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Beyond brain age: Empirically-derived proxy measures of mental health — Source link 🖸

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Beyond brain age: Empirically-derived proxy measures of mental health

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Abstract

Biological aging is revealed by physical measures, e.g., DNA probes or brain scans. Individual differences in personal functioning are instead explained by psychological constructs. Constructs such as intelligence or neuroticism are typically assessed by specialized workforce through tailored questionnaires and tests. Similar to how brain age captures biological aging, intelligence and neuroticism may provide empirical proxies for mental health. Could the combination of brain imaging and sociodemographic information yield measures for these constructs that do not rely on human judgment? Here, we built proxy measures by applying machine learning on multimodal MR images and rich sociodemographic information from the largest brain-imaging cohort to date: the UK Biobank. Objective comparisons revealed that all proxies captured the target constructs and related to health-contributing habits beyond the measures they were derived from. Our results demonstrate that proxies targeting classical psychological constructs reveal facets of mental health complementary to information conveyed by brain age.

Introduction

Individual assessments in psychology and psychiatry rely on observing behavior. Using 2 biological insight to diagnose and treat mental disorders remains a hard problem despite 3 substantial research efforts (Kapur et al., 2012). The field of psychiatry has struggled with 4 purely descriptive and unstable diagnostic systems (Insel et al., 2010), small sample sizes 5 (Szucs and loannidis, 2017), and reliance on dichotomized groups, *i.e.*, patients vs controls 6 (Hozer and Houenou, 2016). Compared to somatic medicine, mental-health research faces 7 the additional roadblock that mental pathologies cannot be measured the same way diabetes can be assessed through plasma levels of insulin or glucose. Psychological constructs, e.g., 9 depressiveness or anxiety can only be probed indirectly through expert-built procedures 10

such as specially-crafted questionnaires and structured interviews. Measuring reliably a 11 given construct is difficult and questionnaires often remain the best option (Enkavi et al., 12 2019). While the field of psychometrics has thoroughly studied the validity of psychological 13 constructs and their measures (Borsboom et al., 2004; Cronbach and Meehl, 1955; Eisenberg 14 et al., 2019), the advent of new biophysical measurements on the brain brings new promises 15 (Engemann et al., 2020; Kievit et al., 2018b; Nave et al., 2018). In particular, the growth of 16 biobanks as well as the advances in statistical-learning techniques opens the door to large-17 scale validation of psychological constructs and measures for neuropsychiatric research 18 (Collins, 2012). 19

In clinical neuroscience, machine learning is increasingly popular, driven by the hope 20 to develop more generalizable models (Woo et al., 2017). Yet, to be reliable, machine 21 learning needs large labeled datasets (Varoquaux, 2018). Its application to learn imaging 22 biomarkers of neuropsychiatric disorders is limited by the availability of large cohorts with 23 high-quality neuropsychiatric assessements (Bzdok and Meyer-Lindenberg, 2018). However, 24 data on populations without diagnosed neuropsychiatric conditions is easier to collect. Such 25 data has driven successes in developing brain-derived aging measures that capturing proxy 26 information on mental health (Cole et al., 2015, 2018; Dosenbach et al., 2010; Engemann 27 et al., 2020; He et al., 2020; Koutsouleris et al., 2014; Liem et al., 2017; Smith et al., 2020). 28 Extrapolating from these successes, we propose to learn more of such proxy measures of 29 health-related individual traits in large datasets. These could then enhance an analysis in 30 a small dataset via links between the proxy measures and the actual clinical endpoint of 31 interest, e.g., diagnosis or drug response. Emerging results validate the usefulness of age 32 as one such proxy measure, leading to the so called brain age delta: the difference between 33 predicted and actual age (Cole et al., 2015; Dosenbach et al., 2010; Smith et al., 2019a). 34 The delta has been shown to reflect physical and cognitive impairment in adults and gives an 35 index of neurodegenerative processes (Gonneaud et al., 2020; Liem et al., 2017). Can this 36 strategy of biomarker-like proxy measures be extended beyond the construct of aging? Can 37 measures derived from other targets than age serve as proxies for latent constructs? 38

Bevond aging, one high-stake target is intelligence, which is measured through socially 39 administered tests and is one of the most extensively studied constructs in psychology. 40 Fluid intelligence refers to the putatively culture-free, heritable and physiological component 41 of intelligence (Cattell, 1963; Cattell and Scheier, 1961). Fluid intelligence is a latent 42 construct designed to capture individual differences in cognitive capacity. It has been 43 robustly associated with neuronal maturation and is typically reflected in cognitive-processing 44 speed and working-memory capacity (Shelton et al., 2010). Compared to brain age, fluid 45 intelligence may yield a proxy measure more specifically indexing cognitive function. It 46 has been associated with psychiatric disorders such as psychosis, bipolar disorder and 47 substance abuse (Keyes et al., 2017; Khandaker et al., 2018). 48

Neuroticism is a second promising target. As a key representative of the extensively stud-49 ied Big Five personality inventory, neuroticism has a long-standing tradition in the psychology 50 of individual differences (Costa and McCrae, 1992; Eysenck et al., 1985). Neuroticism is 51 typically measured using self-assessment questionnaires and conceptualized as capturing 52 dispositional negative emotionality including anxiety and depressiveness (Shackman et al., 53 2016). It has been inter-culturally validated (Cattell and Scheier, 1961; Lynn and Martin, 54 1997) and population-genetics studies have repeatedly linked variance in neuroticism to 55 shared genes (Birley et al., 2006; Hettema et al., 2006; Pedersen et al., 1988). Neuroticism 56 was shown useful in psychometric screening and supports predicting real-world behavior 57 (Lahey, 2009; Tyrer et al., 2015). However, despite strong heritability at the population level 58 (Power and Pluess, 2015; Vukasović and Bratko, 2015), the link with brain function at the 59 level of large-scale network dynamics or the level of molecular mechanisms is being actively 60

researched (Shackman et al., 2016; Yarkoni, 2015).

The advent of large MRI datasets has revealed the complexity of predicting personality 62 traits from brain signals. Current attempts to predict fluid intelligence or neuroticism from 63 thousands of MRI scans argue in favor of overwhelming heterogeneity and rather subtle 64 effects that do not generalize strongly to unseen data (Dubois et al., 2018a,b). This stands in 65 contrast to the remarkable performance obtained when predicting intelligence or neuroticism 66 from other psychometric measures or semantic data qualitatively similar to psychometric 67 questionnaires, e.g., Twitter and Facebook posts (Quercia et al., 2011; Youyou et al., 2015). 68 As MRI acquisitions can be expensive and difficult in clinical settings or populations, the 69 promises of social-media data is appealing. However, in clinical practice or research, such 70 data can lead to measurement and selection biases difficult to control. On the other hand, 71 background sociodemographic characteristics of individuals can be easily accessible and 72 may help inform in similar ways on the heterogeneity of psychological traits, for instance 73 capturing that fluid intelligence decreases with age (Horn et al., 1981). An important question 74 is then whether this data can reveal non-redundant information on the constructs of interest. 75

Another challenge of quantifying psychological traits is the diversity of measurement 76 scales, often categorical or on arbitrary non-physical units, e.g. education degree or monthly 77 income. In fact, society treats individual differences as categorical or continuous, depending 78 on the practical context. Personality has been proposed to span a continuum (Eysenck, 79 1958). Nevertheless, psychiatrists treat certain people as patients and not others (Perlis, 80 2011). The utility of any mental-health measure therefore depends on its practical context: 81 When learning boundaries between gualitatively distinct groups, a measure that performs 82 globally poorly as a continuous scale can nevertheless be sufficient to distinguish subgroups. 83 In fact, a measure may be solely informative around the boundary region between certain 84 classes, e.g., pilots who should fly and who should not. Importantly, the utility of any measure 85 ultimately depends on its signal-to-noise ratio, which may be driven by measurement noise, 86 heterogeneity, as well as the interesting variability of the particular construct measured, e.g., 87 the type of test to assess intelligence. 88

Confronting the promises of population brain imaging with the challenges of measuring 89 psychological traits raises the following questions. 1) How well can various health-related 90 latent constructs be approximated from general-purpose inputs not designed to measure 91 specific latent constructs? 2) What is the relative merit of brain imaging and sociodemo-92 graphics for probing various latent constructs? 3) Can the success of brain age be extended 93 to other proxy measures that capture complementary facets of health-contributing behavior? 94 In this study, we tackled these questions by using machine learning to build proxy mea-95 sures, crafted to approximate well-characterized target measures from brain-imaging and 96 sociodemographic data. As target measures, we studied age, fluid intelligence, and neuroti-97 cism. Figure 1 summarizes our approach. We first assessed how well the proxy measures 98 approximated the target measures, isolating the contributions of the different data types. 99 Second, to assess the intrinsic value of the proxy measures, we studied their associations 100 with health-related habits (alcohol consumption, cumulative tobacco consumption, sleep 101 duration, physical activity). Results suggest that, as with brain age, proxy measures can bring 102 value for the study of mental health that goes beyond approximating an available measure. 103

