosertib 250 mg once daily, on days 1-5 and 8-12 of a 3-week cycle. The next 23 patients received

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adavosertib 300 mg once daily, on the same schedule. Patients took an oral 5HT3 antagonist with each dose, and oral dexamethasone 4 mg was given on day 1 and day 8 of each cycle unless clinically contraindicated.

Because of the mandatory supportive medication for nausea and vomiting for which a placebo was not available, blinding was not possible, and AM was used as the control arm. Patients randomly assigned to AM followed the same follow-up schedule and remained off any other anticancer treatment until clinical or radiologic evidence of disease progression.

Patient tumor status was assessed at the treating hospital every 8 weeks by CT scan, according to RECIST, version  $1.1.^{15}$  Toxicities and symptoms were assessed locally every 4 weeks, using the National Cancer Institute's Common Terminology Criteria for Adverse Events (version 3.0). Patients remained on trial until disease progression occurred, at which point the patient was recommended to restart the same chemotherapy that was used in the induction phase. Treatment was stopped in the event of grade 3 or worse toxic effects or persistent toxicities judged medically significant or not tolerated by the patient, until the toxicity resolved to grade 1 or better.

#### Statistical Methods

A full description of the statistical methods is provided in the Data Supplement. In summary, patients were allocated to either adavosertib or AM, using a 2:1 allocation ratio by minimization with a 20% random element. All analyses were performed according to a predefined statistical analysis plan using Stata (version 16.1; Stata Corporation, TX). The primary outcome measure was progression-free survival (PFS), and the prespecified primary efficacy analysis was a per-protocol analysis (PPA) using Cox regression adjusting for minimization factors. Intention-to-treat (ITT) and unadjusted models were also performed as secondary analyses. Sample size calculations were based upon a target hazard ratio (HR) of 0.5 with 80% power and .025 one-sided alpha requiring a target of 26 PFS events in the control arm for final analysis.

#### **RESULTS**

#### **Recruitment and Patient Characteristics**

The FOCUS4 trial program ran between January 2014 and March 2020. FOCUS4-C ran between April 2017 and March 2020, during which time 817 patients were registered, of whom 718 underwent successful biomarker profiling (Fig 1 and Data Supplement). Two hundred forty-seven patients (34%) had tumors confirmed with both *RAS* and *TP53* mutations (*RAS/TP53*-mut). Of these, 151 had stable or responding disease after 16 weeks of first-line treatment and 69 were randomly assigned using a 2:1 ratio: 44 to adavosertib and 25 to AM. Of the remaining eligible 82, two chose to be randomly assigned into the concurrent FOCUS4-N trial and others chose not to be randomly

assigned into FOCUS4 for reasons such as toxicity from first-line therapy or patient-clinician choice to seek alternative pathways.

Table 1 summarizes the patient baseline characteristics. There were some minor imbalances, which are corrected for in the adjusted analysis (primary model). There were no differences in the frequency of other molecular alterations between the groups. There were no significant differences between the registration period chemotherapy regimens in the adayosertib and AM arms.

## Primary Analysis: PFS (per-protocol)

Five patients were excluded from the PPA: four did not start treatment (adavosertib arm) and one was subsequently found to have had progressive disease at the point of random assignment (AM arm). One patient was censored early when they received fluorouracil as anticancer treatment before progression (AM arm).

Within the primary PPA (n = 64), there were 40 of 40 PFS events in the adavosertib arm and 22 of 24 in the AM arm. Patients treated with adavosertib had a longer PFS than those on AM (3.61 v 1.87 months). Both unadjusted HR (0.52; 95% CI, 0.30 to 0.89; P = .022) and adjusted HR (0.35; 95% CI, 0.18 to 0.68; P = .0022) were statistically significant. Kaplan-Meier curves are provided in Figure 2.

#### PFS (ITT)

All patients were included in the ITT analysis, but four patients were censored the day after random assignment: three in the adavosertib arm (two because of patient withdrawal and one without any post–random assignment CT scan assessments) and one in the AM arm without any post–random assignment CT scan assessments.

There were 41 of 44 PFS events in the adavosertib arm and 23 of 25 in the AM arm. Consistent with the PPA, the ITT PFS analysis shows a PFS advantage with adavosertib over AM in both the unadjusted (HR = 0.55; 95% CI, 0.32 to 0.94; P=.032) and adjusted analyses (HR = 0.40; 95% CI, 0.21 to 0.75; P=.0051).

## Overall Survival (ITT)

There were 27 of 44 deaths in the adavosertib arm and 16 of 25 in the AM arm. There was no significant overall survival (OS) benefit with adavosertib compared with AM (median survival 14.0 v 12.8 months; unadjusted HR = 0.79; 95% CI, 0.42 to 1.48, P = .47; adjusted HR = 0.92; 95% CI, 0.44 to 1.94, P = .93; Fig 2).

## **Tumor Control**

Adavosertib was associated with a higher proportion of patients with disease control compared with AM (47%  $\nu$  28% at any time during the trial), including one patient with a documented partial response to adavosertib (Data Supplement).

**TABLE 1.** Baseline Patient Characteristics by Randomized Group

Characteristic	Active Monitoring (n = 25)	Adavosertib (n = 44)
Mean (SD) age, years	61.9 (12.2)	59.2 (12.8)
Sex, No. (%)		
Male	15 (60)	31 (70)
Female	10 (40)	13 (30)
Current WHO performance status, No. (%)		
0	17 (68)	35 (80)
1	8 (32)	9 (20)
Site of primary tumor, No. (%)		
Right colon	9 (36)	13 (30)
Left colon	6 (24)	13 (30)
Rectum	10 (40)	18 (41)
Current state of primary tumor, No. (%)		
Resected primary	9 (36)	23 (52)
Unresected primary	16 (64)	19 (43)
Unresected local recurrence	0 (0)	2 (5)
Timing of metastases, No. (%)		
Metachronous	4 (16)	13 (30)
Synchronous	21 (84)	31 (70)
No. of metastatic sites, No. (%)		
One	6 (24)	16 (36)
Two or more	19 (76)	28 (64)
Disease assessment at end of first-line treatment, No. (%)		
Complete response	0 (0)	1 (2)
Partial response	13 (52)	26 (59)
Stable disease	12 (48)	17 (39)
First-line treatment regimen, No. (%)		
FOLFOX	7 (28)	15 (34)
FOLFIRI	8 (32)	14 (32)
CAPOX	6 (24)	11 (25)
FOLFOXIRI	3 (12)	3 (7)
Others	1 (4)	1 (2)
PIK3CA mutation status, No. (%)		
Mutation	1 (4)	1 (2)
Wildtype	24 (96)	43 (98)
Total	25 (100)	44 (100)

Abbreviations: CAPOX, capecitabine and oxaliplatin; FOLFIRI, fluorouracil, leucovorin, and irinotecan; FOLFOX, infusional fluorouracil, leucovorin, and oxaliplatin; FOLFOXIRI, folinic acid, fluorouracil, oxaliplatin, and irinotecan; SD, standard deviation.

#### Subgroup Analyses

The impact of adavosertib versus AM on PFS was explored in prespecified subgroups (Fig 3). The most marked difference in effect was for primary tumor location (PTL):

patients with a right PTL had no PFS advantage with adavosertib compared with AM (1.87 v 1.91 months; HR = 1.02; 95% CI, 0.41 to 2.56), whereas those with a left PTL did (3.61 v 1.87 months, HR = 0.24; 95% CI, 0.11 to 0.51; interaction P = .043; Data Supplement).

This prompted an unplanned subgroup analysis of PTL on OS, and although the numbers of events were low, the interaction was even more marked (Data Supplement). Median OS was 14.1 versus 11.3 months for adavosertib versus AM in left PTL (adjusted HR = 0.37; 95% CI, 0.15 to 0.87) but was 6.5 versus 15.5 months in right PTL (HR = 6.5; 95% CI, 0.72 to 6.43; interaction P = .0032). In terms of response, 38% of right-sided adavosertib patients versus 42% of right-sided AM patients reported disease stability or response at least once while on trial, whereas for left-sided tumors, the figures were 53% versus 19%.

Patients who had responded to induction chemotherapy (v stable disease) and who had two or more metastatic sites appeared to benefit more from adavosertib, albeit to a lesser degree (interaction P value = .14 for response to induction; P = .12 for number of metastatic sites; Fig 3).

