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Subcortical Volume Trajectories across the Lifespan: Data from 18,605 healthy individuals aged 3-90 years

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Abstract

Age has a major effect on brain volume. However, the normative studies available are constrained by small sample sizes, restricted age coverage and significant methodological variability. These limitations introduce inconsistencies and may obscure or distort the lifespan trajectories of brain morphometry. In response, we capitalised on the resources of the Enhancing Neuroimaging Genetics through Meta-Analysis (ENIGMA) Consortium to examine the age-related morphometric trajectories of the ventricles, the basal ganglia (caudate, putamen, pallidum, and nucleus accumbens), the thalamus, hippocampus and amygdala using magnetic resonance imaging data obtained from 18,605 individuals aged 3-90 years. All subcortical structure volumes were at their maximum early in life; the volume of the basal ganglia showed a gradual monotonic decline thereafter while the volumes of the thalamus, amygdala and the hippocampus remained largely stable (with some degree of decline in thalamus) until the sixth decade of life followed by a steep decline thereafter. The lateral ventricles showed a trajectory of continuous enlargement throughout the lifespan. Significant age-related increase in inter-individual variability was found for the hippocampus and amygdala and the lateral ventricles. These results were robust to potential confounders and could be used to derive risk predictions for the early identification of diverse clinical phenotypes.

Introduction

Over the last 20 years, studies using structural magnetic resonance imaging (MRI) have confirmed that brain morphometric measures changes with age. In general, whole brain, global and regional gray matter volumes increase during development and decrease with aging (Brain Development Cooperative Group, 2012; Driscoll et al. 2009; Fotenos et al. 2005; Good et al. 2001; Pfefferbaum et al. 2013; Pomponio et al., 2019; Raz et al. 2005; Raznahan et al. 2014; Resnick et al. 2003; Walhovd et al. 2011). However, most published studies are constrained by small sample sizes, restricted age coverage and methodological variability. These limitations introduce inconsistencies and may obscure or distort the lifespan trajectories of brain structures. To address these limitations, we formed the Lifespan Working group of the Enhancing Neuroimaging Genetics through Meta-Analysis (ENIGMA) Consortium (Thompson et al. 2014, 2017) to perform large-scale analyses of brain morphometric data extracted from MRI images using standardized protocols and unified quality control procedures, harmonized and validated across all participating sites.

Here we focus on ventricular, striatal (caudate, putamen, nucleus accumbens), pallidal, thalamic, hippocampal and amygdala volumes. Subcortical structures are crucial for normal cognitive and emotional adaptation (Grossberg, 2009). The striatum and pallidum (together referred to as basal ganglia) are best known for their role in action selection and movement coordination (Calabresi et al. 2014) but they are also involved in other aspects of cognition particularly memory, inhibitory control, reward and salience processing (Chudasama and Robbins 2006; Richard et al. 2013; Scimeca and Badre 2012; Tremblay et al. 2015). The role of the hippocampus has been most clearly defined in connection to declarative memory (Eichenbaum, 2004; Shohamy and Turk-Browne 2013) while the amygdala has been historically linked to affect processing (Kober et al. 2008). The thalamus is centrally located in the brain and acts as a key hub for the integration of motor and sensory information with higher-order functions (Sherman 2005; Zhang et al. 2010). The role of subcortical structures extends beyond normal cognition because changes in the volume of these regions have been reliably identified in developmental (Ecker et al. 2015; Krain and Castellanos 2006), psychiatric (Kempton et al. 2011; Hibar et al. 2016; Schmaal et al. 2016; van Erp et al. 2016) and degenerative disorders (Risacher et al. 2009).

Using data from 18,605 individuals aged 3-90 years from the ENIGMA Lifespan working group we delineated the age-related trajectories of subcortical volumes from early to late life in order to (a) identify periods of volume change or stability, (b) provide normative, age-adjusted centile curves of subcortical volumes and (c) quantify inter-individual variability in

subcortical volumes which is considered a major source of inter-study differences in age-related trajectories derived from smaller samples (Dickie et al. 2013; Raz et al. 2010).

Materials and Methods

Study Samples

The study data comes from 88 samples and comprising 18,605 healthy participants, aged 3-90 years, with near equal representation of men and women (48% and 52%) (Table 1, Figure 1). At the time of scanning, participating individuals were screened to exclude the presence of mental disorders, cognitive impairment or significant medical morbidity. Details of the screening process and eligibility criteria for each research group are shown in Table S1).

Neuroimaging

Detailed information on scanner vendor, magnet strength and acquisition parameters for each sample are presented in Table S1. For each sample, the intracranial volume (ICV) and the volume of the basal ganglia (caudate, putamen, pallidum, nucleus accumbens), thalamus, hippocampus, amygdala and lateral ventricles were extracted using FreeSurfer (<http://surfer.nmr.mgh.harvard.edu>) from high-resolution T₁-weighted MRI brain scans (Fischl et al. 2002, 2012). Prior to data pooling, images were visually inspected at each site to exclude participants whose scans were improperly segmented. After merging the samples, outliers were identified and excluded using Mahalanobis distances. In each sample, the intracranial volume (Figure S1) was used to adjust the subcortical volumes via a formula based on the analysis of the covariance approach: 'adjusted volume = raw volume – b x (ICV – mean ICV)', where b is the slope of regression of a region of interest volume on ICV (Raz et al. 2005). The values of the subcortical volumes were then harmonized between sites using the ComBat method in R (Fortin, et al. 2017; 2018; Radua et al., 2019 this issue). Originally developed to adjust for batch effect in genetic studies, ComBat uses an empirical Bayes to adjust for inter-site variability in the data, while preserving variability related to the variables of interest.

Fractional polynomial regression analyses

The effect of age on each ICV- and site-adjusted subcortical volume was modelled using high order fractional polynomial regression (Royston and Altman 1994; Sauerbrei et al. 2006) in each hemisphere. Because the effect of site (and thus scanner and FreeSurfer version) were adjusted using ComBat, we only included sex as a covariate in the regression models. Fractional polynomial regression is currently considered the most advantageous

modelling strategy for continuous variables (Moore et al. 2011) as it allows testing for a wider range of trajectory shapes than conventional lower-order polynomials (e.g., linear or quadratic) and for multiple turning points (Royston and Altman 1994; Royston et al. 1999). For each subcortical structure, the best model was obtained by comparing competing models of up to three power combinations. The powers used to identify the best fitting model were -2, -1, -0.5, 0.5, 1, 2, 3 and the natural logarithm (ln) function. The optimal model describing the association between age and each of the volumes was selected as the lowest degree model based on the partial F-test (if linear) or the likelihood-ratio test. To avoid over-fitting at ages with more data points, we used the stricter 0.01 level of significance as the cut-off for each respective likelihood-ratio tests, rather than adding powers, until the 0.05 level was reached. For ease of interpretation we centred the volume of each structure so that the intercept of a fractional polynomial was represented as the effect at zero for sex. Fractional polynomial regression models were fitted using Stata/IC software v.13.1 (Stata Corp., College Station, TX). Standard errors were also adjusted for the effect of site in the FP regression.

We conducted two supplemental analyses: (a) we specified additional FP models separately for males and females and, (b) we calculated Pearson's correlation coefficient between subcortical volumes and age in the early (6-29 years), middle (30-59 years), and late-life (60-90 years) age-group. The results of these analyses have been included in the supplemental material.