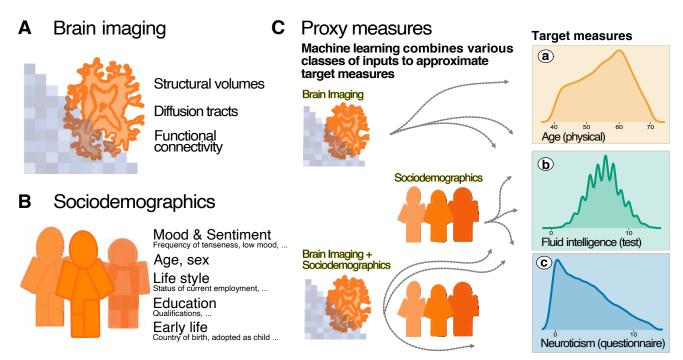


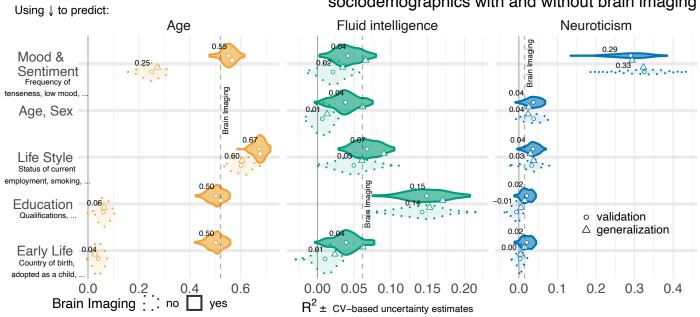
Figure 1. Methods overview: building and evaluating proxy measures We combined multiple brain-imaging modalities (**A**) with sociodemographic data (**B**) to approximate health-related biomedical and psychological constructs (**C**), *i.e.*, brain age (accessed through prediction of chronological age), cognitive capacity (accessed through a fluid-intelligence test) and the tendency to report negative emotions (accessed through a neuroticism questionnaire). We included the imaging data from the 10 000-subjects release of the UK biobank. Among imaging data (**A**) we considered features related to cortical and subcortical volumes, functional connectivity from rfMRI based on ICA networks, and white-matter molecular tracts from diffusive directions (see Table 3 for an overview about the multiple brain-imaging modalities). We then grouped the sociodemographic data (**B**) into five different blocks of variables related to self-reported mood & sentiment, primary demographics, lifestyle, education, and early-life events (Table 4 lists the number of variables in each block). Subsequently, we systematically compared the approximations of all three targets based on either brain images and sociodemographics in isolation or combined (**C**) to evaluate the relative contribution of these distinct inputs. Models were developed on 50% of the data (randomly drawn) based on random forest regression guided by Monte Carlo cross-validation with 100 splits (see section Model Development and Generalization Testing). We assessed generalization using the other 50% of the data as fully independent out-of-sample evaluations (see

Results

Traditional measures of mental health can be empirically approximated

We first performed model comparisons to evaluate the relative performance of proxy mea-106 sures built from brain signals and distinct groups of sociodemographic variables. Figure 2 107 summarizes these model comparisons for approximating three targets: age, fluid intelligence 108 and neuroticism. For the sociodemographic variables (Figure 2, dotted outlines), the analysis 109 revealed that, for each target, there was one principal block of variables explaining most 110 of the prediction performance. Combining all sociodemographic variables did not lead to 111 obvious enhancements (Figure 2 – Figure supplement 2). For age prediction, variables 112 related to current life-style showed by far the highest performance. For fluid intelligence, 113 education performed by far best. Finally, for neuroticism, mood & sentiment clearly showed 114 the strongest performance. 115

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Approximation quality of proxy measures derived from sociodemographics with and without brain imaging

Figure 2. Approximation performance of proxy measures derived from sociodemographic data and MRI. To approximate age, fluid intelligence and neuroticism, we applied random-forest regression on sociodemographic data and brain images as inputs. The data was split into validation data for model construction (see section Model Development and Generalization Testing) and generalization data for statistical inference on out-of-sample predictions with independent data (see section Statistical Analysis). For each block of sociodemographic predictors models were fitted with and without additional predictors derived from brain images. We report the R^2 metric to facilitate comparisons across prediction targets. The cross-validation (CV) distribution (100 Monte Carlo splits) on the validation dataset is depicted by violins. Drawing style indicates whether brain imaging (solid outlines of violins) was included in addition or not (dotted outlines of violins). Dots depict the average performance on the validation data across CV-splits. Pyramids depict the performance of the average prediction (CV-bagging) on held-out generalization datasets. For convenience, the mean performance on the validation set is annotated for each plot. Vertical dotted lines indicate the average performance of the full MRI model. The validation and held-out datasets gave similar picture of approximation performance with no evidence for cross-validation bias Varoquaux et al. (2017a). One can readily see that approximation from sociodemographics (dotted violins) was often markedly better than purely brain-based models (dotted vertical lines) for all three targets. The most important blocks of sociodemographic predictors (annotated with red cross) were lifestyle for age, education for fluid intelligence, and mood & sentiment for neuroticism. The effect of combining sociodemographics with brain-data depended on the target measure. For age, overall performance improved beyond the purely sociodemographics-based or imaging-based analyses. The picture was less consistent for fluid intelligence and neuroticism showing weaker additive effects, if any. For the averaged out-of-sample predictions, the probability of the observed performance under the null-distribution and the uncertainty of effect sizes were formally probed using permutation tests and bootstrap-based confidence intervals (Table 1). Corresponding statistics for the baseline performance of models solely based on brain imaging (vertical dotted lines) are presented in Table S1. For additional findings please consider the supplement of Figure 2: Figure 2 – Figure supplement 1. Prediction of individual differences in proxy measures from MRI.

Figure 2 - Figure supplement 2. Approximation performance using all sociodemographic data.

Target	sociodemographics	R^2_{diff}	p-value	Cl _{low}	Cl _{high}
Age	Early Life	0.494	0.0001	0.473	0.515
Age	Education	0.458	0.0001	0.437	0.479
Age	Life style	0.071	0.0001	0.058	0.085
Age	Mood & sentiment	0.294	0.0001	0.272	0.315
Fluid intelligence	Age, Sex	0.048	0.0001	0.040	0.057
Fluid intelligence	Early Life	0.039	0.0001	0.027	0.050
Fluid intelligence	Education	0.018	0.0001	0.010	0.025
Fluid intelligence	Life style	0.030	0.0001	0.020	0.040
Fluid intelligence	Mood & sentiment	0.031	0.0001	0.019	0.043
Neuroticism	Age, Sex	0.001	0.6789	-0.006	0.008
Neuroticism	Early Life	0.010	0.0697	-0.001	0.021
Neuroticism	Education	0.009	0.0817	-0.001	0.020
Neuroticism	Life style	-0.008	0.1750	-0.020	0.004
Neuroticism	Mood & sentiment	-0.030	0.0001	-0.041	-0.018

Table 1. Paired difference between purely sociodemographic and models including brain imag	ing on
generalization data.	

When combining MRI and sociodemographics (Figure 2, solid outlines), age prediction 116 was enhanced in a systematic and visible way on all four blocks of variables (Table 1), 117 suggesting that the observed differences should reproduce on future data and are unlikely to 118 be due to chance. The benefit of including brain-imaging features, however, was less marked 119 for prediction of fluid intelligence and neuroticism. With fluid intelligence, brain-imaging data 120 improved the performance statistically significantly for all models, yet, with small effect sizes 121 at the scale of a few percent or even lower (Table 1). Further, for neuroticism, no systematic 122 advantage of including brain images alongside sociodemographics emerged. Instead, includ-123 ing brain images seemed to reduce generalization performance when predicting from mood & 124 sentiment variables (Table 1, bottom row). Nevertheless, using only brain data was sufficient 125 for statistically significant approximation of the target measures not only for age but also 126 fluid intelligence and neuroticism (Table S1), suggesting that lifestyle and mood & sentiment 127 explains at least some of the neurobiological variance. For neuroticism, variables on current 128 mood & sentiment were strongly informative for prediction, reflecting that mood & sentiment 129 is strongly related to neuroticism. Overall, predicting fluid intelligence or neuroticism was 130 clearly more successful when sociodemographic was included (Table 1). For subsequent 131 analyses we included all sociodemographic variables (Figure 2 - Figure supplement 2). 132