## External Analyses to Further Characterize the RAS/TP53-Mut Biomarker Population

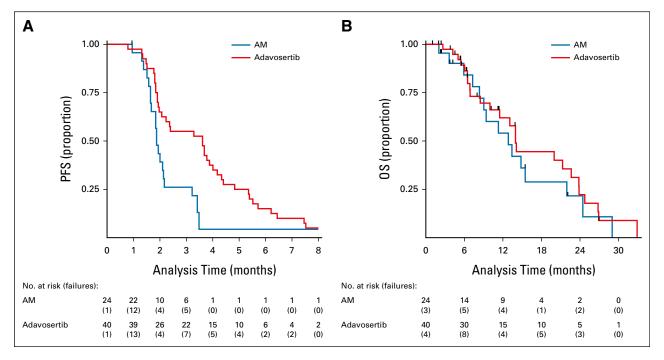
The *RAS/TP53*-mutant population has not been previously described. To understand the prognostic implication of this alteration, we analyzed the outcomes of a subset (n = 438) of patients from the FOCUS trial in whom the S:CORT consortium had analyzed a wider panel of CRC genes including *KRAS*, *NRAS*, *BRAF*, *MSI*, and *TP53*. The *RAS/RAF* wild-type group was the reference population (median OS 21.6 months). The *RAS/TP53*-mutant population is distinct from either mutation alone (*RAS* or *TP53*) and had a worse prognosis than either in isolation with a median OS of 14.9 months (HR = 2.06; 95% CI, 1.08 to 3.93; P = .028; Fig 4). This suggests that the *RAS/TP53*-mut population is a poor-prognosis subgroup but not as marked as for patients with a *BRAF* mutation or microsatellite instabilityhigh tumor.

These data are consistent with the finding that during the registration period of FOCUS4, 33% of patients in the *RAS/TP53*-mut population experienced progression during the first 16 weeks of chemotherapy. This is similar to the rate in the *BRAF*-mutant group (34% progressed) but higher than that seen in *RAS*-mutant (24%) and all wild-type (22%) subgroups (Data Supplement).

## Effect of RAS and TP53 Mutation Subtypes on Adayosertib Activity

We observed that patients with *KRAS* codon 12/13 mutations had a significant benefit from adavosertib (*P* for interaction = .014; Fig 3), whereas no detectable benefit was observed in those with *KRAS* mutations at other codons or with *NRAS* mutation. Furthermore, the interaction effects of *KRAS* subtype and of PTL on PFS may be additive as

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**FIG 2.** (A) PFS (primary analysis) in PPA population: Cox regression, adjusted for minimization factors—HR = 0.35 (95% CI, 0.18 to 0.68), P = .0022. Minimization factors: location of primary tumor (left, right, and rectum), baseline WHO performance status, baseline disease assessment, number of metastases, and first-line therapy (fluoropyrimidine, oxaliplatin or irinotecan, and monoclonal antibody). (B) OS (secondary analysis) in PPA population: Cox regression, adjusted for minimization factors—HR = 0.86 (95% CI, 0.39 to 1.86), P = .70. Minimization factors: location of primary tumor (left, right, and rectum), baseline WHO performance status, baseline disease assessment, number of metastases, and first-line therapy (fluoropyrimidine, oxaliplatin or irinotecan, and monoclonal antibody). AM, active monitoring; HR, hazard ratio; OS, overall survival; PFS, progression-free survival; PPA, per-protocol analysis.

there is a significant benefit from adavosertib within the subgroup of left PTL KRAS codon 12/13 subtypes (HR = 0.16; 95% CI, 0.05 to 0.50) and a clear disbenefit within the subgroup of right PTL noncodon 12/13 subtypes (HR = 1.56; 95% CI, 0.49 to 4.97; Data Supplement). The subtype of TP53 mutation or the co-occurrence of PIK3CA mutation did not affect outcome.

## **Toxicity and Compliance**

There was good compliance with randomized allocation, and adavosertib was generally well-tolerated (Fig 5 and Data Supplement). Compared with AM, adavosertib was associated with increased reported toxicity (≥ grade 1), most notably increased frequency of diarrhea (61% v 28%), fatigue (75% v 56%), nausea (68% v 32%), and vomiting (41% v 4%). However, the majority of such toxicity was of low grade, with 9% in the adavosertib arm reporting diarrhea of  $\geq$  grade 3, 11% fatigue, 5% nausea, and 2% vomiting, versus none of each in the AM arm. As described, during the trial, there was an increase in the dose of adavosertib from 250 mg to 300 mg. The higher dose was associated with an increased frequency of grade 3 diarrhea (14% v 4%), but otherwise the toxicity profile was similar, and with similar rates of dose modifications and delays.

#### Impact of Adavosertib Dosing

As described, during the trial, there was an increase in the dose of adavosertib from 250 mg to 300 mg. PFS was 2.2 months (HR = 0.58; 95% CI, 0.31 to 1.06) with the 250-mg dose and 3.7 months (HR = 0.47; 95% CI, 0.25 to 0.89) with the 300-mg dose; this difference was nonsignificant (P = .48; Data Supplement). Between the 250-mg and 300-mg doses, there was an increased frequency of grade 3 diarrhea (4% v 14%), but otherwise the toxicity profile was similar. There were similar rates of dose modifications between the 250-mg and 300-mg doses: dose delays (16% v 7%), dose reductions (4% v 5%), and dose omissions (19% v 17%). A swimmer plot integrating the effects of adavosertib dose, randomized group, and PTL on PFS is shown in the Data Supplement.

#### **DISCUSSION**

Here, we have reported that FOCUS4-C met its primary end point; patients with *RAS/TP53*-mutant mCRC had PFS advantage with adavosertib compared with AM following induction chemotherapy. These results are particularly encouraging as *RAS/TP53*-mutant mCRC is a poor prognostic population with limited treatment options. Adavosertib was well-tolerated at both doses evaluated.

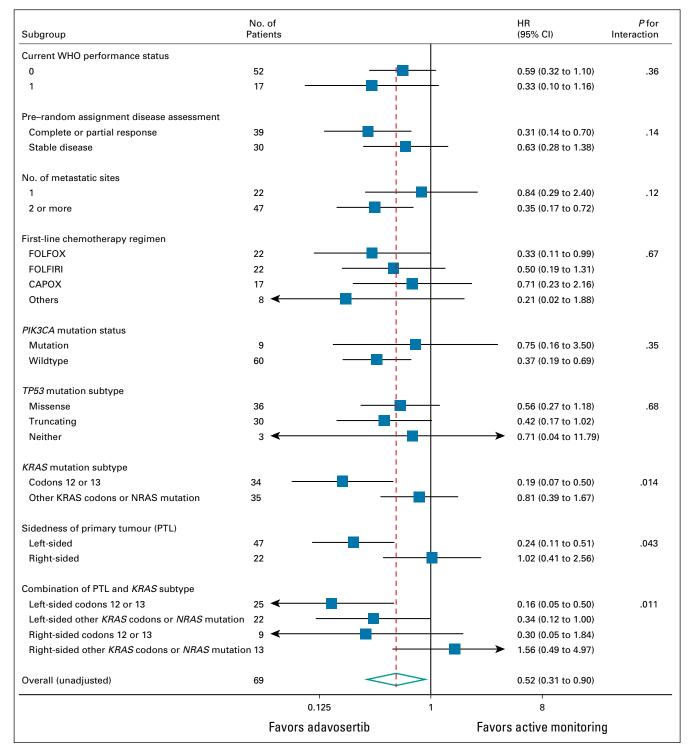


FIG 3. Subgroup analyses for PFS by intention to treat. CAPOX, capecitabine and oxaliplatin; FOLFIRI, fluorouracil, leucovorin, and irinotecan; FOLFOX, infusional fluorouracil, leucovorin, and oxaliplatin; PFS, progression-free survival; PTL, primary tumor location.

test novel agents efficiently with specified biomarker subgroups in mCRC with the multi-arm, multi-stage design allowing for an early signal of drug inactivity<sup>14</sup>; thus, any demonstrated efficacy would require further confirmatory

The overarching aim of the FOCUS4 trial program was to study to lead to practice change. FOCUS4-C represents a success of this approach, efficiently demonstrating promising activity of adavosertib within patients with RAS/ P53-mutant mCRC, and will directly influence research practice in mCRC.