Inter-individual variability

Inter-individual variability was assessed using two complimentary approaches. First, for each subcortical structure we compared the early (6-29 years), middle (30-59 years) and late-life (60-90 years) age-groups in terms of their mean inter-individual variability; these groups were defined following conventional notions regarding periods of development, midlife and aging. The variance of each structure in each age-group was calculated as

$$\ln \left(\frac{\sum \sqrt{e_i^2}}{n_t} \right)$$

where e represents the residual variance of each individual (i) around the non-linear best fitting regression line, and n the number of observations in each age-group (t). The residuals (e_i) were normally distributed suggesting good fit of the model without having over- or under-fitted the data. Upon calculating the square root of the squared residuals we used the natural logarithm to account for the positive skewness of the new distribution. Then the mean inter-

individual variability between early (6-29 years), middle (30-59 years) and late-life (60-90 years) age-groups was compared using between-groups omnibus tests for the residual variance around the identified best-fitting non-linear fractional polynomial model of each structure. The critical alpha value was set at 0.003 following Bonferroni correction for multiple comparisons.

The second approach entailed the quantification of the mean individual variability of each subcortical structure through a meta-analysis of the standard deviation of the adjusted volumes according to the method proposed by Senior et al. 2016.

Centile Curves

Reference curves for each structure by sex and hemisphere were produced from ICV- and site-adjusted volumes as normalized growth centiles using the parametric Lambda (λ), Mu (μ), Sigma (σ) (LMS) method (Cole and Green, 1992) implemented using the Generalised Additive Models for Location, Scale and Shape (GAMLSS) in R (<http://cran.r-project.org/web/packages/gamlss/index.html>) (Rigby and Stasinopoulos, 2005; 2007). LMS allows for the estimation of the distribution at each covariate value after a suitable transformation and is summarized using three smoothing parameters, the Box-Cox power λ , the mean μ and the coefficient of variation σ . GAMLSS uses an iterative maximum (penalized) likelihood estimation method to estimate λ , μ and σ as well as distribution dependent smoothing parameters and provides optimal values for effective degrees of freedom (edf) for every parameter (Indrayan, 2014). This procedure minimizes the Generalized Akaike Information Criterion (GAIC) goodness of fit index; smaller GAIC values indicate better fit of the model to the data. GAMLSS is a flexible way to derive normalized centile curves as it allows each curve to have its own number of edf while overcoming biased estimates resulting from skewed data.

Results

Fractional polynomial regression analyses

The volume of the caudate, putamen, globus pallidus and nucleus accumbens peaked early during the first decade of life and showed a linear decline immediately thereafter (Figure 2, Figures S2-S4). The age-related trajectories of the thalamic, hippocampal and amygdala volumes followed a flattened, inverted U-curve (Figure 3, Figures S5-S6). Specifically, the volumes of these structures were largest during the first 2-3 decades of life, remained largely stable until the 6th decade and declined gradually thereafter (Table S2). The volume of the lateral ventricles bilaterally increased steadily with age (Figure S7). The smallest proportion

of variance explained by age and its FP derivatives was noted in the right amygdala (7%) and the largest in the lateral ventricles bilaterally (38%) (Table S2).

Striatal volumes correlated negatively with age throughout the lifespan with the largest coefficients observed in the middle-life age-group ($r=-0.39$ to -0.20) and the lowest ($|r|<0.05$) in the late-life age-group, particularly in the caudate. The volumes of the thalamus, the hippocampus and the amygdala showed small positive correlations with age ($r\approx 0.16$) in the early-life age-group. In the middle-life age-group, the correlation between age and subcortical volumes became more negative ($r=-0.30$ to -0.27) for the thalamus but remained largely unchanged for the amygdala and the hippocampus. In the late-life age-group, the largest negative correlation coefficients between age and volume were observed for the hippocampus bilaterally ($r=-0.44$ to -0.39). The correlation between age and lateral ventricular volumes bilaterally increased throughout the lifespan from $r=0.19$ to 0.20 in early-life age-group to $r=0.40$ to 0.45 in the late-life age-group (Table S3). No effect of sex was noted for any pattern of correlation between subcortical volumes and age in any age-group.

Inter-individual variability: For each structure, the mean inter-individual variability in volume in each age-group is shown in Table S5. Inter-individual variance was significantly higher for the hippocampus, thalamus amygdala and lateral ventricles bilaterally in the late-life age-group compared to both the early- and middle-life group. These findings were recapitulated when data were analysed using a meta-analytic approach (Figure 4 and Figure S8).

Normative Centile Curves: Centile normative values for each subcortical structure stratified by sex and hemisphere are shown in Tables S6-S8.

Discussion

We analysed subcortical volumes from 18,605 healthy individuals from multiple cross-sectional cohorts to infer age-related trajectories between the ages of 3 to 90 years. Our lifespan perspective and our large sample size complement and enrich previous literature on age-related changes in subcortical volumes.

We found three distinct age-related trajectories. The volume of the lateral ventricles increased monotonically with age. Striatal and pallidal volumes peaked in childhood and declined thereafter. The volumes of the thalamus, hippocampus and amygdala peaked later and showed a prolonged period of stability lasting till the 6th decade of life, before they also started to decline. The trajectories defined here represent a close approximation to a

normative reference dataset and are in line with findings from Pomponio et al (2019) who also used harmonised multi-site MRI data from 10,323 individuals aged 3-96 years. Similar findings were reported by Douaud and colleagues (2014) who analysed volumetric data from 484 healthy participants aged 8 to 85 years; they also noted the similarity in the age-related trajectories of the thalamus, hippocampus and the amygdala. Our results also underscore the acceleration in age-related decline from the 6th decade of life onwards. This effect seemed relatively more pronounced for the hippocampus, compared to the other subcortical regions, as observed in other studies (Jernigan et al. 2001; Pomponio et al. 2019; Raz et al. 2010).

The trajectories of subcortical volumes are shaped by genetic and non-genetic exposures, biological or otherwise (Eyler et al. 2011; Somel et al. 2010; Wardlaw et al. 2011). Our findings of high age-related inter-individual variability in the volumes of the thalamus, hippocampus and amygdala suggest that these structures may be more susceptible to person-specific exposures, or late-acting genes, particularly from the 6th decade onwards.

In medicine, biological measures from each individual are typically categorised as normal or otherwise in reference to a population derived normative range. This approach is yet to be applied to neuroimaging data, despite the widespread use of structural MRI for clinical purposes and the obvious benefit of a reference range from the early identification of deviance (Dickie et al. 2013; Pomponio et al. 2019). Alzheimer's disease provides an informative example as the degree of baseline reduction in medial temporal regions, and particularly the hippocampus, is one of the most significant predictors of conversion from mild cognitive impairment to Alzheimer's disease (Risacher et al. 2009). The data presented here demonstrate the power of international collaborations within ENIGMA for analyzing very large-scale datasets that could eventually lead to normative range for brain volumes for well-defined reference populations. The unique strengths of this study are the availability of age-overlapping cross-sectional data from healthy individuals, lifespan coverage and the use of standardized protocols for volumetric data extraction across all samples. Study participants in each site were screened to ensure mental and physical wellbeing at the time of scanning using procedures considered as standard in designating study participants as healthy controls. Although health is not a permanent attribute, it is extremely unlikely given the size of our sample that our results could have been systematically biased by incipient disease.