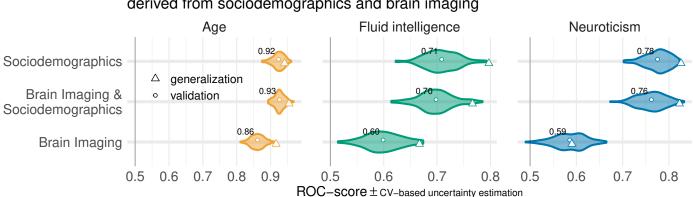
One important challenge with evaluating approximations of psychological measures is 133 that such measures often come without physical scales and units (Stevens et al., 1946). 134 In practice, clinicians and educators use them with specific thresholds for decision making. 135 How useful proxy measures built with predictive models are to separate out discrete extreme 136 groups? To address this question, we performed binary classification of extreme groups 137 obtained from discretizing the targets using the 33rd and 66th percentiles. Moreover, we 138 focused on the AUC as a performance metric which is only sensitive to ranking while 139 ignoring the scale of the error. The results are comparable to the previous regression 140 analysis. Classification performance for extreme groups visibly exceeded the chance level 141 of an AUC of 0.5 for all models (Figure 3). Across proxy measures, models including 142 sociodemographics performed best but the difference between purely sociodemographic and 143 brain-based models was comparably weak, at the order of 0.01-0.02 AUC points (Table 2). 144 Using only brain data resulted in proxy measures that perform less well, yet, still better than 145 chance as revealed by permutation testing (Table S2). It is noteworthy that for both types 146 of models the performance of discrimination reached levels above 0.8, which is considered 147 clinically useful for biomarkers (Perlis, 2011). Overall, the results suggest that moving 148 from the more difficult full-scale regression problem to extreme-group classification problem 149 with purely ranking-based loss functions, the relative differences between brain-based and 150 sociodemographics-based prediction gradually faded away. 151

Table 2. Difference statistics for classification on the held-out set for sociodemographic vs combined approximation.

Target	AUC _{diff observed}	p-value	Cl _{low}	Cl _{high}
Age	0.013	0.0008	0.006	0.021
Fluid intelligence	-0.031	0.0001	-0.044	-0.017
Neuroticism	-0.003	0.4818	-0.013	0.006

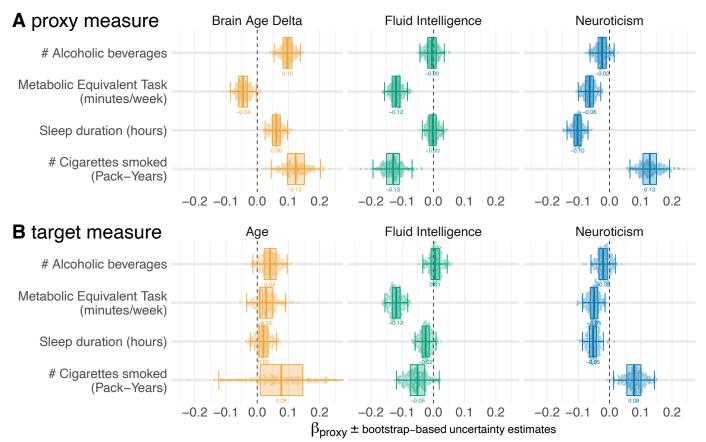
External validity: proxy measures capture ecological health-related factors

Results so far have shown that psychological constructs can be approximated from general-153 purpose inputs such as brain images and sociodemographic variables that are not tailored to 154 measure these latent constructs. Beyond approximating target measures, which are them-155 selves imperfect, can our empirically-derived proxy measures capture complementary facets 156 of real-world behavior? To address this question we studied the link between the three proxy 157 measures studied -built via brain age, fluid intelligence and neuroticism- and various health 158 behavior (sleep, physical exercise, alcohol and tobacco consumption). These behaviors are 159 more ecological probes of mental health than questionnaires or lab-based measures and 160 are potentially linked in multiple ways to our proxy measures. We, hence, modeled them as 161 weighted sums of predicted brain-age delta, fluid intelligence and neuroticism using multiple 162 linear regression. To avoid any form of circularity, we used the out-of-sample predictions for 163 all three proxy measures, applied on the generalization dataset that was not used for building 164 the machine learning models. We derived the brain-age delta by subtracting the actual 165 age from the predicted age. To mitigate brain age bias (Le et al., 2018), we deconfounded 166 health-related habits for their association with actual age (Engemann et al., 2020; Smith 167



Extreme–group classification with proxy measures derived from sociodemographics and brain imaging

Figure 3. Classification analysis from imaging, sociodemographics and combination of both data. For classification of extreme groups instead of continuous regression, we split the data into low vs high groups based on 33_{rd} and 66_{th} percentiles. Visual conventions follow Figure 2. We report the accuracy in AUC. Models including sociodemographics performed visibly better than models purely based on brain imaging. Differences between brain-imaging and sociodemographics appeared less pronounced as compared to the fully-fledged regression analysis. For the average out-of-sample predictions, the probability of the observed performance under the null-distribution and the uncertainty of effect sizes were formally probed using permutation tests and bootstrap-based confidence intervals (Table 2). Corresponding statistics for the baseline performance of models solely based on brain imaging (vertical dotted lines) are presented in Table S2.



Specific associations of proxy and target measures with health-related habits

Figure 4. Proxy measures show systematic and complementary out-of-sample associations with health-related habits. To probe external validity of the proxy measures, we investigated their out-of-sample associations with ecological indicators of mental health (sleep duration, time spent with physical exercise, number of alcoholic beverages and cigarettes consumed). To tease apart complementary and redundant effects, we constructed multiple linear regression models on out-of-sample predictions combining the proxy measures (**A**) from Figure 2. For comparison, we repeated the analysis using the actual target measures (**B**) observed on the generalization data. Regression models are depicted rows-wise. Box plots summarize the uncertainty distribution of target-specific (color) regression coefficients with whiskers indicating two-sided 95% uncertainty intervals (parametric bootstrap). A random subset of 200 out of 10000 coefficient-draws is illustrated by dots. The average coefficient estimate is annotated for convenience. At least two distinct patterns emerged: either the health outcome was specifically associated with one proxy measures (brain age delta and number of alcoholic beverages) or multiple measures showed additive associations with the outcome (*e.g.* number of pack years smoked). Finally, target measures (**B**) show noisier associations than proxy measures (**A**), though none of the significant associations changed direction. For additional findings, please consider the supplement of Figure 4:

Figure 4 – Figure supplement 1. Marginal associations between proxy measures and health-related habits.

et al., 2019a).

The estimated regression coefficients, capturing partial correlations, revealed specific 169 as well as complementary associations between the proxy measures and health-related 170 behavior (Figure 4). A marginal association analysis shows similar patterns, indicating that the 171 relationships hold also when considering the proxy measures in isolation (Figure 4 – Figure 172 supplement 1). Elevated brain-age delta was consistently associated with increased number 173 of alcoholic beverages. These latter proxy measures showed no consistent association with 174 alcohol consumption (Figure 4, first row). Level of physical exercise -measured through the 175 number of minutes spent weekly with metabolic equivalent tasks- consistently associated 176 with the scores in all three predicted targets, suggesting independent associations (Figure 4, 177

> second row). This may seem counter-intuitive but could simply point at the possibility that 178 people with higher test scores, as a tendency, have a more sedentary life style. Sleep 179 duration was independently associated with brain age delta and predicted neuroticism but in 180 opposite directions (Figure 4, third row): increased sleep duration consistently went along 181 with elevated brain age, but lower levels of predicted neuroticism. No consistent effect 182 emerged for fluid intelligence. Increased cumulative numbers of cigarettes smoked was 183 independently and consistently associated with all predicted targets (Figure 4, last row): 184 Intensified smoking went along with elevated brain age delta and elevated neuroticism but 185 lower fluid intelligence. 186

> The question remains whether the proxy measures bring any additional value compared to the original target measures that they were derived from. Studying the association of these original target measures with the health-related habits shows similar trends: associations with the same signs as with the proxy measures (Figure 4, B). However, these associations were more noisy or less marked as those seen with the proxy measures.