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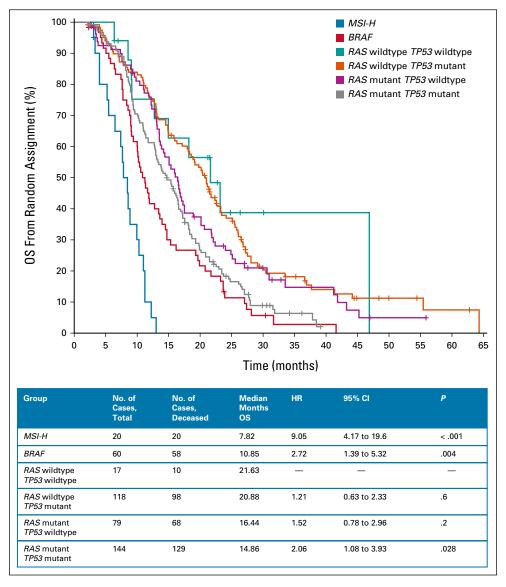


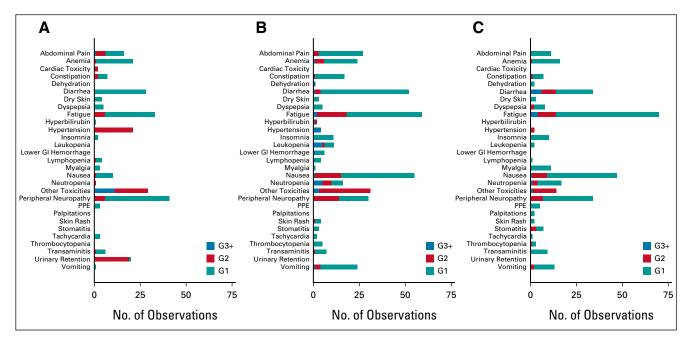
FIG 4. Prognostic impact of biomarker subgroups on OS in previous FOCUS trial. HR, hazard ratio; MSI-H, microsatellite instability-high; OS, overall survival.

The intermittent treatment strategy used in FOCUS4 follows the demonstration of no detriment in OS in the MRC COIN trial. This is now further substantiated by an individual participant data meta-analysis. <sup>16</sup> Thus, AM is an accepted standard of care following a few months of first-line therapy. FOCUS4 was specifically designed to use this window following first-line induction chemotherapy to test novel agents in specified biomarker groups, before the evolution of multiple resistance mechanisms. <sup>14</sup>

A prespecified analysis demonstrated that adavosertib activity was limited to left colon and rectal PTL, with little activity observed in right PTL. Having observed the significant subgroup effects on PFS, we investigated possible impact on OS. It is provocative to see that in the left-sided tumors, OS was significantly improved with median OS from random assignment increasing from 11.3 months to 14.1

months (HR = 0.40; 95% CI, 0.17 to 0.97). There is also a possibility of adverse effect on outcome in patients with right PTL. However, the number of patients and events was limited and thus, any conclusions need to be cautious in relation to this observed effect on OS in both subgroups. Differences in CRC by PTL are well-documented, in terms of biology, prognosis, and treatment response,  $^{17}$  but the mechanisms for differences of treatment efficacy by PTL are not well-understood.

An exploratory analysis showed that adavosertib had the most PFS effect in patients with *KRAS* codons 12/13/*TP53*-mutant tumors, with lesser activity in those with extended *KRAS*, or *NRAS* mutations; functional differences between *RAS* isoforms are documented. <sup>18</sup> Despite the small sample sizes in FOCUS4-C, the PTL and *RAS* subtypes subgroup analyses showed interactions significant at the 5% level.



**FIG 5.** Cumulative reported toxicity, within FOCUS4-C treatment groups and with initial AZD1775 doses separated: (A) active monitoring (n = 25), (B) AZD1775 250 mg (n = 23), and (C) AZD1775 300 mg (n = 21). G, grade; PPE, palmar plantar erythema.

Although these subgroup analyses provide provocative results, we lack a mechanistic explanation for these differences in adavosertib effect; ongoing translational work shall investigate this. We would recommend that further clinical development of adavosertib in the *RAS/TP53*-mut mCRC population should not be limited by PTL or *RAS* subtype but should include close monitoring of patients with right PTL and extended *RAS* mutations to ensure that neither futility nor detriment are observed.

Although the clinical implications of the *RAS/TP53* mutation in mCRC are not well-studied, each alteration is individually well-characterized. Here, we have shown that the double *RAS/TP53*-mutant subgroup carries a moderately poor prognosis (Fig 4) and appears to confer a worse prognosis than either mutation in isolation. This biomarker subgroup has thus shown distinct prognostic and therapeutic relevance and so merits further study in translational work, existing data sets, and ongoing therapeutic trials in mCRC.

Adavosertib has demonstrated an acceptable safety profile; the main toxicity was diarrhea. Efficacy was noted at both the 250-mg and 300-mg doses, with a suggestion of additional activity with the higher dose. We would therefore recommend the 300-mg dosing to progress to further clinical studies in fit patients. However, in the treatment-refractory setting, the 250-mg dose may be more tolerable.

There are limitations to this study. We considered, and would have preferred, a placebo-controlled design; however, at the time of launching FOCUS4-C, high rates of

nausea and vomiting had been observed in other adavosertib trials and high-dose steroid antiemetics were considered necessary. For this reason, both clinicians and patient representatives considered a placebo design unfeasible. Given the favorable safety data for single-agent adavosertib in FOCUS4-C, placebo-controlled design could be considered in the future. It is possible therefore that the PFS effect observed was influenced by investigator and patient preference to restart first-line chemotherapy sooner in the AM arm. However, a marked difference in effect was observed between the right and left PTL groups treated with adavosertib, suggesting a lesser effect on the primary analysis because of this potential bias. Additionally, the PFS end point was not centrally reviewed, but assessed in individual sites by RECIST criteria. A further limitation is that by testing adavosertib in the maintenance setting and requiring stability following induction chemotherapy, we have excluded the RAS/TP53 patients with the worse outcome. We therefore cannot generalize the effect of adayosertib within this entire biomarker group.

In conclusion, adavosertib (AZD1775) has demonstrated promising activity compared with AM in patients with *RAS/TP53*-mut mCRC. This treatment benefit may relate to PTL and *KRAS* subtype. Given this clear demonstration of efficacy in an RCT and acceptable toxicity profile, future clinical development of adavosertib is warranted particularly as it may represent a future treatment opportunity in this sizable population of unmet need.

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#### **EQUAL CONTRIBUTION**

J.F.S., L.C.B., and D.F. contributed equally to this work as first authors. M.S. and T.S.M. contributed equally to this work as last authors.

#### **SUPPORT**

FOCUS4-C was cofunded by the MRC/NIHR Efficacy and Mechanism Evaluation (EME) program and CRUK with funding commencing in April 2013. Drug supply and distribution of adavosertib and an educational grant were provided by AstraZeneca Ltd. The MRC Clinical Trials Unit at UCL receives core funding from the Medical Research Council within UKRI (Budget: MC\_UU\_12023/20).

#### **CLINICAL TRIAL INFORMATION**

ISRCTN#90061546

## AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

Disclosures provided by the authors are available with this article at DOI https://doi.org/10.1200/JCO.21.01435.

#### **DATA SHARING STATEMENT**

Individual deidentified participant data (including data dictionaries) can be shared upon appropriate application to the MRC CTU at any time from full publication. Study protocols and statistical analysis plan have been provided in the Data Supplement with this manuscript. Going forward, it is proposed that data will be shared with an appropriate international collaborative repository to enable future IPD meta-analysis.

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#### **ACKNOWLEDGMENT**

The full list of FOCUS4 Trial Investigators can be found in Appendix 1.

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#### **AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST**

Inhibition of WEE1 Is Effective in TP53- and RAS-Mutant Metastatic Colorectal Cancer: A Randomized Trial (FOCUS4-C) Comparing Adavosertib (AZD1775) With Active Monitoring

The following represents disclosure information provided by authors of this manuscript. All relationships are considered compensated unless otherwise noted. Relationships are self-held unless noted. I = Immediate Family Member, Inst = My Institution. Relationships may not relate to the subject matter of this manuscript. For more information about ASCO's conflict of interest policy, please refer to www.asco.org/rwc or ascopubs.org/ico/authors/author-center.

Open Payments is a public database containing information reported by companies about payments made to US-licensed physicians (Open Payments).