A similar longitudinal design would be near infeasible in terms of recruitment and retention both of participants and investigators. Although multisite studies have to account for differences in scanner type and acquisition, lengthy longitudinal designs encounter similar issues due to inevitable changes in scanner type and strength and acquisition parameters

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over time. In this study, the use of age-overlapping samples from multiple different countries has the theoretical advantage of diminishing systematic biases reflecting cohort and period effects (Glenn, 2003; Keyes et al. 2010) that are likely to operate in single site studies.

In conclusion, we used existing data to derive age-related trajectories of regional subcortical volumes. The size and age-coverage of the analysis sample has the potential to disambiguate uncertainties regarding developmental and aging changes in subcortical volumes while the normative centile values could be further developed to derive clinically meaningful predictors of risk of adverse health outcomes.

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sample: HUBIN was supported by the Swedish Research Council (K2007-62X-15077-04-1, K2008-62P-20597-01-3, K2010-62X-15078-07-2, K2012-61X-15078-09-3), the regional agreement on medical training and clinical research between Stockholm County Council, and the Karolinska Institutet, and the Knut and Alice Wallenberg Foundation. P.M.T., N.J., M.J.W., S.E.M., O.A.A., D.A.R., L.S., D.J.V., T.G.M. v.E., D.G., and D.P.H. were supported in part by a Consortium grant (U54 EB020403 to P.M.T.) from the NIH Institutes contributing to the Big Data to Knowledge (BD2K) Initiative.

Conflict of interest

None of the authors reports any conflict of interest in connection to this manuscript.

Data Availability Statement

The ENIGMA Lifespan Working Group welcomes expression of interest from researchers in the field who wish to use the ENIGMA samples. Data sharing is possible subsequent to consent for the principal investigators of the contributing datasets. Requests should be directed to the corresponding authors.

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| Table 1. ENIGMA samples | | | | | | |
|--------------------------------|------------------------------|----------------------------|------------------------------|--------------------------------|---------------------|-----------------------|
| Sample | Age, Mean (Years) | Age, SD (Years) | Age Range (Years) | Sample Size (N) | Male (N) | Female (N) |
| ADHD NF | 13.4 | 0.9 | 12-15 | 12 | 6 | 6 |
| AMC | 23 | 3.4 | 17-32 | 94 | 60 | 34 |
| Barcelona 1.5T | 15 | 1.8 | 11-17 | 25 | 12 | 13 |
| Barcelona 3T | 14.5 | 2.2 | 11-17 | 37 | 18 | 19 |
| Betula | 62.5 | 12.8 | 26-81 | 263 | 123 | 140 |
| BIG | 28.4 | 14.3 | 13-82 | 1311 | 651 | 660 |
| BIG | 24.1 | 8.1 | 18-71 | 1275 | 537 | 738 |
| BIL&GIN | 26.7 | 7.7 | 18-57 | 451 | 219 | 232 |
| CAMH | 43.6 | 19.3 | 18-86 | 141 | 72 | 69 |
| Cardiff | 25.2 | 7.1 | 18-58 | 290 | 78 | 212 |
| CEG | 15.6 | 1.7 | 13-19 | 32 | 32 | 0 |
| CIAM | 26.1 | 4.8 | 19-40 | 27 | 13 | 14 |
| CLiNG | 25.2 | 5.3 | 18-58 | 316 | 130 | 186 |
| CODE | 39.7 | 13.3 | 20-64 | 72 | 31 | 41 |
| ENIGMA-HIV | 24.7 | 4.5 | 19-33 | 30 | 15 | 15 |
| ENIGMA-OCD (1) | 14.9 | 2 | 12-17 | 6 | 2 | 4 |
| ENIGMA-OCD (2) | 34.5 | 12.7 | 18-61 | 23 | 11 | 12 |
| ENIGMA-OCD (3) | 38.9 | 10.9 | 26-63 | 20 | 8 | 12 |
| ENIGMA-OCD (4) | 39.5 | 12.4 | 26-63 | 17 | 3 | 14 |
| ENIGMA-OCD (5) | 33.9 | 9.3 | 24-53 | 19 | 6 | 13 |
| ENIGMA-OCD (6) | 39.7 | 8.2 | 24-53 | 17 | 8 | 9 |
| ENIGMA-OCD (7) | 31.5 | 10.9 | 20-56 | 22 | 8 | 14 |
| FBIRN | 37.6 | 11.3 | 19-60 | 173 | 123 | 50 |
| FIDMAG | 37.5 | 10.1 | 19-64 | 123 | 54 | 69 |
| GSP | 27.2 | 16.5 | 18-90 | 1996 | 882 | 1114 |
| HMS | 39.6 | 12.2 | 19-64 | 55 | 21 | 34 |
| HUBIN | 42 | 8.8 | 19-56 | 102 | 69 | 33 |
| IMH | 31.9 | 10.1 | 20-59 | 78 | 49 | 29 |
| Indiana 1.5T | 62.5 | 11.7 | 37-84 | 49 | 9 | 40 |
| Indiana 3T | 27.7 | 20.1 | 6-87 | 199 | 95 | 104 |
| KaSP | 27.5 | 5.7 | 20-43 | 31 | 14 | 17 |

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| Table 1. ENIGMA samples | | | | | | |
|--------------------------------|------------------------------|----------------------------|------------------------------|--------------------------------|---------------------|-----------------------|
| Sample | Age, Mean (Years) | Age, SD (Years) | Age Range (Years) | Sample Size (N) | Male (N) | Female (N) |
| MAS | 78.4 | 4.7 | 70-90 | 528 | 240 | 288 |
| MCIC | 32.5 | 12.1 | 18-60 | 91 | 61 | 30 |
| Melbourne | 19.9 | 2.9 | 15-26 | 79 | 46 | 33 |
| METHCT | 26.5 | 6.8 | 18-53 | 59 | 45 | 14 |
| NESDA | 40.3 | 9.7 | 21-56 | 65 | 23 | 42 |
| NeuroIMAGE | 16.7 | 3.6 | 8-28 | 345 | 155 | 190 |
| Neuroventure | 13.7 | 0.6 | 12-15 | 130 | 55 | 75 |
| NU | 32.8 | 14.8 | 14-68 | 79 | 46 | 33 |
| NUIG | 36.1 | 11.6 | 18-58 | 92 | 53 | 39 |
| NYU | 31.1 | 8.7 | 19-52 | 49 | 29 | 20 |
| Olin | 36 | 13.1 | 21-87 | 590 | 236 | 354 |
| Oxford | 16.2 | 1.4 | 14-19 | 38 | 18 | 20 |
| QTIM | 22.6 | 3.4 | 16-30 | 306 | 92 | 214 |
| Sao Paolo (3) | 30.5 | 8.4 | 18-50 | 76 | 42 | 34 |
| SCORE | 25.5 | 4.3 | 19-39 | 44 | 17 | 27 |
| SHIP-2 | 54.4 | 12 | 32-81 | 190 | 99 | 91 |
| SHIP TREND | 50 | 13.5 | 22-79 | 425 | 229 | 196 |
| StagedDep | 46.9 | 8.4 | 27-58 | 19 | 4 | 15 |
| Stanford | 37 | 10.6 | 19-61 | 40 | 13 | 27 |
| STROKEMRI | 45.2 | 22.1 | 18-78 | 52 | 19 | 33 |
| Sydney | 39.1 | 22.1 | 12-84 | 157 | 65 | 92 |
| TOP | 35.4 | 9.9 | 18-73 | 303 | 159 | 144 |
| TS-Eurotrain | 10.9 | 1.1 | 9-13 | 45 | 29 | 16 |
| Tuebingen | 40.5 | 11.9 | 24-61 | 50 | 22 | 28 |
| UMCU | 39.7 | 14.5 | 19-65 | 66 | 26 | 40 |
| UNIBA | 27.4 | 9.1 | 18-63 | 128 | 64 | 64 |
| UPENN | 36.3 | 14 | 16-85 | 176 | 77 | 99 |
| Yale | 14.2 | 2.3 | 10-18 | 22 | 11 | 11 |
| Total | 32.9 | 18.3 | 6-90 | 11550 | 5334 | 6216 |