> These results demonstrates that the proxy measures capture well health-related habits, 192 potentially better than the original target measures, and in a complementary way across the three measures. 194

Discussion

In this study, we have extended the brain-age approach for neuroimaging to the wider notion 196 of empirical proxy measures. Guided by machine learning, we have derived empirical 197 approximations of traditional, extensively validated target measures from psychology. Beyond 198 biological age, we focused on cognitive capacity (accessed by the fluid-intelligence test) and 199 negative emotionality (accessed by the neuroticism questionnaire). Our proxy measures 200 were derived from data not explicitly designed to assess specific latent constructs: brain 201 imaging data and heterogeneous sociodemographic descriptors. We observed that the 202 combination of brain imaging and target-specific sociodemographic inputs often improved 203 approximation performance. On the held-out data that was not used for model construction, 204 we found important associations between all proxy measures and ecological health indicators. 205 These associations were often complementary and useful beyond the information conveyed 206 by the approximated targets. 207

Constructs of mental-health can be accessed from general-purpose data

Brain age has served as landmark in this study, both conceptually and empirically. It has 209 been arguably the most discussed candidate for a surrogate biomarker in the neuroimaging 210 literature so far (Cole et al., 2015; Dosenbach et al., 2010; Smith et al., 2019a). With mean 211 absolute errors around 4 years, up to 67% variance explained, and AUC-scores up to 0.93 in 212 the classification setting, our results compare favorably to the recent brain-age literature within 213 the UK Biobank (Cole et al., 2017; Smith et al., 2020) and in other datasets (Engemann et al., 214 2020; Liem et al., 2017), though we relied on non-optimized standard inputs and algorithms 215 and not deep learning (He et al., 2018). Applying the same approach to other behavioral 216 outcomes that probe psychological constructs, namely fluid intelligence and neuroticism, we 217 found that these were considerably harder to approximate from general brain imaging data 218 or sociodemographic descriptors. 219

It is important to recapitulate that approximation quality on the three targets investigated has a different meaning, as these are measured differently. On the one hand, age is a physical variable measured with meaningful units (years) on a ratio scale (Stevens et al., 1946) (Selma 222 221 222

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is twice as old as Bob). On the other hand, psychometric scores such as fluid intelligence 223 -measured via socially-administered performance tests- and neuroticism -measured by self-224 assessment via questionnaires- are unit-free scores resulting from operationalized counting, 225 which provokes ambiguity regarding the level of measurement (Borsboom, 2005). Their 226 implied scales may be considered as interval (the difference between Bob's and Selma's 227 intelligence is -0.1 standard deviations) if not rather ordinal (Bob's intelligence was ranked 228 below Selma's) (Stevens et al., 1946). In day-to-day psychological practice, these scores 229 are often used via practically-defined thresholds, e.g. school admission or pilot candidate 230 selection in aviation (Carretta, 2011; Carretta and Ree, 1994). Approximations of these 231 measures via empirically-defined proxies should thus be subjected to different standards: 232 Brain-age prediction should be gauged accordingly to its natural continuous scale; we 233 observed more than 50% of the variance explained. Instead, approximation of psychometric 234 scores might be more appropriately gauged via implicit thresholds, hence, discrimination 235 tasks. With the corresponding metrics, the receiver-operator characteristics (ROC) and its 236 AUC-score, all proxy measures approached or exceeded a performance of 0.80 deemed 237 relevant in biomarker development (Perlis, 2011), though to be fair, they approximated 238 established psychometric targets (proxy measures themselves) and not a medical condition. 239

Nevertheless, the out-of-sample associations of the approximated constructs - the proxy 240 measures- with health-related habits (Figure 4) paint a more complete picture of their value. 241 Sleep duration, minutes spent exercising, and the amount of alcoholic drinks or cigarettes 242 consumed were specifically and complementarily associated with all proxy measures on 243 more than 4000 held-out individuals. In other words, we found multiple statistically important 244 associations with proxy measures fluid intelligence and neuroticism that were not accounted 245 for by brain age. Compared to the traditional measures (Figure 4 B), the associations 246 between these proxy measures and ecological behavioral traits were less noisy, hence more 247 consistent, regardless of their approximation quality (Figure 4 A). This may seem surprising 248 at first, but the target measures are themselves noisy and of imperfect ecological validity. 249 Conversely, the proxy measures are assembled via a richer phenotyping than the target 250 measures, drawing from both fine sociodemographics and brain signals, which can help 251 refining them. 252

The benefits offered by brain data depend on the approximated construct

All brain-derived approximations were statistically meaningful. Yet, only for age prediction, 254 imaging data by itself led to convincing performance levels. Combining brain-imaging data 255 to sociodemographics led to systematically enhanced performance for predicting age and, 256 less strongly, fluid intelligence (Table 1). On the other hand, for neuroticism, including brain 257 imaging never substantially improved the approximation. Does this mean that brain imaging 258 could be avoided in practice when approximating latent constructs? Such a view is probably 259 misleading as the numerical quality of the approximation is not the only thing that matters in a 260 proxy measure. The interest in building a proxy measure of age from brain imaging is justified 261 by its interpretation as an index of precocious or accelerated biological aging (Cole et al., 262 2015, 2017; Smith et al., 2020). In contrast, it is not yet clear that an age delta built from 263 sociodemographic inputs -along the lines of a "social age"- supports such interpretation. 264 From this point of view one may even prefer purely brain-based assessment of individual 265 aging, though sociodemographics probably provide important context to the brain images. 266

For fluid intelligence and neuroticism the situation seemed more complex. For both targets, ²⁶⁷ the best performing sociodemographic model was based on inputs semantically close to ²⁶⁸ the construct of interest, *i.e.*, education details for fluid intelligence and mood & sentiment ²⁶⁹ for neuroticism. While those results reinforce the construct validity of the measure, they ²⁷⁰

also come with a certain risk of circularity. In particular, the causal role of those predictors 271 is not necessarily clear as better educational attainment is heritable itself (Krapohl et al., 272 2014) and may reinforce existing cognitive abilities rather than simply resulting from them. 273 Similarly, prolonged emotional stress due to life events may exacerbate existing dispositions 274 to experience negative emotions captured by neuroticism (Colodro-Conde et al., 2018), traits 275 which in turn commonly help accumulate stressful life events (Lahey, 2009). Nevertheless, for 276 fluid intelligence but not neuroticism, brain imaging added incremental value when combined 277 with various sociodemographic predictors. This may suggest that the cues for neuroticism 278 conveyed by brain imaging were already present in various sociodemographic predictors, 279 potentially hinting at common causes. 280

It may be worthwhile to revisit the frequently reported difficulty to predict complex traits 281 from brain imaging- especially fMRI (Dubois et al., 2018a,b; Liem et al., 2017; Maglanoc 282 et al., 2020). This may not be entirely surprising at a theoretical level as it has even been 283 argued that psychometric measures of complex traits may not map to biological mechanisms 284 in simple ways (Yarkoni, 2015). Of course, this does not preclude the investigation of their 285 brain correlates and mechanisms (Cole et al., 2015; Cox et al., 2019a; Kievit et al., 2018a; 286 Shackman et al., 2016). It rather emphasizes the importance of searching for appropriate 287 signals and representations supporting the given modeling goals (Bzdok and Ioannidis, 2019). 288 As a speculation, some traits could be tightly linked to the current predominant behavior that 289 may be poorly reflected by resting-state recordings. To consider an extreme counter example, 290 disorders of consciousness— a stable trait induced by severe brain injuries— manifest 291 themselves in systematically and intensely altered brain activity, hence, can be robustly 292 detected from fMRI- and EEG-signals regardless of the present stimulation (Demertzi et al., 293 2019; Engemann et al., 2018). In this context, the recent turn towards naturalistic stimuli and 294 movies (Hasson et al., 2010; Jääskeläinen et al., 2016; Nummenmaa et al., 2012; Sonkusare 295 et al., 2019; Venkatesh et al., 2020) may be promising as trait-level differences in emotion 296 and cognition may need to be systematically provoked by potent stimuli, e.g., emotionally 297 charging or cognitively demanding cinematic content. 298

Empirically-derived proxy measures: From validity to practical utility

The validity of constructs and their measures remains a challenging question (Borsboom, 300 2005; Borsboom et al., 2004; Cronbach and Meehl, 1955). Here, we have demonstrated 301 reasonable out-of-sample generalization for our proxy measures. Yet, generalization per-302 formance in itself, arguably, only yields an upper bound for validity of the measure for a 303 target construct, comparable internal-consistency checks and re-test reliability in classical 304 psychometrics. Even a perfect approximation may be limited by the quality of the target 305 measure as fluid intelligence and neuroticism are notoriously difficult to measure without 306 noise. In our study, the construct validity of the corresponding proxy measures is supported 307 by the substantial gain in prediction performance brought by related information, namely 308 education history and mental-health variables respectively (Figure 2). Moreover, association 309 with health-relevant habits brings external validity to the proxy (Figure 4). For example, the 310 complementary patterns that emerged can be related to traditional construct semantics: High 311 consumption of cigarettes is typically associated with neuroticism (Terracciano and Costa Jr, 312 2004) and excessive drinking may lead to brain atrophy and cognitive decline (Topiwala et al., 313 2017), both common correlates of brain age (Liem et al., 2017; Wang et al., 2019). 314