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	Lynsey	Bourner		Sandra	Jenkins
	Richard	Bowers	-	Sai Juan	Jia
	Hollie	Brooke		Jennifer	Keith
	Ellis	Burton		Donna	Kelly
	Julie	Burton		Audrey	Leonard
	Deborah	Cooper		Gail	Lynch
	Elizabeth	Clayton		Alex	McDonald
	Jane	Eastwood		Jordan	McGill
	Aimee	Fletcher		Anne	McKillop
	Rebecca	Foster		Austin	McInnes
	Darren	Gomersall		Fiona	McQueen
	Hassan	Hameed		Nazia	Mohammed
				Paul	Mooney

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Maria	Nygren		Fay	Annison
	Shilpa	Thapar		Sharon	Armstrong
	Kirsty	Ross		Abimbola	Barango
	Patricia	Roxburgh		Balazs	Binnyei
	Pavlina	Spiliopoulou		Gillian	Brand
	Eileen	Soulis		Kay	Campbell
	Kirsteen	Stuart		Angie	Cheyne
	Rasheed	Syed		Michael	Christie
	Ashita	Waterston		Kathryn	Connolly
	Cheryl	Wilson		Pat	Cooper
Ysbyty Gwynedd	Catherine	Bale (PI)		Amber	Johnson
	Kelly	Andrews		Susan	Martin
	Naomi	Boyle		Celia	Meneses
	Claire	Fuller		Graeme	Murray
	John	Grant		Nicola	Price
	Emma	Hall		Sue	Rodwell
	Anna	Mullard		Mhairi	Scott
	Wendy	Saxton		Margaret	Smith
	Nick	Stuart		Bartosz	Was
	Alice	Thomas		Mehmood	Zaidi
	Linzi	Williams		Ishtiaq	Zubairi
A## 1	Rachel	Williams	Cheltenham General Hospit	Kim al	Benstead (PI)
Withybush General Hospital	Sarah	Gwynne (PI)		Jaqueline	Aberdeen
	Maung	Moe (PI)		Rehana	Bakawala
	Fawwaz	Arikat		Sarah	Beazer
	Denisa	Asandei		Colin	Binks
	Sandra	Evans		Lucy	Blake
	Eirianydd	Garrard		Bethan	Cartwright
	Sophie	Glynn-Williams		Samuel	Croly
	Colette	Griffiths		Lin	Crossley
	Rachel	Hughes		Rachel	Durrant
	Catherine	MacPhee		David	Farrugia
	John	Murphy		Janet	Forkes
	Kirsty	Pope		Emma	Gilbert
	Rocio	Riba	-	Fabrizio	Mauri
	Sally-Ann	Rolls	-	Elaine	Pratten
	Abigail	Taylor	-	Elisabeth	Read
	Carol	Thomas	-	Nick	Reed
	Helen	Thomas		Rachel	Sayers
	Vallipuram	Vigneswaran		Neil	Shepherd
Aberdeen Royal	Leslie	Samuel (PI)	-	Stephen	Shepherd
Infirmary		. , ,		(continued on following	

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Jennifer	Smith		Emma	Robjohns
	Sarah	Stanley		Patrick	Sarsfield
	Catherine	Stuart-Grumbar	_	Ingrid	Seath
	Bilal	Торіа		Shirley	Todd
	Kate	Trigg-Hogarth		Jane	Thompson
Clatterbridge	Nasim	Ali (PI)		Fiona	Walters (nee Hall)
Centre for				Claire	Webb
Oncology	\M/a ala	V -1;-1	_	Julia	Weston
	Wesley	Artist	Southampton	Tim	Iveson (PI)
	Shaker	Abdallah	General Hospital		
	Alexandra	Bailey		Liane	Armstrong
	Danielle	Campbell		Andrew	Bateman
	Maggie	Cantrell		Adrian	Bateman
	Joanne	Cliff (nee Mooney)		Emma	Brown
	Thomas	Davies		Holly	Burton
	Helen	Flint		Tracey	Callen
	Amy	Ford		Bethany	Caruana
	Barbara	King		Caroline	Chau
	Ayman	Madi		Tracey	Day
	Samah	Massalha		Efe	Evbuomwan
	Laura	McAllister		Meg	Gale
	Amir	Montazeri		Julie	Gwilt
	Joanne	Mullen		Sara	Hosseini-Moein
	Julie	O'Hagan		Alice	Johnson
	Anna	Olsson-Brown		Leah	Long
	Katharine	Pelton		Steve	McKenzie
	Kelly	Richardson		Charlotte	Rees
	Sandra	Robinson		Rasha	Said
	Joseph	Sacco	University College	John	Bridgewater (PI)
	Sarah	Stuart	Hospital		
	Hollie	Wilson		Adrienne	Abioye
	Pembe	Yesildag		Mahfuja	Ahmed
	Mariah	Zavery		Shamima	Akther
Royal Devon and	Melanie	Osborne (PI)		Maise	Al Bakir
Exeter Hospital				Adelaide	Austin
	Kizzy	Baines		Holly	Baker
	Tamika	Chapter		Jaytee	Barnett
	Elizabeth	Davey		Nina	Bason
	Susan	Downer		Isabelle	Brown
	Dawn	Edwards		Alexa	Childs
	Theresa	Lawless		Louise	Coyle
	James	Leavy		Patricia	Danaswamy
	Mark	Napier		Kanishka	Dissansayke
	(continued in next co	lumn)		continued on following	

lospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Rosina	Donovan		Paula	Botham
	Lola	Enemuwe		June	Carr
	Victor	Eneh		Louise	Devlin
	Gabrielle	Gould		Katie	Douglas
	Todd	Gumbleton		Grainne	Dunn
	Selina	Gurung		Mohammed	El-Abdullah
	Gemma	Hector		Lynn	Glass
	Sonya	Hessey		Kirsteen	Hamill
	Daniel	Hochhauser		Susan	Hastings
	Sabrina	Holohan		Rebecca	Heron
	Michelle	Hung		Chloe	MacDonald
	Georgios	Imseeh		Steven	Marshall
	Adoracion	Jayme		Laura	Miller
	Sarah	Kerr		Geradline	O'Dowd
	Khurum	Khan		Aqilah	Othman
	Jennifer	Laude		Diana	Park
	Xiao	Lu		Angela	Scullion
	Gina	Margai		Denise	Vigni
	Katie	Matthews		Kai	Yahya
	Eman	Mohamad	Charing Cross	Harpreet	Wasan (PI)
	Fatima	Mohamed	Hospital		
	Sam	Morris		Thalia	Afxentiou
	Anna	Nikopoulou		Riz	Ahmed
	Mayur	Patel		Melloney	Allnutt
	Maria	Power		Gareth	Barker
	Prakash	Rao		Abigail	Caldow
	Manuel	Rodriguez-Justo		Jolene	Carioni
	Derya	Sahin		Sarah	Chilcott-Burns
	Kai Keen	Shiu		Andrea	Davis-Cook
	Luke Owen	Steventon		Yomi	Fatola
	Mark	Sunga		Chee	Goh
	Hinesh	Tailor		Dorothy	Gujral
	Anisa	Tariq		Gillian	Hornzee
	Varji	Thayalan		Eleni	Josephides
	Jennifer	Thomas		Charlotte	Kelly
	Christopher	Wanstall		Daleep	Kumar
	Kristian	Warnes		Priya	Limbu
	Christopher	Whitton		Luzviminda	Llemit Ramos
	Georgina	Wood		Charles	Lowdell
onklands	Lisa	Rogers (PI)		Sophia	Magwaro
Hospital	2.00			Rochelle	McIntyre
	Anne	McKillop (PI)		Philippa	Nutkins
	Ashita	Waterston (PI)		Shola	Ogegbo

lospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Anna	Osei-Kofi		Lai Mun	Wang
	Susan	Ramsey		Andrew	Weaver
	Pippa	Riddle		Sandie	Wellman
	Amalia	Saucan		Anthony	Wilson
	Helen	Saxby		Rebecca	Wiltshire
	Chantelle	Simpson		Martha	Woodward
	Aspa	Spyrou		Kirsten	Wynn
	Kirsty	Tunna	Leicester Royal	Anne	Thomas (PI)
	lman	Yahya	Infirmary		
	Adrian	Zebrowski		Will	Steward (PI)
hurchill Hospital,	Tim	Maughan (PI)		Elizabeth	Andrzejewski
Oxford				Tracey	Alexander
	David	Badcock		Sarah	Attridge
	Magdalena	Benysek		Julie	Barlow
	Rosita	Broderick		Theresa	Beaver
	Anne	Butterfield		Amy	Branson
	Evelyn	Chan		Meera	Chauhan
	Philip	Charlton		Aurora	Del Pozo
	David	Church		Hadia	Haque
	Richard	Cousins		Hannah	Holdsworth
	Louise	Cowen		Rahima	Ibrahim
	Joanne	Davies		Chinenye	lwuji
	Steven	Davis		Mohammed	Karolia
	Alfonso	Gonzalez Blas		Lydianne	Lock
	Will	Goodman		Mohammed	Mahgoub
	Nikki	Hayward		Adrian	Nicholson
	Clare	Jacobs		Ahmed	Osman
	Patrycja	Jastrzebska		Katherine	Perkins
	Evanthia	Komninidou		Sarah	Porter
	Jonathan	Lau		Thiaghrajon	Sridhar
	Carolina	Lepiato		Judith	Underwood
	Clare	Marken		Balaji	Varadhan
	Kerrie	Marston		Julia	Walker
	Mark	Middleton		Kevin	West
	Ann	Murphy		Joanna	Wood
	Rebecca	Muirhead	Raigmore Hospital	Walter	Mmeka (PI)
	Adrian	Nicholson		Anglise	Addison
	Robin	Peach-Toon		Seonaid	Arnott
	Navin	Pol		Karen	Callum
	Sally	Rich		Denise	Campbell
	Nicola	Stoner		Fiona	Campbell
	James	Wakelin		Kay	Kelly