N=number; SD= standard deviation; Abbreviations of sample names: ADHD-NF=Attention Deficit Hyperactivity Disorder- Neurofeedback Study; AMC=Amsterdam Medisch Centrum; Barcelona=University of Barcelona; Betula=Swedish longitudinal study on aging, memory, and dementia; BIG=Brain Imaging Genetics; BIL&GIN=a multimodal multidimensional

database for investigating hemispheric specialization; CAMH=Centre for Addiction and Mental Health; Cardiff=Cardiff University; CEG=Cognitive-experimental and Genetic study of ADHD and Control Sibling Pairs; CIAM=Cortical Inhibition and Attentional Modulation study; CLING=Clinical Neuroscience Göttingen; CODE=formerly Cognitive Behavioral Analysis System of Psychotherapy (CBASP) study; ENIGMA-HIV=Enhancing Neuroimaging Genetics through Meta-Analysis-Human Immunodeficiency Virus Working Group; ENIGMA-OCD=Enhancing Neuroimaging Genetics through Meta-Analysis-Obsessive Compulsive Disorder Working Group; FBIRN=Function Biomedical Informatics Research Network; FIDMAG=Fundación para la Investigación y Docencia Maria Angustias Giménez; GSP=Brain Genomics Superstruct Project; HMS=Homburg Multidiagnosis Study; HUBIN=Human Brain Informatics; IMH=Institute of Mental Health, Singapore; Indiana=Indiana University School of Medicine; KaSP=The Karolinska Schizophrenia Project; MAS=Memory and Ageing Study; MCIC=MIND Clinical Imaging Consortium formed by the Mental Illness and Neuroscience Discovery (MIND) Institute now the Mind Research Network; Melbourne=University of Melbourne; Meth-CT=methamphetamine use, University of Cape Town; NESDA=The Netherlands Study of Depression and Anxiety; NeuroIMAGE=Dutch part of the International Multicenter ADHD Genetics (IMAGE) study; Neuroventure: the imaging part of the Co-Venture Trial funded by the Canadian Institutes of Health Research (CIHR); NU=Northwestern University; NUIG=National University of Ireland Galway; NYU=New York University; Olin=Olin Neuropsychiatric Research Center; Oxford=Oxford University; QTIM=Queensland Twin Imaging; Sao Paulo=University of Sao Paulo; SCORE: University of Basel Study; SHIP-2 and SHIP TREND=Study of Health in Pomerania; Staged-Dep= Stages of Depression Study; Stanford=Stanford University; StrokeMRI=Stroke Magnetic Resonance Imaging; Sydney=University of Sydney; TOP=Tematisk Område Psykoser (Thematically Organized Psychosis Research); TS-EUROTRAIN=European-Wide Investigation and Training Network on the Etiology and Pathophysiology of Gilles de la Tourette Syndrome; Tuebingen=University of Tuebingen; UMCU=Universitair Medisch Centrum Utrecht; UNIBA=University of Bari Aldo Moro; UPENN=University of Pennsylvania; Yale=Yale University

FIGURES

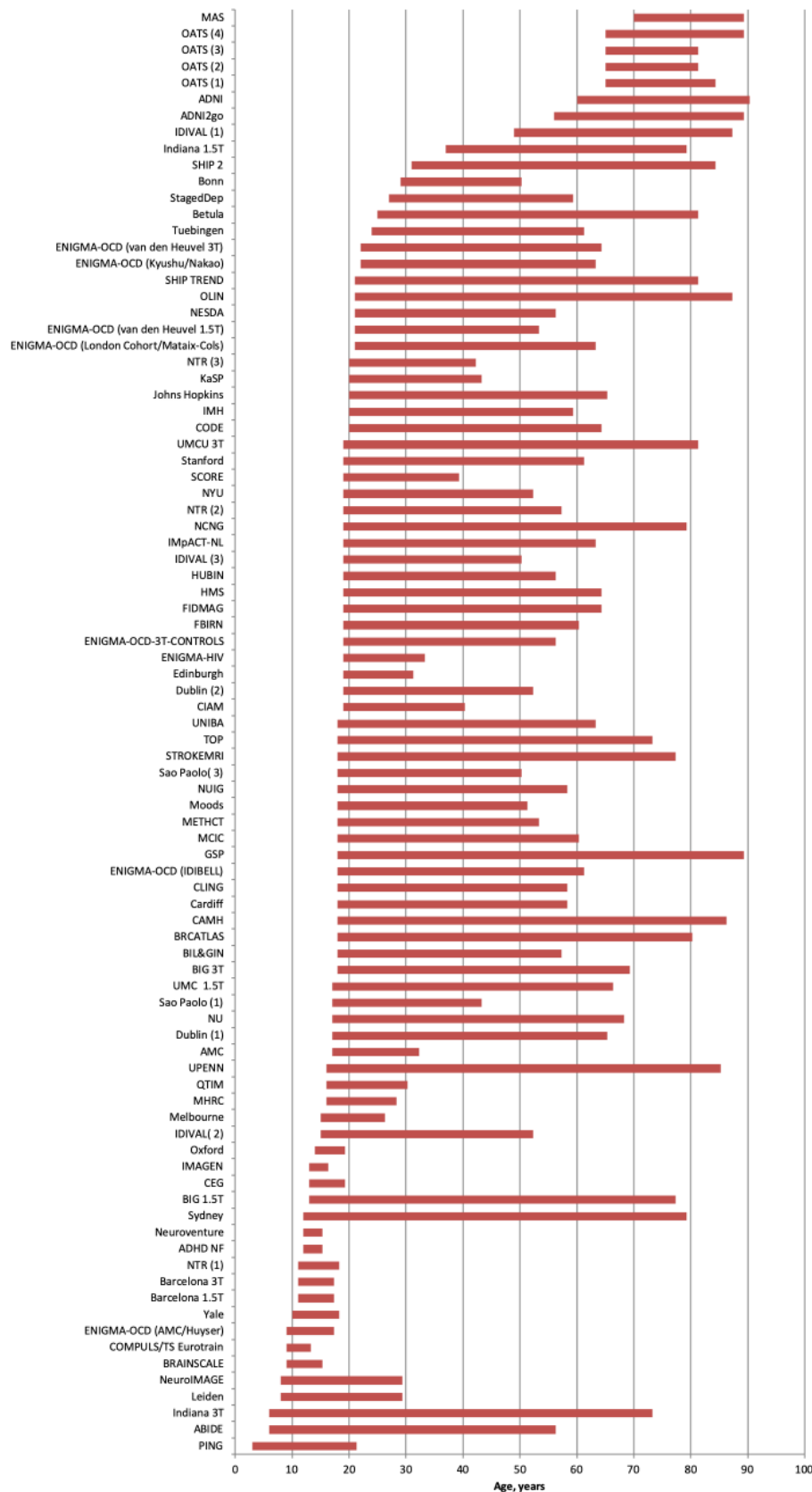


Figure 1. ENIGMA Lifespan Samples

Details of each sample are provided Table 1 and in the supplemental material. Abbreviations are provided in Table 1.