This raises the question of the practical utility of such empirically-derived proxy measures: 315 Can these empirically-derived proxy measures substitute specific psychometric instruments? 316 The present study does not claim to give an unequivocal answer to this question as the 317 utility of proxy measures will depend on the practical context. A specialized mental-health 318

professional may prefer an established routine for clinical assessment, relying on scores such 319 as intelligence tests and personality-scales like neuroticism, and potentially applying implicit 320 experience-based thresholds. Based on our findings, inclusion of brain imaging may even 321 seem to yield diminishing returns when approximating high-level psychological traits. Yet, it 322 mays simply be a matter of time until more effective acquisition protocols will be discovered 323 alongside signal representations supporting predictive modeling. While the cost of including 324 brain imaging may seem exorbitant, whenever available, its inclusion seems to be a "safe 325 bet" as machine learning is capable at selecting relevant inputs (Engemann et al., 2020) and 326 costs of MRI-acquisition can be amortized by baseline clinical usage. Moreover, our study 327 shows that the associations of the proxy measures to health habits compare favorably to the 328 original target measures. As such, the proxy measures may open new doors when tailored 329 assessment of latent constructs is not applicable to due lack of specialized mental-health 330 workforce or sheer cost. For instance, they may bring mental-health assessment in research 331 endeavors on large populations, e.g., for etiology, nosology, or typical epidemiology questions 332 such as risk factors or treatment evaluation. In addition, results derived on large populations 333 can be transferred to clinical data with finer mental-health assessment, e.g., smaller cohorts, 334 possibly leveraging dedicated methods (He et al., 2020; Pan and Yang, 2009). Relying on 335 three proxy measures rather than the brain age alone promises a wider array of applications. 336

Limitations

This study has validated proxy measures of three target constructs. The selection of these 338 targets was guided by literature review as well as the goal to find representative health-related 339 measures with complementary semantics. Additional constructs and psychometric tools 340 could have been visited. Intelligence can be characterized by multiple facets. The broader 341 construct of intelligence as a general factor -g-factor- is often estimated using latent factor 342 models on multiple correlated tests. While g-factor modeling can be interesting for its own 343 sake, we are less interested in normative assessment of intelligence but rather in capturing 344 inter-individual variance related to cognitive capacity as a situational fitness signal. Such 345 variations have been repeatedly linked to mental-health conditions (Khandaker et al., 2018). 346 Likewise, there is a wealth of questionnaires designed to measure negative emotionality 347 and neuroticism specifically. Yet, we could study only that available in the UK-Biobank 348 data, the EPQ neuroticism scale. A complementary approach, leading to different scientific 349 guestions, would be to estimate latent factors by pooling all non-imaging data semantically 350 related to neuroticism (Maglanoc et al., 2020). Rather, we chose to consider established 351 target measures "as is" instead of derivatives to avoid bringing in additional measure-validity 352 considerations. Nevertheless, our framework encourages future studies targeting more 353 sophisticated representations of latent constructs. 354

Second, while the study was clinically motivated, it falls short of directly testing the clinical relevance of estimated proxy measures. Indeed, even in a very large generalpopulation cohort such as the UK Biobank, there are only a few hundred diagnosed cases of mental disorders (ICD-10 mental-health diagnoses from the F chapter) with brain-imaging data available. This challenge highlights the practical importance of studying mental as a continuous, in addition to diagnosed conditions. In this direction, our analysis of health-related habits does provide some clinical relevance. 361

Finally, our study falls short of presenting fine-grained spatial analysis of the imaging data. This work has focused on the approximation quality of proxy measures, relying on methods that are not designed for fine-grained inference on predictors (Bzdok et al., 2018), though future work could explore post-hoc explanations (Biecek, 2018). Our analysis comparing the quality of models helps isolating major explanatory factors, yet does not provide brain 366

mapping (Cole, 2020; Cox et al., 2019a; Kievit et al., 2018a).

Conclusion

Empirically-derived proxy measures targeting age, fluid intelligence and neuroticism reveal 369 complementary facets of real-world behavior that contribute to maintaining mental health. As 370 the relative importance of brain imaging and sociodemographics varies with the approximated 371 target, we recommend generously including all available data and approximating as many 372 targets as possible while letting machine learning perform the labor of integration. We believe 373 that further developing and using proxy measures for constructs that are difficult to assess is 374 a promising agenda for mental-health research. Therefore, we have made all data analysis 375 and visualization source code available on Github: https://github.com/KamalakerDadi/ 376 proxy_measures_2020. 377

Materials and Methods

Dataset

The United Kingdom Biobank (UKBB) database is to date the most extensive large-scale 380 cohort aimed at studying the determinants of the health outcomes in the general adult 381 population. The UKBB is openly accessible and has extensive data acquired on 500 000 382 individuals aged 40-70 years covering rich phenotypes, health-related information, brain-383 imaging and genetic data (Collins, 2012). Participants were invited for repeated assessments, 384 some of which included MR imaging. For instance, cognitive tests that were administered 385 during an initial assessment were also assessed during the follow-up visits. This has enabled 386 finding for many subjects at least one visit containing all heterogeneous input data needed to 387 develop the proposed proxy measures. The study was conducted using the UKBB Resource 388 Applixaction 23827.

Participants

All participants gave informed consent. The UKBB study was examined and approved by the 391 North West Multi-centre Research Ethics Committee. We considered participants who have 392 responded to cognitive tests, questionnaires, and have access to their primary demographics 393 and brain images (Sudlow et al., 2015). Out of the total size of UKBB populations, we found 394 11 175 participants who had repeated assessments overlapping with the first brain imaging 395 release (Miller et al., 2016). The demographics are 51.6% female (5572) and 48.3% male 396 (5 403), and an age range between 40-70 years (with a mean of 55 years and standard 397 deviation of 7.5 years). Out of the complete analysis set, 5587 individuals were used in 398 the study to train the model and remaining subjects were set aside as a held-out set for 399 generalization testing (see section Model development and generalization testing). 400

To establish specific comparisons between models based on sociodemographics, brain 401 data or their combinations we exclusively considered the cases for which MRI scans were 402 available. The final sample sizes used for model construction and generalization testing 403 then depended on the availability of MRI: For age and fluid intelligence, our random splitting 404 procedure (Model development and generalization testing) yielded 4203 cases for model 405 building and 4157 for generalization. For cases with valid neuroticism assessment, fewer 406 brain images were available, which yielded 3550 cases for model building and 3509 for 407 generalization. 408

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Data acquisition

Sociodemographic data (non-imaging) was collected with self-report measures administered 410 through touchscreen questionnaires, complemented by verbal interviews, physical measures, 411 biological sampling and imaging data. MRI data were acquired with the Siemens Skyra 3T 412 using a standard Siemens 32-channel RF receiver head coil (Alfaro-Almagro et al., 2018). 413 We considered three MR imaging modalities as each of them potentially captures unique 414 neurobiological details: structural MRI (sMRI/T1), resting-state functional MRI (rs-fMRI) and 415 diffusion MRI (dMRI). For technical details about the MR acquisition parameters, please 416 refer to Miller et al. (2016). We used image-derived phenotypes (IDPs) of those distinct 417 brain-imaging modalities, as they provide actionable summaries of the brain measurements 418 and encourage comparability across studies. 419

Target measures

As our target measures for brain age modelign, we use an individual's age at baseline 421 recruitment (UKBB code "21022-0.0"). Fluid intelligence, was assessed using a cognitive 422 battery designed to measure an individual's capacity to solve novel problems that require 423 logic and abstract reasoning. In the UK Biobank, the fluid intelligence test (UKBB code 424 "20016-2.0") comprises thirteen logic and reasoning questions that were administered via 425 the touchscreen to record a response within two minutes for each question. Therefore, 426 each correct answer is scored as one point with 13 points in total¹. Neuroticism (UKBB code "20127-0.0") was measured using a shorter version of the revised Eysenck Personality 428 Questionnaire (EPQ-N) comprised of 12-items (Eysenck et al., 1985). Neuroticism was 429 assessed during Biobank's baseline visit. The summary of the individual's scores ranges 430 from 0 to 12 that assess dispositional tendency to experience negative emotions². 431

Sociodemographic data

In this work, we refer to non-imaging variables broadly as sociodemographics excluding the 433 candidate targets fluid intelligence and neuroticism. To approximate latent constructs from 434 sociodemographics, we included 86 non-imaging inputs (Table S3) which are the collection 435 of variables reflecting each participant's demographic and social factors *i.e.*, sex, age, date 436 and month of birth, body mass index, ethnicity, exposures at early life -e.g. breast feeding, 437 maternal smoking around birth, adopted as a child-education, lifestyle-related variables -e.g. 438 occupation, household family income, household people living at the same place, smoking 439 habits-, and mental-health variables. All these data were self-reported. We then assigned 440 these 86 variables to five groups based on their relationships. Based on our conceptual 441 understanding of the variables, we name assigned them to one out of five groups: 1) mood & 442 sentiment, 2) primary demographics as age, sex, 3) lifestyle, 4) education, 5) early life. We 443 then investigated the intercorrelation between all 86 variables to ensure that the proposed 444 grouping is compatible with their empirical correlation structure Figure S1. 445