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Alison	Macdonald		Michelle	Tingley
	Angela	Macgregor		Linzi	Wilson
	Carol	Macgregor	Princess Alexandra	John	Bridgewater (PI)
	Zoe	Maciver	Hospital (Harlow)		
	Laura	Maclennan		Gemma	Cook
	Jude	Madeleine		Amelia	Daniel
	Melanie	McIlroy		Venkatesh	Gajapathy
	Mary	McKenzie		Evelyn	Holmes
	Neil	McPhail		Tayo	Jaiyesimi
	Alison	Nicholls		Joanne	Kellaway
	Marion	Paterson		Teresa	Light
	Leslie	Samuel		Lucinda	Melcher
	Georgina	Simpson		Cait	Rees
	Glenda	Sinclair		Vasi	Sundaresan
	Feng Yi	Soh	Royal Surrey County Hospital	Tony	Dhillon (PI)
	Grant	Stenhouse	County Hospital	Mazhar	 Ajaz
	Joan	Stewart		Nawa	Amin
	Una	Taylor		Humyraa	Aziz
	Zoe	Urquhart		Izhar	Bagwan
/ictoria Hospital (Kirkcaldy)	Sally	Clive (PI)		Catherine	Blake
<u> </u>	Brian	Adamson		Fiona	Butler
	Julie	Aitken		Penny	Champion
	John	Brush		Karen	Chan
	Rebecca	Cain		Sebastian	Cummins
	Lesley	Cargill		Tineke	Edmunds
	Shona	Cheyne		Sharadah	Essapen
	Clare	Cliff		Andrew	Furness
	Hazel	Cree		Laura	Gordon
	Karen	Gray		Di	Grainger
	Sophie	lwanikiw		Helen	Graves
	Fiona	Johnston		Imogen	Heenan
	Alastair	Matthews		Kirsty	Horwood
	Wendy	McCorry		Daniel	Jennings
	Catriona	Mclean		Natasha	Kamboh
	Fiona	Murdoch		Aga	Kehinde
	Ibrahim	Nawroz		Karla	Lee
	Julie	Penman		Sibylle	Lintott
	Anna	Scott		Gaybrielle	Livingstone
	Maria	Simpson		Cheryl	Marriott
	Deepak	Subedi		Catherine	Medcalf
	Jennifer	Tait	·	Aruna	Medisetti
	3011111101	IUIL			

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Gayathri	Nagarajan		Deborah	Abrams
	Sarah	Oakes		Debbie	Austin
	Sue	Sargent		Carlos	Gonzalez
	Alexandra	Stewart		Matthew	Howlett
	Hasina	Thandar		Natalie	Lloyd
	Claire	Thompson		Rita	Ng
	Katharine	Webb		Paul	Ridley
	Rosalyne	Westley		Kirubah	Selvaraj
	Julia	Whittle		Liz	Sherwin
	Julie	Wilkinson		Bamini	Sivarajah
	Rebecca	Wills		Susan	Upson
St Helens Hospital	Zahed	Khan (PI)		Angharad	Williams
	Rachel	Cassidy		Jason	Wong
	Jenny	Cotton	Royal Hampshire	Luke	Nolan (PI)
	Lisa	Dobson	County Hospital	Louiso	Beattie
	Nicola	Hornby	-	Louise Julie	Conti
	Sheila	Kelly	-		
	Amanda	McCairn		Duncan	Cooke
	Jeanette	Ribton		Victoria	Corner
	Michelle	Robinson		Adrienn	Fazekasne Fulep
	Carol	Ross		Angela	Frith
	Victoria	Thomas		Julie	Gwilt
Chesterfield Royal Hospital	Vanessa	Wilshaw (PI)		Samantha Liz	Hammond Happle
	 Ibrahim	Al-Modaris		Lesley	Hollister
	Rebecca	Clark	-	Roger	Hudson
	Aurora	Del Pozo	-	Abigail	Hughes
	Alice	Dewdney	-	Lauriane	Kerwood
	Nicky	Ford		Matthew	Pitt
	Rachel	Gascoyne		Balvinder	Shoker
	Neeta	Gogna		Rao	Vuyyuru
	Charlotte	Hoult	Peterborough City	Catherine	Jephcott (PI)
	Emma	Hudson	Hospital	Odificilite	Jepheoti (11)
	Kelly	Pritchard		Terri-Anne	Baker
	Martin	Shepherd		Helen	Bowyer
	Lesley	Stevenson		Kerrie	Cavanagh
	Danesh	Taraporewalla		Rebecca	Chilvers
	Julie	Toms		Marilyna	Chong
	Katie	Wallace		Laura	Costello
	Julie	Whitehead		Abigail	Hollingdale
	Lucinda	Wilson		Steph	Lawrence
pswich Hospital		Srinivasan (PI)		Heather	Maccoll
ipswich Hospital	Gopalakrishnan Zoltan	Szucs (PI)		Carla	Martino
	continued in next co		(6	continued on following	

Participating Hospitals in Descending Order of the Number of Patients Registered With All Staff Listed (N = 2,076) (continued)

Surname (principal

lospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	investigator [PI]
озртан	Claire	Palombo		Julie	Froud
	Stuart	Richmond		Bojidar	Goranov
	Richard			Irene	Harvey
		Skells		Maggie	Kalita
	Laura	Simon		Sarah	Kingdon
	Claire	Snowden		Mike	Marner
	Lisa	Wilde		Laura	Marks
	Louise	Wilmer		Susan	McFarlane
alderdale Royal Hospital	Jo	Dent (PI)		Chelsea	Morton
	Mohammad Irfan	Alam		Anna	Mucha
	Nick	Brown		Sarah	Prance
	Nicky	Daker		Olivia	Reed-Poysden
	Sam	Dale		Peter	Sankey
	Denise	Hancock		Helen	Smith
	James	Harris	Macclesfield	Victoria	Lavin (PI)
	Lisa	Horner	District General Hospital		
	Jeremy	Hyde		Ganesh	Radhakrishna (PI)
	Rebecca	Jenkins		Catherine	McBain (PI)
	Christopher	Knight		Victoria	Adinkra
	Mandy	Madigan	-	Dane	Bradwell
	Adam	Mawer		Lisa	Brookes
	Belinda	McLean	-	Helen	Burns
	Sabiha	Ravat		Nicola	Dawson
	Hannah	Riley	-	Catherine	Fenson
	Jodie	Rowan		Lisa	Hardstaff
	Simone Deborah	Ryan		Abbi	Henderson
	Lisa	Shaw	-	Christy	Henderson
	Selina	Shaw		Pippa	Hill
	Kathryn	Smith		Debra	Jowle
	Christine	Turner		Mark	Lawrence
	Georgina	Turner		Joanna	Longden
	Hayley	Webster		Nicola	Lunt
	Tracy	Wood		Marilyn	McCurrie
erriford Hospital	David	Sherriff (PI)		Karen	Rotchell
2.2oop.tan	Rebecca	Aaron		Barbara	Townley
	Bridget	Aire		Helen	Wassall
	Baffour	Amo-Takyi		Julie	Whitehead
	Erin	Brennan		Lesley	Wilkinson
	Lucy	Cadmore		-	Woodhouse
	Leonie	Eastlake	Torboy District	lain	
	Laura	Evenden	Torbay District General Hospital	Nangi	Lo (PI)
				Michele	Allison
	Kay	Facey		continued on following	
	Olivia	Fraser		Johnnaca on Iollowill	D Mage/

lospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Kenneth	Almedilla		Sarah	Clark
	Emmie	Arbury		Joseph	Gallagher
	Lauren	Blunt		Svitlana	lyevkova
	Jo	Blurton	-	Rashmi	Jadon
	Catherine	Brookman		Catherine	Jephcott
	lan	Buley		Natalie	Jones
	Shelley	Chamberlain		Hannah	Loveday
	Stacey	Davies		Jane	Macdonald
	Angela	Foulds		Betania	Mahler-Araujo
	Meadow	Fisher-Crisp	-	Debra	Mansergh
	Joanne	Garfield-Smith		Ultan	McDermott
	Petra	Gee		Lindsay	Piper
	Caera	Good		Amy	Strong
	Hannah	Griffin		Catherine	Thorbinson
	Andrew	Harford-Brown		Saji	Victor
	Prithvi	Jampana		Naval	Vyse
	Ingrid	Koehler		Amanda	Walker
	Tyler	Lowe		Emma	Wong
	Sally	Maddison		Zsuzsa	Zaborszky
	Mitchell	McMillan	Guy's Hospital	Paul	Ross (PI)
	Louise	Medley	(London)		
	Lyn	Micklewright		Samantha	Barrett
	Louise	Paatz		Eva	Batovska
	Maeve	Pomeroy		Jessica	Brady
	Helen	Randall		Maribel	Boyce
	Fleur	Rogers		Laura	Camburn
	Lorraine	Thornton		Lorna	Caplis
	Christine	Tsang		Noan Minh	Chall
	Elaine	Vandecandalaere		Jason	Chow
	Sarah	Wright		Chi Yee	Chung
ddenbrooke's	Hugo	Ford (PI)		Sophie	Clark
Hospital				Sarah	Cleary
	Athar	Ahmad		Victoria	Donovan
	Alexandra	Azevedo		Sandra	Esteban Moreno
	Lesley	Bennett		Adrienn	Fazekasne Fulep
	Elizabeth	Blake		Lucy	Featherstone
	Mark	Bolton		Michael	Flanagan
	Rebecca	Bradley		Laura	Green
	Jane	Bushen		Sara	Hulf
	Joanna	Calder		Arun	Karnad
	Anita	Chhabra		Sara	Kazemzadeh
	Kathy	Chin		Vevangaune	Ketjiperue