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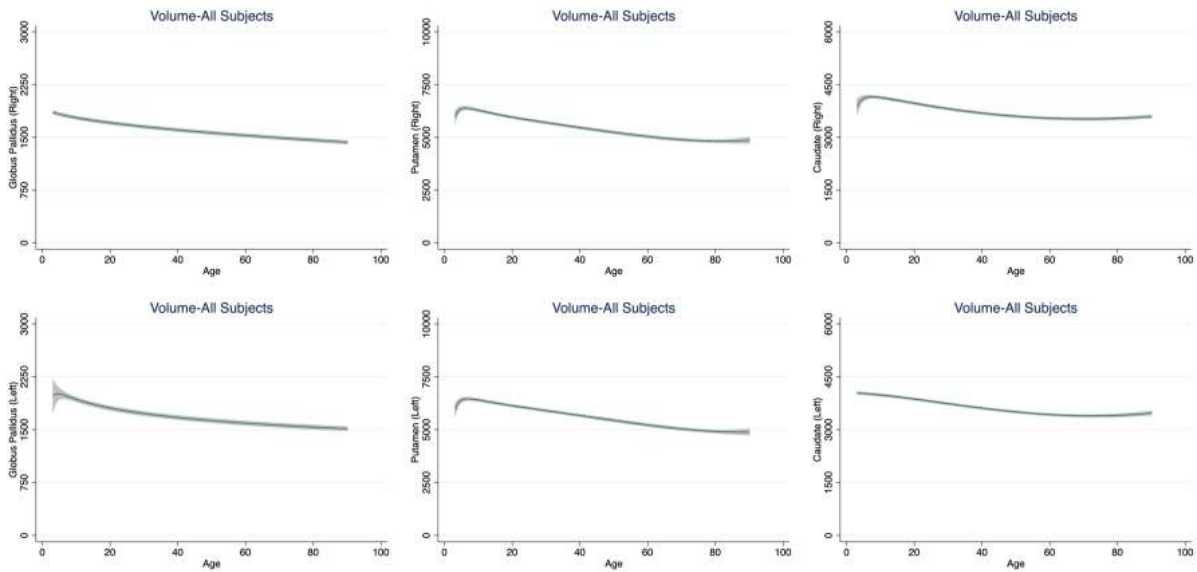


Figure 2. Fractional Polynomial Plots for the Volume of the Basal Ganglia

Fractional Polynomial plots of adjusted volumes (mm^3) against age (years) with a fitted regression line (solid line) and 95% confidence intervals (shaded area).

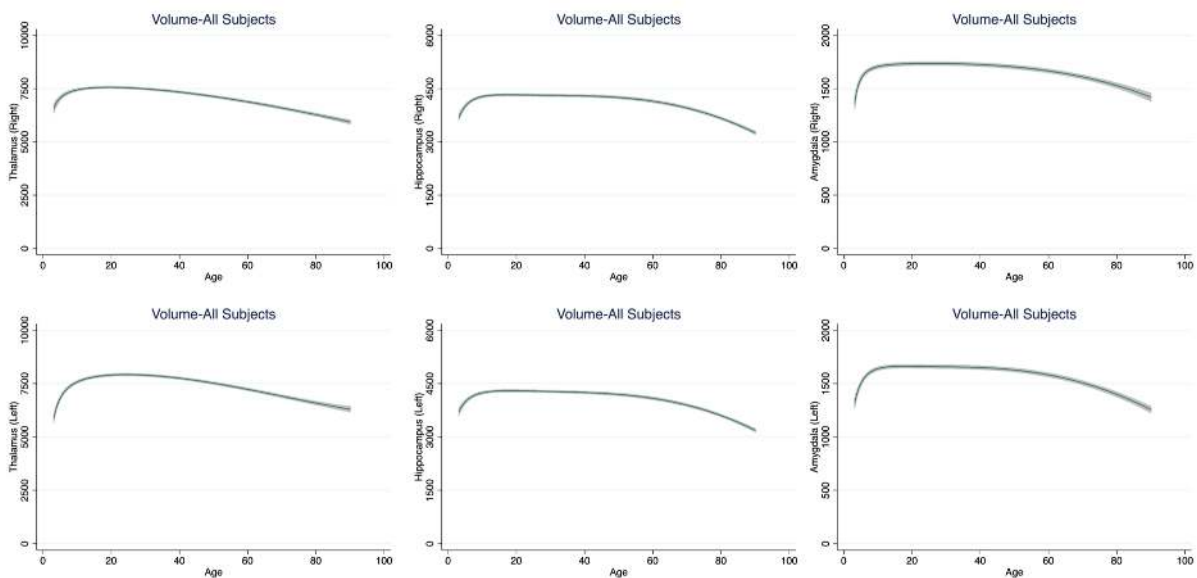


Figure 3. Fractional Polynomial Plots for the Volume of the Thalamus, Hippocampus and Amygdala

Fractional Polynomial plots of adjusted volumes (mm^3) against age (years) with a fitted regression line (solid line) and 95% confidence intervals (shaded area).

Short title: Subcortical trajectories throughout life

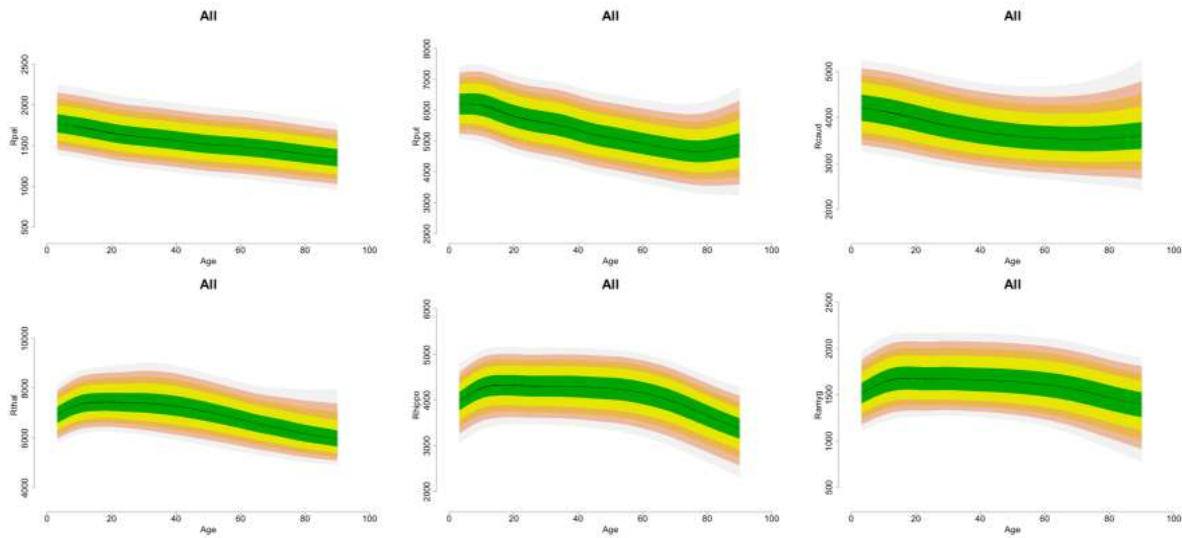


Figure 4. Mean Inter-individual Variability of Subcortical Volumes

Mean individual variability for each subcortical structure was estimated by means of a meta-analysis of the standard deviation of the adjusted volumes in each age-group.