The sociodemographic groups had varying amounts of missing data. For e.g. the source 446 of missingness is concerned with the participants lifestyle habits such as smoking and mental 447 health issues (Fry et al., 2017). To deal with this missingness in the data using imputation 448 (Little and Rubin, 1986), we used column-wise replacement of missing information with the 449 median value calculated from the known part of the variable. We subsequently included 450

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¹A complete overview of the 13 individual fluid intelligence items can be seen from this manual https: //biobank.ctsu.ox.ac.uk/crystal/crystal/docs/Fluidintelligence.pdf

²For a complete list of Neuroticism questionnaires can be seen from this manual https://biobank.ctsu. ox.ac.uk/crystal/crystal/docs/MentalStatesDerivation.pdf

an indicator for the presence of imputed for down-stream analysis. Such imputation is well 451 suited to predictive models (Josse et al., 2019). 452

Image processing to derive phenotypes for machine learning

MRI data preprocessing were carried out by UKBB imaging team. The full technical details 454 are described elsewhere (Alfaro-Almagro et al., 2018; Miller et al., 2016). Below, we describe 455 briefly the custom processing steps that we used on top of the already preprocessed inputs. 456

Structural MRI

This type of data analysis on T1-weighted brain images are concerned with morphometry of 458 the gray matter areas *i.e.* the quantification of size, volume of brain structures and tissue 459 types and their variations under neuropathologies or behavior (Lerch et al., 2017). For 460 example, volume changes in gray matter areas over lifetime are associated with: brain aging 461 (Ritchie et al., 2015), general intelligence (Cox et al., 2019b) and brain disease (Thompson 462 et al., 2007). Such volumes are calculated within pre-defined ROIs composed of cortical 463 and sub-cortical structures (Desikan et al., 2006) and cerebellar regions (Diedrichsen et al., 464 2009). We included 157 sMRI features consisting of volume of total brain and grey matter 465 along with brain subcortical structures³. All these features are pre-extracted by UKBB brain 466 imaging team (Miller et al., 2016) and are part of data download. We concatenated all inputs 467 alongside custom-built fMRI features for predictive analysis (feature union). 468

Diffusion weighted MRI

Diffusion MRI enables to identify white matter tracts along principal diffusive direction of water 470 molecules, as well as the connections between different gray matter areas (Behrens et al., 471 2003; Conturo et al., 1999). The study of these local anatomical connections through white 472 matter are relevant to the understanding of neuropathologies and functional organization 473 (Saygin et al., 2016). We included 432 dMRI skeleton features of FA (fractional anisotropy), 474 MO (tensor mode) and MD (mean diffusivity), ICVF (intra-cellular volume fraction), ISOVF 475 (isotropic volume fraction) and OD (orientation dispersion index) modeled on many brain 476 white matter structures extracted from neuroanatomy⁴. For extensive technical details, please 477 refer to de Groot et al. (2013). The skeleton features we included were from category134 478 shipped by the UKBB brain-imaging team and we used them without modification. 479

Functional MRI

Resting-state functional MR images capture low-frequency fluctuations in blood oxygenation 481 that can reveal ongoing neuronal interactions in time forming distinct brain networks (Biswal 482 et al., 1995). Functional connectivity within these brain network can be linked to clinical status 483 (Greicius et al., 2004), to behavior (Miller et al., 2016), or to psychological traits (Dubois 484 et al., 2018b). We also included resting-state connectivity features based on the time-series 485 extracted from Independent Component Analysis (ICA) with 55 components representing 486 various brain networks extracted on UKBB rfMRI data (Miller et al., 2016). These included 487 the default mode network, extended default mode network and cingulo-opercular network, 488 executive control and attention network, visual network, and sensorimotor network. We 489

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³Regional grey matter volumes http://biobank.ctsu.ox.ac.uk/crystal/label.cgi?id=1101 Subcortical volumes http://biobank.ctsu.ox.ac.uk/crystal/label.cgi?id=1102

⁴Diffusion-MRI skeleton measurements http://biobank.ctsu.ox.ac.uk/crystal/label.cgi?id=134

Index	Name	# variables	# groups
1	brain volumes (sMRI)	157	1
2	white matter (dMRI)	432	1
3	functional connectivity (fMRI)	1485	1
4	sMRI, dMRI	589	2
5	sMRI, fMRI	1642	2
6	dMRI, fMRI	1917	2
7	sMRI, dMRI, fMRI (full MRI)	2074	3

Table 3.	Imaging-based models.
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Table 4. Non-imaging baseline models or sociodemographic models based on single group. Variables in each group are described at corresponding section: Sociodemographic data.

Index	Name	# variables
1	Mood & Sentiment (MS)	25
2	Age, Sex (AS)	5
3	Life style (LS)	45
4	Education (EDU)	2
5	Early Life (EL)	9

measured functional connectivity in terms of the between-network covariance. We estimated the covariance matrices using Ledoit-Wolf shrinkage (Ledoit and Wolf, 2004). To account for the fact that covariance matrices live on a particular manifold, *i.e.*, a curved non-Euclidean space, we used the tangent-space embedding to transform the matrices into a Euclidean space (Sabbagh et al., 2019; Varoquaux et al., 2010) following recent recommendations (Dadi et al., 2019; Pervaiz et al., 2020). For predictive modeling, we then vectorized the performed with NiLearn (Abraham et al., 2014).

Comparing predictive models to approximate target measures

Imaging-based models

First, we focused on purely imaging-based models based on exhaustive combinations of the three types of MRI modalities (see Table 3 for an overview). This allowed us to study potential overlap and complementarity between the MRI-modalities. Preliminary analyses revealed that combining all MRI data gave reasonable results with no evident disadvantage over particular combinations of MRI modalities (Figure 2 – Figure supplement 1), hence, for simplicity, we only focused on the full MRI model in subsequent analyses.

Sociodemographic models

We composed predictive models based on non-exhaustive combinations of different types of sociodemographic variables. To investigate the relative importance of each class of sociodemographic inputs, we performed systematic model comparisons. We were particularly interested in studying the relative contributions of early-life factors as compared to factors related to more recent life events such as education as well as factors related to current circumstances such as mood & sentiment and life-style. The resulting models based on distinct groups of predictors are listed in Table 4 (for additional details see Table S3 and Figure S1).

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Hyperparameter	Values
Impurity criterion	Mean squared error
Maximum tree depth	5, 10, 20, 40, full depth
Fraction of features for split	1, 5, "log2", "sqrt", "complete"
Number of trees	250

Table 5. Random forest hyperparameters and tuning with grid search (5 fold cross-validation).

Table 6. Number of samples for classification analysis (N).

# groups	Age	Fluid intelligence	Neuroticism
1		1108	1054
2	1200	898	1020

Combined imaging and sociodemographic models

In the next step, we were interested in how brain-related information would interact within each of these sociodemographic models. For example, information such as the age of an individual, or the level of education, may add important contextual information to brain images. We therefore considered an alternative variant for each of the models in Table 4 that included all MRI-related features (2 074 additional features) as described at section Image processing to derive phenotypes for machine learning.

Predictive model

Linear models are recommended as default choice in neuroimaging research (Dadi et al., 522 2019; Poldrack et al., 2020) especially when datasets include fewer than 1000 data points. 523 In this study approximated targets generated by distinct underlying mechanisms based on 524 multiple classes of heterogenous input data with several thousands of data points. We 525 hence chose the non-parametric random forest algorithm that can be readily applied on 526 data of different units for non-linear regression and classification (Breiman, 2001) with mean 527 squared error as impurity criterion. To improve computation time we fixed tree-depth to 528 250 trees, a hyper-parameter that is not usually not tuned but set to a generous number 529 as performance plateaus beyond a certain number of trees (Hastie et al., 2005, ch. 15). 530 Preliminary analyses suggested that additional trees would not have led to substantial 531 improvements in performance. We used nested cross-validation (5-fold grid search) to tune 532 the depth of the trees as well as the number of variables considered for splitting (see Table 5 533 for a full list of hyper-parameters considered). 534

Classification analysis. We also performed classification analysis on the continuous 535 targets. For this purpose, we discretized the targets into extreme groups based on the 33rd 536 and 66_{th} percentiles (see Table 6 for the number of classification samples per group). We 537 were particularly interested in understanding whether model performance would increase 538 when moving toward classifying extreme groups. For this analysis, we considered all 539 three types of models (full MRI 2074 features from imaging-based models see section 540 Imaging-based models, all sociodemographics variables, total 86 variables see section 541 Sociodemographic models), combination of full MRI and all sociodemographics, a total 2160 542 variables see section Combined imaging and sociodemographic models. When predicting 543 age, we excluded the age & sex sociodemographic block from all sociodemographic variables 544 which then yielded a total of 81 variables. To assess the performance for classification 545 analysis, we used the area under the curve (AUC) of the receiver operator characteristic 546 (ROC) as an evaluation metric (Poldrack et al., 2020). 547

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Model development and generalization testing

Before any empirical work, we generated two random partitions of the data, one validation dataset for model construction and one held-out generalization dataset for studying out-of-sample associations using classical statistical analyses. 551

For cross-validation, we then subdivided the validation set into 100 training- and testing splits following the Monte Carlo resampling scheme (also referred to as shuffle-split) with 10% of the data used for testing. To compare model performances based on paired tests, we used the same splits across all models. Split-wise testing performance was extracted and carried forward for informal inference using violin plots (Figure 2, Figure 3). For generalization testing, predictions on the held-out data were generated from all 100 models from each cross-validation split.