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Choi Chin	Lau		Alistair	Ellis-Jones
	Nick	Maisey		Emma	Hall
	Simranjit	Mehta		Rachel	Hughes
	Ngozi	Muoneke		Ravi	Kodavatiganti
	Theodorah	Nago		Arwel	Lloyd
	Rita	Njoku		Bethan Wyn	Owen
	Vitalis	Nwokorie		Beryl	Roberts
	Temi	Olusi		Charley-Anne	Rutter
	Kishen	Patel		Jane	Stockport
	Amy	Quinn		Gemma	Szabo
	Catherine	Rogers		lan	Walker
	Hannah	Rush		Claire	Watkins
	Susie	Slater		Glesni	Williams
	Anita	Soma		Linzi	Williams
	Chara	Stavraka	Glan Clwyd	Simon	Gollins (PI)
	Harriet	Waine	Hospital		
	Sally	Walker		Elizabeth	Allan
St George's	Fiona	Lofts (PI)		Jill	Andrews
Hospital				Kelly	Andrews
(London)	Doroid	Alrifa		Lisa	Ashley
	Doraid			Llinos	Davies
	Nia	Alsamarrai		Rachel	Davies
	Jason	Chow		Clair	Domeney
	Alice	Dainty		Sarah	Evans
	Lorette	Ffolkes		Emma	Hall
	Caroline	Finlayson		Jane	Heron
	Claire	Gilmartin		Ravi	Kodavatiganti
	Anne	Haldeos		Joanne	Lewis
	Sam	Hollingworth		Arwel	Lloyd
	Geoffrey	Howell		Carey	Macdonald-Smith
	Robert	Ingham		Claire	McGregor
	Kay	Laurent		Bethan Wyn	Owen
	Vitalis	Nwokorie		Tracy	Parry-Jones
	Antonio	Pesino		Fiona	Redmond
	Mark	Quarrell		Beryl	Roberts
	Agne	Sekmokaite		Charley-Anne	Rutter
	Jesusa	Toledo		Libby	Thackray
Vrexham Maelor Hospital	Simon	Gollins (PI)		lan	Walker
i iospilai	Stacy	Ackerley		Jill	Westlake-Guy
	Stacy			Linzi	Williams
	Achrof	Alkhaldi			
	Ashraf Kelly	Alkhaldi Andrews		Stephanie	Wynne

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
ames Cook University Hospital	Nick	Wadd (PI)	Royal Cornwall Hospital	Richard	Ellis (PI)
Hospital		_		Linda	Allsop
	Andrea	Boyce		Nicholas	Ashley
	Alison	Chilvers		Kerry	Atkinson
	Anthony	Donnelly		Nigel	Bailey
	Helen	Dunn		Thea	Barlow
	Vicky	Hanlon		Kayleigh	Bennett
	Charlotte	Jacobs		Carolyn	Brode
	Steven	Liggett		Thomas	Cornell
	Craig	Mower		Alexander	Dengler
	Lisa	Peacock		Emma	Duley
	Jacqueline	Richards		Sophia	Eloi
	Agnieszka	Skotnicka		Caroline	Goddard
	Danielle	Sweeney		Aaron	Gould
	Jane	Thompson		Anne	Griffiths
	Hans	Van der Voet		Karina	Harris
	Gill	Wheater		Peter	Helliwell
	David	Wilson		Claire	Hill
	Jason	Wong		Louise	Johns
Poole Hospital	Amelie	Harle (PI)		Tinnaya	King
	Tamas	Hickish (PI)		Samantha	Lomax
	Michael	Adrio		Kirsty	Maclean
	Maria	Alban		John	Madine
	Julian	Alexander		Joe	Mathew
	Lyn	Allen		John	McGrane
	Mary	Apps		Fiona	Minear
	Beth	Aubrey		Sharon	Moore
	Helen	Bradley		Anna	Oakes
	Savina	Elitova		Caroline	Parnell
	Daniel	Fielding		Kerena	Partridge
	Maxine	Flubacher		Sallyanne	Platt
	Deborah	Forster		Kirsty	Prout
	Melanie	Foster		William	Pynsent
	Louise	Heckford		Rebecca	Rogers
	Jill	Hobson		Jenifer	Row
	Hannah	James		Laura	Royle
	Min Yee	Lee		Johanna	Skewes
	Helen	Morling		David	Smith
	Victoria	Osborne		David	Snell
	Sharon	Power		Luke	Townley
	Victoria	True	Royal Free Hospital		Krell (PI)
	Craig	Vincent	Troyal Free Hospital	Astrid	Mayer (PI)
	Roger	Wheelwright			
	(continued in next co		(c	ontinued on following	g page)

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Tahmin	Ahmed		Edwin	Cooper
	lan	Clark		Sarah	De Bruijn
	Jen	Fraser-Fish		David	Donaldson
	Roopinder	Gillmore		Tracey	Duckett
	Sara	Hamilton		Adam	Edwards
	Ben	Marks		Shirley	Fox
	Leah	Meaden		Karen	Flynn
	Aarti	Nandani		Michelle	Kotze
	Tesha	Suddason		Michaela	Nock
	Sharon	Thompson		Jess	Perry
	Elizabeth	Woodford		Lucy	Pippard
outh Tyneside District Hospital	Ashraf	Azzabi (PI)		Kerry	Rennie
District Frospitar	Amy	Burns			Rowsell
	Kumud	Jain			Smith
	Judith	Moore			Thomas
	Ruth	Tindle		Barbara	Williams-Yesson
t Bartholomew's	David	Propper (PI)	Lincoln County Hospital	Zuzana	Stokes (PI)
Hospital (London)				Edwin Sarah David Tracey Adam Shirley Karen Michelle Michaela Jess Lucy Kerry Amber Rufus Lesley Barbara Junty Zuzana Antoinette Suzanne Sarah Jayne Sarah Olesya Annette Kathryn Carol Maryanne Rhiannan Manuel Thomas Andrew Caroline Ruth Alyson	Adu
(LONGOTT)	Waheeda	Abida			Archer
	Hayley	Blackgrove		Sarah	Bell
	Joanne	Chin-Aleong		Jayne	Borley
	Nikolaos	Diamantis		Sarah	Coombs
	Resmi	Jayachandran		Olesya	Francis
	Sumaiya	Kamora		Annette	Hilldrith
	Cheryl	Lawrence		Kathryn	Hoare
	Alia	Mahboob		Carol	Lockwood
	Juan	Navarro		Maryanne	Okubanjo
	Tanjil	Nawaz		Rhiannan	Pegg
	Pratistha	Panday		Manuel	Ruiz-Echarri
	Hannah	Payne		Thomas	Sheehan
	Stephen	Russell		Anuradha	Sheth
	Sarah	Slater		Andrew	Sloan
eovil District	Andrew	Allison (PI)		Caroline	Taylor
Hospital	Andrew	Allisori (F1)		Ruth	Thoy
	Erica	Beaumont (PI)		Alyson	Wilson
	Matthew	Sephton (PI)	Maidstone Hospital	Mark	Hill (PI)
	Joanna	Allison		Doraid	Alrifa
	Zenaida	Armstrong		Elizabeth	Angus
	Claire	Barron		Paulette	Basham
	Nigel	Beer		Lisa	Brown
	Kate	Beesley		Tracey	Chambers
	(continued in next co			Alison	Davison