References

- Allen JS, Bruss J, Brown CK, Damasio H (2005): Normal neuroanatomical variation due to age: the major lobes and a parcellation of the temporal region. *Neurobiology of Aging* 26:1245–1260.
- Brain Development Cooperative Group (2012): Total and regional brain volumes in a population-based normative sample from 4 to 18 years: The NIH MRI Study of Normal Brain Development. *Cereb Cortex* 22:1–12.
- Calabresi P, Picconi B, Tozzi A, Ghiglieri V, Di Filippo M (2014): Direct and indirect pathways of basal ganglia: a critical reappraisal. *Nat Neurosci* 17:1022-30.
- Chudasama Y, Robbins TW (2006): Functions of frontostriatal systems in cognition: comparative neuropsychopharmacological studies in rats, monkeys and humans. *Biol Psychol* 73:19-38.
- Cole TJ, Green PJ (1992): Smoothing Reference Centile Curves: The LMS Method and Penalized Likelihood. *Stat Med* 11:1305-1319.
- Dickie DA, Job DE, Gonzalez DR, Shenkin SD, Ahearn TS, Murray AD, Wardlaw JM (2013): Variance in brain volume with advancing age: implications for defining the limits of normality. *PLoS One* 8:e84093.
- Douaud G, Groves AR, Tamnes CK, Westlye LT, Duff EP, Engvig A, Walhovd KB, James A, Gass A, Monsch AU, Matthews PM, Fjell AM, Smith SM, Johansen-Berg H (2014): A common brain network links development, aging, and vulnerability to disease. *Proc Natl Acad Sci U S A* 111:17648-17653.
- Driscoll I, Davatzikos C, An Y, Wu X, Shen D, Kraut M, Resnick SM (2009): Longitudinal pattern of regional brain volume change differentiates normal aging from MCI. *Neurology* 72:1906–1913.
- Ecker C, Bookheimer SY, Murphy DG (2015): Neuroimaging in autism spectrum disorder: brain structure and function across the lifespan. *Lancet Neurol* 14:1121-1134.
- Eichenbaum H (2004): Hippocampus: cognitive processes and neural representations that underlie declarative memory. *Neuron* 44:109-120.
- Eyler LT, Prom-Wormley E, Fennema-Notestine C, Panizzon MS, Neale MC, Jernigan TL, Fischl B, Franz CE, Lyons MJ, Stevens A, Pacheco J, Perry ME, Schmitt JE, Spitzer NC, Seidman LJ, Thermenos HW, Tsuang MT, Dale AM, Kremen WS (2011): Genetic patterns of correlation among subcortical volumes in humans: results from a magnetic resonance imaging twin study. *Hum Brain Mapp* 32:641-653.
- Fischl B, Salat DH, Busa E, Albert M, Dieterich M, Haselgrove C, van der Kouwe A, Killiany R, Kennedy D, Klaveness S, Montillo A, Makris N, Rosen B, Dale AM (2002): Whole brain segmentation: automated labeling of neuroanatomical structures in the human brain. *Neuron* 33:341-355.

Fischl B (2012): FreeSurfer. *Neuroimage* 62:774-781.

Fortin JP, Parker D, Tunc B, Watanabe T, Elliott MA, Ruparel K, Roalf DR, Satterthwaite TD, Gur RC, Gur RE, Schultz RT, Verma R, Shinohara, RT (2017): Harmonization of multi-site diffusion tensor imaging data. *Neuroimage* 161:149-170.

Fortin JP, Cullen N, Sheline YI, Taylor WD, Aselcioglu I, Cook PA, Adams P, Cooper C, Fava M, McGrath PJ, McInnis M, Phillips ML, Trivedi MH, Weissman, MM, Shinohara RT (2018): Harmonization of cortical thickness measurements across scanners and sites. *Neuroimage* 167:104-120.

Fotenos AF, Snyder AZ, Girton LE, Morris JC, Buckner RL (2005): Normative estimates of cross-sectional and longitudinal brain volume decline in aging and AD. *Neurology* 64:1032–1039.

Glenn ND (2003): Distinguishing Age, Period, and Cohort Effects. In: JT Mortimer & MJ Shanahan, editors. *Handbook of the Life Course*. New York: Springer US. p 465-476.

Good CD, Johnsrude IS, Ashburner J, Henson RN, Friston KJ, Frackowiak RS (2001): A voxel-based morphometric study of ageing in 465 normal adult human brains. *NeuroImage* 14:21–36.

Grossberg S (2009): Cortical and subcortical predictive dynamics and learning during perception, cognition, emotion and action. *Philos Trans R Soc Lond B Biol Sci* 364:1223–1234.

Hibar DP, Westlye LT, van Erp TG, Rasmussen J, Leonardo CD, Faskowitz J, Haukvik UK, Hartberg CB, Doan NT, Agartz I, Dale AM, Gruber O, Krämer B, Trost S, Liberg B, Abé C, Ekman CJ, Ingvar M, Landén M, Fears SC, Freimer NB, Bearden CE; Costa Rica/Colombia Consortium for Genetic Investigation of Bipolar Endophenotypes, Sprooten E, Glahn DC, Pearlson GD, Emsell L, Kenney J, Scanlon C, McDonald C, Cannon DM, Almeida J, Versace A, Caseras X, Lawrence NS, Phillips ML, Dima D, Delvecchio G, Frangou S, Satterthwaite TD, Wolf D, Houenou J, Henry C, Malt UF, Bøen E, Elvsåshagen T, Young AH, Lloyd AJ, Goodwin GM, Mackay CE, Bourne C, Bilderbeck A, Abramovic L, Boks MP, van Haren NE, Ophoff RA, Kahn RS, Bauer M, Pfennig A, Alda M, Hajek T, Mwangi B, Soares JC, Nickson T, Dimitrova R, Sussmann JE, Hageaars S, Whalley HC, McIntosh AM, Thompson PM, Andreassen OA (2016): Subcortical volumetric abnormalities in bipolar disorder. *Mol Psychiatry* 21:1710-1716.

Indrayan A (2014): Demystifying LMS and BCPE methods of centile estimation for growth and other health parameters. *Indian pediatrics* 51: 37-43.

Jernigan TL, Archibald SL, Fenema-Notestine C, Gamst AC, Stout JC, Bonner J, Hesselink JR (2001): Effects of age on tissues and regions of the cerebrum and cerebellum. *Neurobiol Aging* 22:581-594.