On the held-out set, unique subject-wise predictions were obtained by averaging across folds and occasional duplicate predictions due to Monte Carlo sampling which could produce multiple predictions per subject⁵. Such strategy is known as CV-bagging (Varoquaux et al., 2017b) and can improve both performance and stability of results⁶. The resulting averages were reported as point estimates in Figures 2,3, and 2 – Figure supplement 1 and used as proxy measures in the analysis of health-related behaviors Figure 4. 564

Statistical analysis

Resampling statistics for model comparisons on the held-out data

To assess the statistical significance of the observed model performance and the differences in performance between the models, we computed resampling statistics of the performance metrics on the held-out generalization data not used for model construction (Gemein et al., 2020). Once unique subject-wise predictions were obtained on the held-out generalization data by averaging the predictions emanating from each fold of the validation set (cv-bagging), we computed null- and bootstrap-distributions of the observed test statistic on the held-out data, i.e., R^2 score for regression and *AUC* score for classification. 567

Baseline comparisons. To obtain a p-value for baseline comparisons (*could the prediction performance of a given model be explained chance?*) on the held-out data, we permuted targets 10 000 times and then recomputed the test statistic in each iteration. P-values were then defined as the probability of the test statistic under null distribution being larger than the observed test statistic. To compute uncertainty intervals, we used bootstrap, recomputing the test statistic after resampling 10 000 times with replacement and reporting the 2.5 and 97.5 percentiles of the resulting distribution.

Pairwise comparisons between models. For model comparisons, we considered the 581 out-of-sample difference in R^2 or AUC between any two models. To obtain a p-value for model 582 comparisons (could the difference in prediction performance between two given models be 583 explained chance?) on the held-out data, we permuted the scores predicted by model A and 584 model B for every single prediction 10 000 times and then recomputed the test statistic in 585 each iteration. We omitted all cases for which only predictions from one of the models under 586 comparison was present. P-values were then defined as the probability of the absolute of the 587 test statistic under null distribution being larger than the absolute observed test statistic. The 588 absolute was considered to account for differences in both directions. Uncertainty intervals 589 were obtained from computing the 2.5 and 97.5 percentiles of the bootstrap distribution 590

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⁵We ensured prior to computation that with 100 CV-splits, predictions were available for all subjects.

⁶The use of CV-bagging can explain why on figures 2,3, and 2 – Figure supplement 1 the performance was sometimes slightly better on the held-out set compared to the cross-validation on the validation test.

Family	eid	Variables
Alcohol*	1568-0.0	Average weekly red wine intake
	1578-0.0	Average weekly champagne plus white wine intake
	1588-0.0	Average weekly beer plus cider intake
	1598-0.0	Average weekly spirits intake
	1608-0.0	Average weekly fortified wine intake
	5364-0.0	Average weekly intake of other alcoholic drinks
Physical activity	22040-0.0	Summed MET minutes per week for all activity
Smoking	20161-0.0	Pack years of smoking
Sleep	1160-0.0	Sleep duration

Table 7. Extra health variables used for correlation analysis with subject-specific predicted scores.

*We computed a compound drinking score by summing up all variables from the alcohol family

based on 10 000 iterations. Here, predictions from model A and model B were resampled using identical resampling indices to ensure a meaningful paired difference. 592

Out-of-sample association between proxy measures and health-related habits

Computation of brain age delta and de-confounding For association with health-contributing habits (Table 7), we computed the brain age delta as the difference between predicted age and actual age:

$$BrainAge \Delta = Age_{predicted} - Age \tag{1}$$

As age prediction is rarely perfect, the residuals will still contain age-related variance which commonly leads to brain age bias when relating the brain age to an outcome of interest, e.g., sleep duration (Le et al., 2018). To mitigate leakage of age-related information into the statistical models, we employed a de-confounding procedure in line with Smith et al. (2019b) and (Engemann et al., 2020, eqs. 6-8) consisting in residualizing a measure of interest (e.g. sleep duration) with regard to age through multiple regression with quadratic terms for age. To minimize computation on the held-out data, we first trained a model relating the score of interest to age on the validation set to then derive a de-confounding predictor for the held-out data amounts to computing an age-residualized predictor *measure*_{resid} from the measure of interest (*e.g.* sleep duration) by applying the following quadratic fit on the validation data:

$$measure_{validation} = age_{validation} \times \beta_{val1} + age_{validation}^2 \times \beta_{val2} + \epsilon$$
(2)

The de-confounding predictor was then obtained by evaluating the weights β_{val1} and β_{val2} ⁵⁹⁴ obtained from Equation 2 on the generalization data: ⁵⁹⁵

measure_{deconfounding} =
$$age_{generalization} \times \beta_{val1} + age_{generalization}^2 \times \beta_{val2}$$
 (3)

We performed this procedure for all target measures, to study associations not driven by the effect of age. 597

Health-related habits regression We then investigated the joint association between proxy measures of interest and health-related habits (Table 7) using multiple linear regression. For simplicity, we combined all brain imaging and all sociodemographics variables (Figure 2, Figure 2 – Figure supplement 1, Figure 2 – Figure supplement 2). The ensuing model can be denoted as

 $measure = measure_{deconfounding} \times \beta_1 + BrainAge \times \Delta \beta_2 + PredFluidInt \times \beta_3 + PredNeurot \times \beta_4 + \epsilon,$

(4)

where *outcome_{resid}* is given by Equation 2. Prior to model fitting, rows with missing inputs were omitted. For comparability, we then applied standard scaling on all outcomes and all predictors.

The parametric bootstrap was a natural choice for uncertainty estimation, as we used standard multiple linear regression which provides a well defined procedure for mathematically quantifying its implied probabilistic model. Computation was carried out using sim function from the arm package as described in Gelman and Hill (2006, Ch.7,pp.142-143). This procedure can be intuitively regarded as yielding draws from the posterior distribution of the multiple linear regression model under the assumption of a uniform prior. For consistency with previous analyses, we computed 10000 draws.

Software

Preprocessing and model building were carried out using Python 3.7. The NiLearn library 614 was used for processing MRI inputs (Abraham et al., 2014). We used the scikit-learn library for 615 machine learning (Pedregosa et al., 2011). For statistical modeling and visualization we used 616 the R-language (R Core Team, 2019) (version 3.5.3) and its ecosystem: data.table for high-617 performance manipulation of tabular data, ggplot (Clarke and Sherrill-Mix, 2017; Wickham, 618 2016) for visualization and the arm package for parametric bootstrapping (Gelman and Su, 619 2020). All data analysis code is shared on GitHub: https://github.com/KamalakerDadi/ 620 proxy_measures_2020. 621

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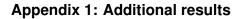
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Supporting Information



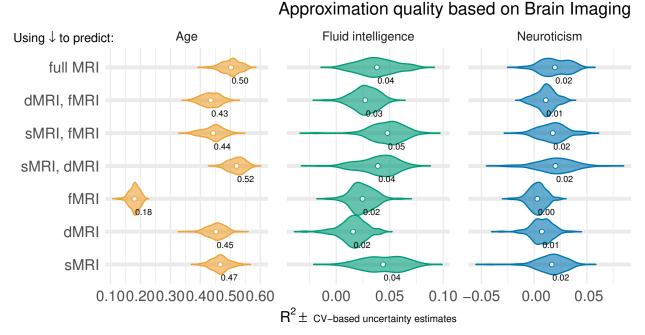


Figure 2 – Figure supplement 1. Prediction of individual differences in proxy measures from MRI. Approximation performance using multiple MR modalities on the validation dataset: sMRI, dMRI, rfMRI and their combinations (see Table 3). Visual conventions as in Figure 2. One can see that prediction of age was markedly stronger than prediction of fluid intelligence or prediction of neuroticism. As a general trend, models based on multiple MRI modalities tended to yield better prediction. For simplicity, we based subsequent analyses on the full model based on all MRI data.

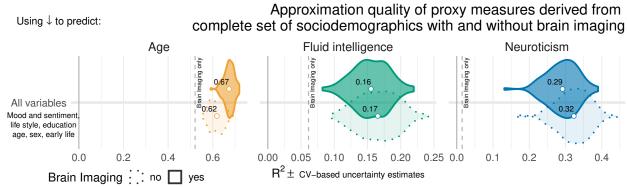


Figure 2 – Figure supplement 2. Approximation performance using all sociodemographic data. Approximation performance using all sociodemographic variables with or without brain imaging included on the validation dataset. Visual conventions as in Figure 2. The performance was highly to the best performing models within each target Figure 2, *i.e.*, life style for age, education for fluid intelligence and mood & sentiment for neuroticism. This suggests that for each target those specific blocks of predictors were sufficiently explaining the performance. For simplicity, we based subsequent analyses in Figure 3 and Figure 4 on all sociodemographic variables.