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Jackie	Evans		Rosalind	Roberts
	Sanjina	Kathuria		Maria	Scott
	Samantha	Kestenbaum		Rafael	Silverman
	Tiana	Kordbacheh		Ananth	Sivanandan
	Satish	Kumar		Tania	Slater
	Barbara	LeBrocq		Anita	Stevenson
	Gemma	McCormick		Richard	Swinden
	Christos	Mikropoulos		Jackie	Worville
	lan	Pamphlett		Georgina	Walker
	Joanne	Patterson		Andrew	Wright
	Caroline	Rodger	Hinchingbrooke	Cheryl	Palmer (PI)
	Holly	Slater	Hospital		
	Charlotte	Stevens		Shilamba	Bramham
	Jeff	Summers		Sue	Donnelly
	Alicia	Synowiec		Simon	Duke
	Katy	Taylor		Vanessa	Goss
	Lisa	Tribe		Beverley	Haynes
Nottingham	Cristina	Lopez Escola (PI)		Rebecca	Lam
University		,		Elizabeth	Lee
Hospitals				Sarah	Littlechild
	Rebecca	Ashton		Adam	McGeoch
	Suha	Atabani		Suzanne	Miller
	Alex	Blades		Agnieska	Osmanska
	Emma	Blades	North Middlesex	John	Bridgewater (PI)
	Lauren	Blackburn	Hospital		
	Pauline	Brookes		Ernesto	Balaguer-Ruiz
	Eliot	Chadwick		Girish	Bhome
	Caroline	Coulson		Moira	Durdy
	Michelle	Cunnell		Lorraine	Hurl
	James	Donworth		Shardul	Kulkarni
	Jade	Eggleton		Simranjit Kaur	Mehta
	Susan	Elliott		Lucinda	Melcher
	Joanne	Hobbs		Julia	Rees
	Shaymaa	Hosni		Jamila	Roehrig
	Laura	Kirk		Rahi	Shah
	Emma	Marshall		Chloe	Van Someren
	Balwir	Matharoo-Ball	Queen Alexandra	Ann	O'Callaghan (PI)
	Kayleigh	Mills	Hospital		
	Jamie	Mills		Oluwatobi	Adeagbo
	Jeanette	Mulhurn		Suhail	Baluch
	Karen	Newcombe		Kathy	Blight
	Vanessa	Potter		Sherilee	Cook
	Tin	Sang-Tsang		Heather	Cuell

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principal investigator [PI])
	Tracey	Dobson		Sonya	Goriah
	Mya	Gyi		Praba	Gupta
	Antony	Higginson		Ann	Hewins
	Samuel Luke	Hill		John	Murphy
	Chloe	Holden		Zohra	Omar
	Tracey	Lee		Bryan	Phillips
	Jayne	McCartney		Meena	Raj
	Badrriyya	Mohamedali		Kelly	Reed
	Sethupathi	Muthuramalingam		Rocio	Riba
	Andras	Nagy	Royal Albert	Francisca Marti	Marti (PI)
	Eleanor	Taylor	Edward Infirmary		
	Mary	Wands		Elena	Takeuchi (PI)
	Robert	Williams		Jennifer	Cannon
	Carole	Wragg		Kate	Chilman
Weston General	Stephen	Falk (PI)		Shien	Chow
Hospital				Louise	Devereaux
	Paola	Di Nardo (PI)		Alison	Doran
	Marjorie	Tomlinson		Diane	Forrest
	Kathy	Beard		Karen	Moss
	Sandra	Beech		Monica	Patel
	Hannah	Berry		Angela	Power
	Debbie	Coles		Wendy	Stevens
	Donna	Cotterill	Sunderland Royal	Ashraf	Azzabi (PI)
	Harvey	Dymond	Hospital		
	Symeon	Eleftheriadis		Hayley	Anderson
	Rajesh	Gamare		Rod	Beard
	Christine	Graham		Jane	Cole
	Serena	Hilman		Michelle	Edwards
	Sarah	Kidd		Adam	Hassani
	Denise	Leighton-Price		James	Henry
	Hugh	Lloyd-Jones		Vivienne	Hullock
	Andrew	McKendrick		Stephen	Laybourne
	Kathryn	Munday		Paula	Newton
	Vivienne	Pixton		Rachel	Pearson
	Glenn	Saunders		lan	Pedley
	Ed	Sheffield		lan	Pepley
	Dawn	Simmons		Melanie	Robertson
	Axel	Walther		Fiona	Wakinshaw
	Rachel	Warinton		Kathryn	Wright
	Tom	Wells	Basingstoke and	Charlotte	Rees (PI)
Glangwili General	Mau-Don	Phan (PI)	North Hampshire Hospital		
23.10.di	Samantha	Coetzee	Ποσρικαι	Louise	Beattie
	(continued in next co		-	ontinued on following	

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principal investigator [PI])
	Victoria	Corner		Shiv	Gayadeen
	Abigail	Edwards		Rob	Glynne-Jones
	Adrienn	Fazekasne Fulep		Marcia	Hall
	Angela	Frith		Rakhi	Jain
	Julie	Gwilt		Colleen	Murray
	Liz	Happle		Julie	Russell
	Roger	Hudson		Waqar	Saleem
	Andrew	Jackson		Anand	Sharma
	Lauriane	Kernwood		Margaret	Stone
	Lauriane	Kerwood		Harsha	Vara
	Kathryn	Leach	Queen Elizabeth	Gary	Middleton (PI)
	Emma	Magras	Hospital		
	Asmat	Mustajab	(Birmingham)	Cabia	Alaban
	Christina	Narh		Sabia	Akhtar
	Pennie	Porter		Amisha	Desai
	Arun	Selvaraju		Colm	Forde
	Jackie	Smith		Kam	Gareja
	Claire	Williams		Sharon	Hackett
Forth Valley Royal	Dawn	Storey (PI)		Sam	Hopkins (nee Poole
Hospital		,		Mary	Kotadia
	Joanne	Blackburn		Victoria	Kunene
	Stephanie	Brogan		Catherine	Prest
	Raj	Burgul		Helen	Preston
	Eilidh	Henderson		Donna	Smith
	Jane	Keddie		Phillipe	Taniere
	Linnet	McGeever	Queen's Hospital Burton	Manjusha	Keni (PI)
	Kaye	McIlvar	Buiton	Ann	Adams
	David	McIntosh	-	Mosan	Ashraf
	Caroline	Mcleary		Jo	Burns
	Lynn	Prentice	-	Helen	Cox
	Annette	Riley	-		
	Joanne	Robinson		Katy	English Fleet
	Anne	Todd		Annette	
	Patricia	Turner		Sarah	Hathaway-Lees
	Sally	Young		Elizabeth	Kemp
Mount Vernon	Mark	Harrison (PI)		Hayley	Lewis
Hospital			-	Clare	Mewies
	Farhan	Ahmed	-	Jennifer	Moyes
	Nicola	Anyamene	-	James	Price
	Nicky	Barnes		Scott	Sanders
	Neel	Bhuva	-	Adrian	Smith
	Sam	Bosompem		Alison	Tilley
	Kari	Evans	(	continued on following	g page)

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
Russells Hall	Ankit	Jain (PI)		Julie	Turner
Hospital				Nia	Viney
	Simon	Grumett (PI)		Dawn	Withers
	Joann	Atkinson	University Hospital	Vanessa	Potter (PI)
	Daniel	Bull	Coventry		
	Donna	Cleal		Jason	Allen
	Lesley	Edwards		Senthil Kumar	Athmanathan
	Kath	Harrow		Rachel	Bazeley
	Stacey	Jennings		Susan	Bird
	Lucy	Kadiki		Yasmin	Brough
	Karen	Kanyi		Maggie	Brown
	Sally	Keates-Porter		Dannielle	Burgess
	Pek	Keng-Koh		Luanne	Carey
	Margaret	Marriott		Philippa	Clark
	Julie	Matthews		Peter	Correa
	Karen	McGarry		Kishore	Gopalakrishnan
	Vanessa	Moore		Cheryl	Hunter
	Andrew	Moores		Sian	Kempster
	Manesh	Patel		Mohammed	Khan
	Veena	Shinde		Fiona	McGurk
	Lucie	Smith		Jade	McKelvie
	Lucy	Smith		Lucy	Miller
	Angela	Watts		Sarah	O'Toole
Singleton Hospital	Sarah	Gwynne (PI)		Karandeepu	Pachoo
	Cristina	Lopez (PI)		Noor	Shaw
	Alya	Al-Affan		Laura	Stanley
	Philip	Bryant		Charlie-marie	Suddens
	Karen	Chesters		Rachel	Thompson
	Sharon	Davies		Maria	Truslove
	Jenna	Edwards		Linda	Wimbush
	Stuart	Evans		Jane	Wording
	Tracey	Ford	University Hospital	Madhavi	Adusumalli (PI)
	Ricky	Frazer	of North Tees		
	Judith	Gooding		David	Wilson (PI)
	Olivia	Hatcher		Alison	Chilvers
	Gillian	Jones		Helen	Dunn
	Lewis	Jones		Sarah	Essex
	Maung	Moe		Mohammad	Hegab
	Karen	Phillips		Hyder	Latif
	Euan	Pratt		Moira	Percival
	Alex	Richards		Sarah	Pitcairn
	Louise	Thomas		Lynda	Poole
	(continued in next co			Pam	Race