- Kempton MJ, Salvador Z, Munafò MR, Geddes JR, Simmons A, Frangou S, Williams SC (2011): Structural neuroimaging studies in major depressive disorder. Meta-analysis and comparison with bipolar disorder. *Arch Gen Psychiatry* 68:675-690.
- Keyes KM, Utz RL, Robinson W, Li G (2010): What is a cohort effect? Comparison of three statistical methods for modelling cohort effects in obesity prevalence in the United States, 1971–2006. *Social Science & Medicine* 70:1100-1108.
- Kober H, Barrett LF, Joseph J, Bliss-Moreau E, Lindquist K, Wager TD (2008): Functional grouping and cortical-subcortical interactions in emotion: A meta-analysis of neuroimaging studies. *Neuroimage* 42:998-1031.
- Krain AL, Castellanos FX (2006): Brain development and ADHD. *Clin Psychol Rev* 26:433–444.
- Moore L, Hanley JA, Turgeon AF, Lavoie A (2011): A Comparison of Generalized Additive Models to Other Common Modeling Strategies for Continuous Covariates: Implications for Risk Adjustment. *J Biomet Biostat* 2:109.
- Pfefferbaum A, Rohlfing T, Rosenbloom MJ, Chu W, Colrain IM, Sullivan EV (2013): Variation in longitudinal trajectories of regional brain volumes of healthy men and women (ages 10 to 85 years) measured with atlas-based parcellation of MRI. *Neuroimage* 65:176-193.
- Pomponio, R., Erus, G., Habes, M., Doshi, J., Srinivasan, D., Mamourian, E., Bashyam, V., Nasrallah, I.M., Satterthwaite, T.D., Fan, Y., Launer L.J., Masters C.L., Maruff P., Zhuo, C., Völzke, H, Johnson, SC, Fripp, J, Koutsouleris, N., Wolf, D.H., Gur, R., Gur, R., Morris, J., Albert, M.S., Grabe, H.J., Resnick, S.M., Bryan, R.N., Wolk, D.A., Shinohara, R.T., Shou, H., Davatzikos, C. (2019) Harmonization of large MRI datasets for the analysis of brain imaging patterns throughout the lifespan. *Neuroimage*, doi: 10.1016/j.neuroimage.2019.116450
- Raz N, Lindenberger U, Rodrigue KM, Kennedy KM, Head D, Williamson A, Dahle C, Gerstorf D, Acker JD (2005): Regional brain changes in aging healthy adults: general trends, individual differences and modifiers. *Cereb Cortex* 15:1676-1689.
- Raz N, Ghisletta P, Rodrigue KM, Kennedy KM, Lindenberger U (2010): Trajectories of brain aging in middle-aged and older adults: regional and individual differences. *NeuroImage* 51:501– 511.
- Raznahan A, Shaw PW, Lerch JP, Clasen LS, Greenstein D, Berman R, Pipitone J, Chakravarty MM, Giedd JN (2014): Longitudinal four-dimensional mapping of subcortical anatomy in human development. *Proc Natl Acad Sci U S A* 111:1592-1597.
- Resnick SM, Pham DL, Kraut MA, Zonderman AB, Davatzikos C (2003): Longitudinal magnetic resonance imaging studies of older adults: a shrinking brain. *J. Neurosci* 23:3295–3301.

- Richard JM, Castro DC, Difeliceantonio AG, Robinson MJ, Berridge KC (2013): Mapping brain circuits of reward and motivation: in the footsteps of Ann Kelley. *Neurosci Biobehav Rev* 37:1919-1931.
- Rigby RA, Stasinopoulos DM (2005): Generalized additive models for location, scale and shape. *Appl. Statist* 54:507–554.
- Risacher SL, Saykin AJ, West JD, Shen L, Firpi HA, McDonald BC; Alzheimer's Disease Neuroimaging Initiative (2009): Baseline MRI predictors of conversion from MCI to probable AD in the ADNI cohort. *Curr Alzheimer Res* 6:347-361.
- Royston P, Altman DG (1994): Regression using fractional polynomials of continuous covariates: parsimonious parametric modelling. *Appl Stat* 43:429-467.
- Royston P, Ambler G, Sauerbrei W (1999): The use of fractional polynomials to model continuous risk variables in epidemiology. *Int J Epidemiol* 28:964-974.
- Sauerbrei W, Meier-Hirmer C, Benner A, Royston P (2006): Multivariable regression model building by using fractional polynomials: description of SAS, STATA and R programs. *Comput Stat Data An* 50: 3464-3485.
- Schmaal L, Veltman DJ, van Erp TG, Sämann PG, Frodl T, Jahanshad N, Loehrer E, Tiemeier H, Hofman A, Niessen WJ, Vernooij MW, Ikram MA, Wittfeld K, Grabe HJ, Block A, Hegenscheid K, Völzke H, Hoehn D, Czisch M, Lagopoulos J, Hatton SN, Hickie IB, Goya-Maldonado R, Krämer B, Gruber O, Couvy-Duchesne B, Rentería ME, Strike LT, Mills NT, de Zubicaray GI, McMahon KL, Medland SE, Martin NG, Gillespie NA, Wright MJ, Hall GB, MacQueen GM, Frey EM, Carballedo A, van Velzen LS, van Tol MJ, van der Wee NJ, Veer IM, Walter H, Schnell K, Schramm E, Normann C, Schoepf D, Konrad C, Zurowski B, Nickson T, McIntosh AM, Papmeyer M, Whalley HC, Sussmann JE, Godlewska BR, Cowen PJ, Fischer FH, Rose M, Penninx BW, Thompson PM, Hibar DP (2016): Subcortical brain alterations in major depressive disorder: findings from the ENIGMA Major Depressive Disorder working group. *Mol Psychiatry* 21:806-812.
- Scimeca JM, Badre D (2012): Striatal contributions to declarative memory retrieval. *Neuron* 75:380-392.
- Senior, A.M., Gosby, A.K., Lu, J., Simpson, S.J., Raubenheimer, D. (2016) Meta-analysis of variance: an illustration comparing the effects of two dietary interventions on variability in weight. *Evol Med Public Health*, 2016:244-55.
- Sherman SM (2005): Thalamic relays and cortical functioning. *Prog Brain Res* 149:107-126.
- Shohamy D, Turk-Browne NB (2013): Mechanisms for widespread hippocampal involvement in cognition. *J Exp Psychol Gen* 142:1159-1170.
- Somel M, Guo S, Fu N, Yan Z, Hu HY, Xu Y, Yuan Y, Ning Z, Hu Y, Menzel C, Hu H, Lachmann M, Zeng R, Chen W, Khaitovich P (2010): MicroRNA, mRNA, and protein

expression link development and aging in human and macaque brain. *Genome Res* 20:1207-1218.

Stasinopoulos DM, Rigby RA (2007): Generalized Additive Models for Location Scale and Shape (GAMLSS) in R. *J Stat Soft* 23:1-46.

Thompson PM, Andreassen OA, Arias-Vasquez A, Bearden CE, Boedhoe PS, Brouwer RM, Buckner RL, Buitelaar JK, Bulaeva KB, Cannon DM, Cohen RA, Conrod PJ, Dale AM, Deary IJ, Dennis EL, de Reus MA, Desrivieres S, Dima D, Donohoe G, Fisher SE, Fouche JP, Francks C, Frangou S, Franke B, Ganjgahi H, Garavan H, Glahn DC, Grabe HJ, Guadalupe T, Gutman BA, Hashimoto R, Hibar DP, Holland D, Hoogman M, Hulshoff Pol HE, Hosten N, Jahanshad N, Kelly S, Kochunov P, Kremen WS, Lee PH, Mackey S, Martin NG, Mazoyer B, McDonald C, Medland SE, Morey RA, Nichols TE, Paus T, Pausova Z, Schmaal L, Schumann G, Shen L, Sisodiya SM, Smit DJ, Smoller JW, Stein DJ, Stein JL, Toro R, Turner JA, van den Heuvel M, van den Heuvel OA, van Erp TG, van Rooij D, Veltman DJ, Walter H, Wang Y, Wardlaw JM, Whelan CD, Wright MJ, Ye J; ENIGMA Consortium (2017): ENIGMA and the individual: Predicting factors that affect the brain in 35 countries worldwide. *NeuroImage* 145:389-408.