939

Target	R ² _{observed}	p-value	Cl _{low}	Cl _{high}
Age	0.521	1×10^{-4}	0.502	0.538
Fluid intelligence	0.061	1×10^{-4}	0.052	0.070
Neuroticism	0.015	1×10^{-4}	0.005	0.024

Table S1. Regression statistics on the held-out set for purely MRI-based approximation.

 Table S2. Classification difference statistics on the held-out set for MRI-based approximation.

Target	AUCobserved	p-value	Cl _{low}	Cl _{high}
Neuroticism	0.590	1×10^{-4}	0.566	0.614
Age	0.916	1×10^{-4}	0.905	0.927
Fluid intelligence	0.667	1×10^{-4}	0.643	0.690

Marginal associations of proxy and target measures with health-related habits

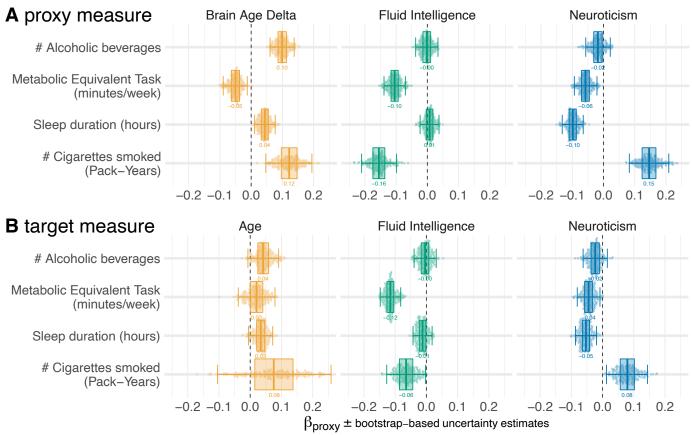


Figure 4 – Figure supplement 1. Marginal associations between proxy measures and health-related habits. Marginal (instead of conditional) estimates using univariate regression. Same visual conventions as in Figure 4.

Appendix 2: Sociodemographic variables

Group	UKBB code	Variables
Mood & Sentiment	2040-2.0	Risk taking
	4526-2.0	Happiness
	4537-2.0	Work/job satisfaction
	4548-2.0	Health satisfaction
	4559-2.0	Family relationship satisfaction
	4570-2.0	Friendships satisfaction
	4581-2.0	Financial situation satisfaction
	4598-2.0	Ever depressed for a whole week
	4609-2.0	Longest period of depression
	4620-2.0	Number of depression episodes
	4631-2.0	Ever unenthusiastic/disinterested for a whole week
	4642-2.0	Ever manic/hyper for 2 days
	4653-2.0	Ever highly irritable/argumentative for 2 days
	2050-2.0	Frequency of depressed mood in last 2 weeks
	2060-2.0	Frequency of unenthusiasm / disinterest in last 2 weeks
	2070-2.0	Frequency of tenseness / restlessness in last 2 weeks
	2080-2.0	Frequency of tiredness / lethargy in last 2 weeks
	2090-2.0	Seen doctor (GP) for nerves, anxiety, tension or depression
	2100-1.0	Seen a psychiatrist for nerves, anxiety, tension or depressio
	5375-2.0	Longest period of unenthusiasm / disinterest
	5386-2.0	Number of unenthusiastic/disinterested episodes
	5663-2.0	Length of longest manic/irritable episode
	5674-2.0	Severity of manic/irritable episode
	6145-2.0	Illness, injury, bereavement, stress in last 2 years
	6156-2.0	Manic/hyper symptoms
Age, Sex	31-0.0	Sex
	34-0.0	Year of birth
	52-0.0	Month of birth
	21022-0.0	Age at recruitment
	21003-2.0	Age when attended assessment centre
Education	6138-2.0	Qualifications
	845-2.0	Age completed full time education
	1647-2.0	Country of birth (UK/elsewhere)
	1677-2.0	Breastfed as a baby
	1687-2.0	Comparative body size at age 10
	1697-2.0	Comparative height size at age 10
Early life	1707-2.0	Handedness (chirality/laterality)
-	1767-2.0	Adopted as a child
	1777-2.0	Part of a multiple birth

Table S3. List of variables contained in each block of sociodemographic models: mood & sentiment (MS), Age, Sex (AS), Education (EDU), Early life (EL).

	Table	<i>S3</i>	continued
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	1787-2.0	Maternal smoking around birth
Lifestyle	670-2.0	Type of accommodation lived in
	680-2.0	Own or rent accommodation lived in
	6139-2.0	Gas or solid-fuel cooking/heating
	699-2.0	Length of time at current address
	709-2.0	Number in household
	6141-2.0	How are people in household related to participant
	728-2.0	Number of vehicles in household
	738-2.0	Income before tax
	796-2.0	Distance between home and job workplace
	757-2.0	Time employed in main current job
	767-2.0	Length of working week for main job
	777-2.0	Freq. of travelling from home to job workplace
	6143-2.0	Transport type for commuting to job workplace
	6142-2.0	Current employment status
	806-2.0	Job involves mainly walking or standing
	816-2.0	Job involves heavy manual or physical work
	826-2.0	Job involves shift work
	3426-2.0	Job involves night shift work
	1031-2.0	Freq. of friend/ family visits
	6160-2.0	Leisure/social activities
	2110-2.0	Able to confide
	1239-2.0	Current tobacco smoking
	1249-2.0	Past tobacco smoking
	1259-2.0	Smoking/smokers in household
	1269-2.0	Exposure to tobacco smoke at home
	1279-2.0	Exposure to tobacco smoke outside home
	2644-2.0	Light smokers, at least 100 smokes in lifetime
	2867-2.0	Age started smoking in former smokers
	2877-2.0	Type of tobacco previously smoked
	2887-2.0	Number of cigarettes previously smoked daily
	2897-2.0	Age stopped smoking
	2907-2.0	Ever stopped smoking for 6+ months
	2926-2.0	Number of unsuccessful stop-smoking attempts
	2936-2.0	Likelihood of resuming smoking
	3436-2.0	Age started smoking in current smokers
	3446-2.0	Type of tobacco currently smoked
	3456-2.0	Number of cigarettes currently
		smoked daily (current cigarette smokers)
	3466-2.0	Time from waking to first cigarette
	3476-2.0	Difficulty not smoking for 1 day
	3486-2.0	Ever tried to stop smoking

Table S3 continued

3496-2.0	Wants to stop smoking
3506-2.0	Smoking compared to 10 years previous
5959-2.0	Previously smoked cigarettes on most/all days
6157-2.0	Why stopped smoking
6158-2.0	Why reduced smoking

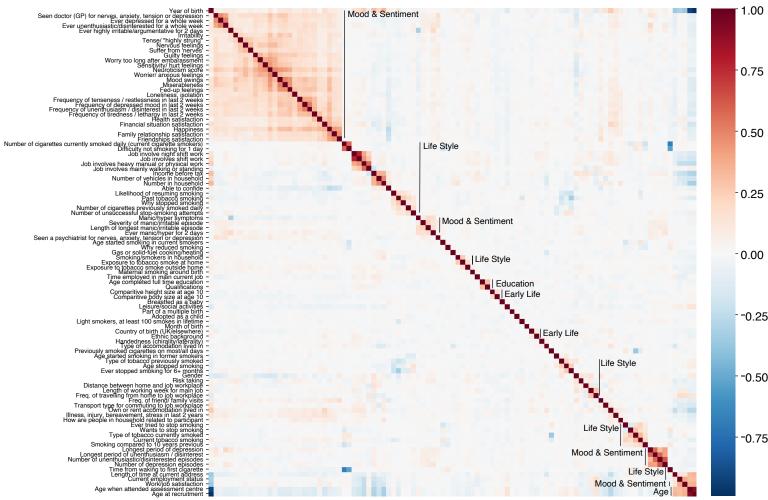


Figure S1. Intercorrelations between sociodemographic inputs. To check the plausibility of the proposed grouping of variables into blocks, we investigated the inter-correlations among the sociodemographic inputs (Table S3). We first applied Yeo-Johnson power transform to the variables yield approximately symmetrical distributions. Then we computed Pearson correlations. One can see that a large majority of variables shows low if any inter-correlations. Strongly inter-correlated blocks emerged, in particular for Mood & Sentiment and Life Style. Note that within the Life Style category many smaller blocks with strong inter-correlation occurred, some of which were obviously related to the circumstance of living such as household or employment status.