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Andrew	Sigsworth		Elizabeth	Reay
	Eleni Andriana	Trigka	·	Timothy	Simmons
	Helen	Wardle	_	Carole	Stobbart
	Bill	Wetherill	-	Jonathan	Stoddart
	Pauline	Leonard (PI)	-	Nichola	Waugh
Hospital			-	Hesther	Wilson
(London)	5		Leighton Hospital	Michael	Braun (PI)
	Rashidat	Adeniba		Vanessa	Adamson
	Dhili	Arul		Carole	Bennion
	Jonathan	Flor		Kim	Best
	Kavita	Kantilal		Leanne	Everall
	Xiao Lou	Lu		Julia	Gemmell
	Mulyati	Mohamed		Laura	Hanton
	Michelle	Saull		Christy	Henderson
	Nuray	Temiz		Adele	Hough
	Azmina	Verjee		Chris	
	Simon	Wan			Hough
Freeman Hospital,	Ashraf	Azzabi (PI)		Cyndy	Jackson
Newcastle	0.1			Taya	Jones
	Craig	Alderson		Tracy	Larcombe
	Chris	Barron		Carolyn	Mansfield
	Michelle	Borthwick		Emma	Margerum
	Julie	Burton		Julie	Meir
	Kay	Carson		Andrew	Ritchings
	Fiona	Chapman		Paul	Simcock
	Sarah	Cook		Sarah	Tinsley
	Fareeda	Coxon		Caroline	Walker
	Sue	Farrell	Ninewells Hospital, Dundee	Sharon	Armstrong (PI)
	Elaine	Greaves	- Dundee	Jennifer	Allison
	Ahmed	Hashmi	-		
	Amanda	Henderson		Rachael	Banks
	Kathryn	Hewitt		Anne	Black
	Ben	Hood	-	Louise	Brannan
	Thomas	Jarvis		Frank	Carey
	Irene	Jobson		Shona	Carson
	Najibah	Mahtab		Helen	Cumming
	Lesley	Naik		Debbie	Forbes
	Stephanie	Needham		Audrey	Lyall
	Gemma	O'Neill		AJ	Munro
	lan	Pedley		Moira	Rogers
	Sindhu	Ramamurthy		lan	Sanders
	Zarine	Razvi		Gail	Weir
	(continued in next co		Westmorland General Hospital	David	Eaton (PI)
				Rebecca	Anderson

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principal investigator [PI])
	Syed	Asghar		Richard	Osborne (PI)
	Manal	Atwan		Pauline	Ashcroft
	Claire	Bartlett		Corrado	d'Arrigo
	Ashoke	Biswas		Maxine	Flubacher
	Jennifer	Bowler		Jackie	Gibbins
	Karen	Burns		Karen	Hogben
	Rebecca	Calvert		Arabis	Oglesby
	Amy	Ford		Andrew	Rees
	Laura	Healey		Simon	Wilsher
	Nima	Herlekar	Great Western	Sarah	Lowndes (PI)
	Maria	Kassi	Hospital		
	Lauren	Kilifin		Graham	Brown
	Jo	Kilkenny		Christopher	Clarke
	Nicola	Mackenzie		Amanda	Colston
	Aileen	Menzies		Jan	Dodge
	Helen	Morris		Eva	Fraile
	Debbie	Power		Sarah	Grayland
	Jane	Ritchie		Lesley	Haxton
	Mary	Robinson		Lawrence	John
	Vickie	Rose		Jean	Kordula
	Rachel	Simmons		Lynsey	Kyeremeh
	Andrew	Taylor		Donna	Lake
	Hilary	Thatcher		Catherine	Lewis Clarke
	Gail	Wiley		Sarah	Long
Belfast City Hospital		Coyle (PI)		Dorota	Marciniak
	Conal	Askin		Laura	McCafferty
	Ellen	Brown		Darren	McFadden
	Karen	Campfield		Sue	Meakin
	Catherine	Davidson		Chanelle	Meyer
	Michael	Hanna		Tim	Owen
	Diane	Law		Cerila	Parajes
	Alison	McKeever		Ronak	Patel
	Aine	McKeown		Suzannah	Pegler
	Damian	McManus		Caroline	Pensotti
	Linda	McNeice	<del></del>	Joseph	Stevens
	Karen	Parsons	Milton Keynes	Wasiru	Saka (PI)
	Miranda	Reid	University		
	Fiona	Tarpey	Hospital	Ann	Ahraham
	Joanne	Todd		Ann	Abraham
	Paul	Ward		Hannah	Ansell
				Sam	Bosompem
Doroot Court	Richard	Wilson		Matthew	Burnett
Dorset County Hospital	Amelie	Harle (PI)		Chris	Ford

Hospital	First Name	Surname (principal investigator [PI])	Hospital	First Name	Surname (principa investigator [PI])
	Chloe	Green		Victoria	Knight
	Sara	Greig		Tara	Lawrence
	Penni	Hawkins		Beverley	Mashegede
	Chamene	Hicks		Helen	Palmer
	Aarzoo	llyas		Kerry	Pettitt
	Charity	Masvaure		Gunjan	Phalod
	Louise	Moran		Manuel	Ruiz-Echarri
	Mala	Nathvani		Gemma	Sankey
	Cheryl	Padilla-Harris		Thomas	Sheehan
	Vijay	Patel		Rebecca	Spencer
	Shahriar Mohammed	Reza		Kinga	Szymiczek
	Syed Azhar Javed	Rizvi		Isobel	Thomas
	Abby	Skillington	Rotherham District	Joanne	Hornbuckle (PI)
	Jeannette	Smith	General Hospital		
	Oliver	Spring		Matthew	Barnes
	Heather	Thomas		Sarah	Besley
	Stephanie	Thorp		Meredyth	Harris
	Valerie	Webb		Kath	Lowe
	Dona	Wingfield		Scott	Nicol
	Christopher	Woodard		Susan	Oakley
lew Cross Hospital	Simon	Grumett (PI)		Amy	Rees
	Syed	Asghar		Charlotte	Widdop
	Vanda	Carter	Royal Bournemouth	Tamas	Hickish (PI)
	Sandeep	Dhillon	Hospital		
	Anna	Grant		Jocelyn	Ablorde
	Clare	Hammond		Omolade	Bakarey
	Kelly	Kauldhar		Rachel	Bower
	Margaret	King		Zoe	Clark
	Christine	Kirk		Nicole	Davies
	Claire	Lomas		Alison	Hogan
	Manel	Mangalika		Stephanie	Jones
	Gurminder	Sahota		Tiffany	Joyce
	Elaine	Wylde		Maria	Lane
Pilgrim Hospital	Zuzana	Stokes (PI)		Sharon	Megson
	Antoinette	Adu		Sandy	Pressdee
	Simon	Archer		Linda	Purandare
	Gloria	Barone		Taslima	Rabbi
	Jayne	Borley		Emma	Sharland
	Wendy	Deamer		Esther	Una Cidon
	Jo	Fletcher		Luke	Vamplew
	Matthew	Flook		Jasmin	Webb
		Kirkby		ontinued on following	

Surname (principal Hospital First Name investigator [PI]) Royal Marsden lan Chau (PI) Hospital (London) Helen Breeze Clifton Shirley Saoirse Dolly Sandra Esteban Moreno Lucy Featherstone Shelby Hatt Blanka Hezelova Lee Alexander Hazel Lote Lizzie Love Nnenna Ngwu Isma Rana Gihan Ratnayake Penny Rogers Clare Saffery Anna Scott Izelle Ueckermann Chloe Westrip Chau lan Abdelmalik Sally Gayahri Anandappa Joo Ern Ang Thushasa Ansari Sheila Azaiji-Benjamin Annette Bryant Shirley Clifton Richard Crux Cunningham David Sara Diffley Julie Duncan Laurice Edwards Sandra Esteban Moreno Featherstone Lucy Monika Ferencova Angela Gillbanks Kaur Sarnjeet Naila Kaudeer Shelize Khakoo Shannon Kidd Retchel Lazaro Alcausi (continued in next column)

Hospital	First Name	Surname (principal investigator [PI])	
	Hazel	Lote	
	Jacqueline	Oates	
	Bijal	Patel	
	Minal	Patel	
	Brenda	Pem	
	Sijy	Pillai	
	Clare	Saffery	
	Francesco	Sclafani	
	Gillian	Smith	
	Eleanor	Temple	
	Jan	Thomas	
	Andrea	Turner	
	Izelle	Ueckermann	
	David	Watkins	