Thompson PM, Stein JL, Medland SE, Hibar DP, Vasquez AA, Renteria ME, Toro R, Jahanshad N, Schumann G, Franke B, Wright MJ, Martin NG, Agartz I, Alda M, Alhusaini S, Almasy L, Almeida J, Alpert K, Andreasen NC, Andreassen OA, Apostolova LG, Appel K, Armstrong NJ, Aribisala B, Bastin ME, Bauer M, Bearden CE, Bergmann O, Binder EB, Blangero J, Bockholt HJ, Bøen E, Bois C, Boomsma DI, Booth T, Bowman IJ, Bralten J, Brouwer RM, Brunner HG, Brohawn DG, Buckner RL, Buitelaar J, Bulayeva K, Bustillo JR, Calhoun VD, Cannon DM, Cantor RM, Carless MA, Caseras X, Cavalleri GL, Chakravarty MM, Chang KD, Ching CR, Christoforou A, Cichon S, Clark VP, Conrod P, Coppola G, Crespo-Facorro B, Curran JE, Czisch M, Deary IJ, de Geus EJ, den Braber A, Delvecchio G, Depondt C, de Haan L, de Zubicaray GI, Dima D, Dimitrova R, Djurovic S, Dong H, Donohoe G, Duggirala R, Dyer TD, Ehrlich S, Ekman CJ, Elvsåshagen T, Emsell L, Erk S, Espeseth T, Fagerness J, Fears S, Fedko I, Fernández G, Fisher SE, Foroud T, Fox PT, Francks C, Frangou S, Frey EM, Frodl T, Frouin V, Garavan H, Giddaluru S, Glahn DC, Godlewska B, Goldstein RZ, Gollub RL, Grabe HJ, Grimm O, Gruber O, Guadalupe T, Gur RE, Gur RC, Göring HH, Hagenaars S, Hajek T, Hall GB, Hall J, Hardy J, Hartman CA, Hass J, Hatton SN, Haukvik UK, Hegenscheid K, Heinz A, Hickie IB, Ho BC, Hoehn D, Hoekstra PJ, Hollinshead M, Holmes AJ, Homuth G, Hoogman M, Hong LE, Hosten N, Hottenga JJ, Hulshoff Pol HE, Hwang KS, Jack CR Jr, Jenkinson M, Johnston C, Jönsson EG, Kahn RS, Kasperaviciute D, Kelly S, Kim S, Kochunov P, Koenders L, Krämer B, Kwok JB, Lagopoulos J, Laje G, Landen M, Landman BA, Lauriello J, Lawrie SM, Lee PH, Le Hellard S, Lemaître H, Leonardo CD, Li CS, Liberg B, Liewald DC, Liu X, Lopez LM, Loth E, Lourdusamy A,

Luciano M, Macciardi F, Machielsen MW, Macqueen GM, Malt UF, Mandl R, Manoach DS, Martinot JL, Matarin M, Mather KA, Mattheisen M, Mattingsdal M, Meyer-Lindenberg A, McDonald C, McIntosh AM, McMahon FJ, McMahon KL, Meisenzahl E, Melle I, Milaneschi Y, Mohnke S, Montgomery GW, Morris DW, Moses EK, Mueller BA, Muñoz Maniega S, Mühleisen TW, Müller-Myhsok B, Mwangi B, Nauck M, Nho K, Nichols TE, Nilsson LG, Nugent AC, Nyberg L, Olvera RL, Oosterlaan J, Ophoff RA, Pandolfo M, Papalampropoulou-Tsiridou M, Pappmeyer M, Paus T, Pausova Z, Pearlson GD, Penninx BW, Peterson CP, Pfenning A, Phillips M, Pike GB, Poline JB, Potkin SG, Pütz B, Ramasamy A, Rasmussen J, Rietschel M, Rijpkema M, Risacher SL, Roffman JL, Roiz-Santiañez R, Romanczuk-Seiferth N, Rose EJ, Royle NA, Rujescu D, Ryten M, Sachdev PS, Salami A, Satterthwaite TD, Savitz J, Saykin AJ, Scanlon C, Schmaal L, Schnack HG, Schork AJ, Schulz SC, Schür R, Seidman L, Shen L, Shoemaker JM, Simmons A, Sisodiya SM, Smith C, Smoller JW, Soares JC, Sponheim SR, Sprooten E, Starr JM, Steen VM, Strakowski S, Strike L, Sussmann J, Sämann PG, Teumer A, Toga AW, Tordesillas-Gutierrez D, Trabzuni D, Trost S, Turner J, Van den Heuvel M, van der Wee NJ, van Eijk K, van Erp TG, van Haren NE, van 't Ent D, van Tol MJ, Valdés Hernández MC, Veltman DJ, Versace A, Völzke H, Walker R, Walter H, Wang L, Wardlaw JM, Weale ME, Weiner MW, Wen W, Westlye LT, Whalley HC, Whelan CD, White T, Winkler AM, Wittfeld K, Woldehawariat G, Wolf C, Zilles D, Zwiers MP, Thalamuthu A, Schofield PR, Freimer NB, Lawrence NS, Drevets W; Alzheimer's Disease Neuroimaging Initiative, EPIGEN Consortium, IMAGEN Consortium, Saguenay Youth Study (SYS) Group (2014): The ENIGMA Consortium: large-scale collaborative analyses of neuroimaging and genetic data. *Brain Imaging Behav* 8:153-182.

Tremblay L, Worbe Y, Thobois S, Sgambato-Faure V, Féger J (2015): Selective dysfunction of basal ganglia subterritories: From movement to behavioral disorders. *Mov Disord* 30:1155-1170.

van Erp TG, Hibar DP, Rasmussen JM, Glahn DC, Pearlson GD, Andreassen OA, Agartz I, Westlye LT, Haukvik UK, Dale AM, Melle I, Hartberg CB, Gruber O, Kraemer B, Zilles D, Donohoe G, Kelly S, McDonald C, Morris DW, Cannon DM, Corvin A, Machielsen MW, Koenders L, de Haan L, Veltman DJ, Satterthwaite TD, Wolf DH, Gur RC, Gur RE, Potkin SG, Mathalon DH, Mueller BA, Preda A, Macciardi F, Ehrlich S, Walton E, Hass J, Calhoun VD, Bockholt HJ, Sponheim SR, Shoemaker JM, van Haren NE, Hulshoff Pol HE, Ophoff RA, Kahn RS, Roiz-Santiañez R, Crespo-Facorro B, Wang L, Alpert KI, Jönsson EG, Dimitrova R, Bois C, Whalley HC, McIntosh AM, Lawrie SM, Hashimoto R, Thompson PM, Turner JA (2016): Subcortical brain volume abnormalities in 2028 individuals with schizophrenia and 2540 healthy controls via the ENIGMA consortium. *Mol Psychiatry* 21:547-553

Short title: Subcortical trajectories throughout life

Walhovd KB, Westlye LT, Amlien I, Espeseth T, Reinvang I, Raz N, Agartz I, Salat DH, Greve DN, Fischl B, Dale AM, Fjell AM (2011): Consistent neuroanatomical age-related volume differences across multiple samples. *Neurobiol Aging* 32:916-932.

Wardlaw JM, Bastin ME, Valdés Hernández MC, Maniega SM, Royle NA, Morris Z, Clayden JD, Sandeman EM, Eadie E, Murray C, Starr JM, Deary IJ (2011): Brain aging, cognition in youth and old age and vascular disease in the Lothian Birth Cohort 1936: rationale, design and methodology of the imaging protocol. *Int J Stroke* 6:547-559.

Zhang D, Snyder AZ, Shimony JS, Fox MD, Raichle ME (2010): Noninvasive functional and structural connectivity mapping of the human thalamocortical system. *Cereb Cortex* 20:1187-1194.