Induced Numerical Instabilities in Analytical Pipelines Lead to Impactful Variability in Brain Networks

Gregory Kiar¹, Yohan Chatelain², Pablo de Oliveira Castro³, Eric Petit⁴, Ariel Rokem⁵, Gaël Varoquaux⁶, Bratislav Misic¹, Alan C. Evans^{1†}, Tristan Glatard^{2†}

Abstract

The analysis of brain-imaging data requires complex processing pipelines to support findings on brain function or pathologies. Recent work has shown that variability in analytical decisions, small amounts of noise, or computational environments can lead to substantial differences in the results, endangering the trust in conclusions¹⁻⁷. We explored the instability of results by instrumenting a connectome estimation pipeline with Monte Carlo Arithmetic^{8,9} to introduce random noise throughout. We evaluated the reliability of the connectomes, their features 10,11, and the impact on analysis 12,13. The stability of results was found to range from perfectly stable to highly unstable. This paper highlights the potential of leveraging induced variance in estimates of brain connectivity to reduce the bias in networks alongside increasing the robustness of their applications in the classification of individual differences. We demonstrate that stability evaluations are necessary for understanding error inherent to brain imaging experiments, and how numerical analysis can be applied to typical analytical workflows both in brain imaging and other domains of computational science. Overall, while the extreme variability in results due to analytical instabilities could severely hamper our understanding of brain organization, it also leads to an increase in the reliability of datasets.

Keywords

Stability — Reproducibility — Network Neuroscience — Neuroimaging

¹Montréal Neurological Institute, McGill University, Montréal, QC, Canada; ²Department of Computer Science and Software Engineering, Concordia University, Montréal, QC, Canada; ³ Department of Computer Science, Université of Versailles, Versailles, France; ⁴ Exascale Computing Lab, Intel, Paris, France; 5 Department of Psychology and eScience Institute, University of Washington, Seattle, WA, USA; ⁶ Parietal project-team, INRIA Saclay-ile de France, France; †Authors contributed equally.

The modelling of brain networks, called connectomics, 9 but potentially pave the way for therapeutics 19-23. ² has shaped our understanding of the structure and function 3 of the brain across a variety of organisms and scales over 4 the last decade 11, 14-18. In humans, these wiring diagrams are 6 and show promise towards identifying biomarkers of disease. 7 This can not only improve understanding of so-called "connec-

However, the analysis of brain imaging data relies on complex computational methods and software. Tools are trusted to ₁₂ perform everything from pre-processing tasks to downstream 5 obtained in vivo through Magnetic Resonance Imaging (MRI), 13 statistical evaluation. While these tools undoubtedly undergo 14 rigorous evaluation on bespoke datasets, in the absence of 15 ground-truth this is often evaluated through measures of re-8 topathies", such as Alzheimer's Disease and Schizophrenia, 16 liability^{24–27}, proxy outcome statistics, or agreement with

 $_{17}$ existing theory. Importantly, this means that tools are not $_{53}$ individual and 100 samples total (25 \times 2 \times 2 samples). Struc-24 and it is likely that software instabilities played a role.

29 cations of the observed instabilities on downstream analyses 30 were quantified. We accomplished this through the use of 31 Monte Carlo Arithmetic (MCA)⁸, a technique which enables 32 characterization of the sensitivity of a system to small nu-33 merical perturbations. This is importantly distinct from data 34 perturbation experiments where the underlying datasets are 35 manipulated or pathologies may be simulated, and allows 36 for the evaluation of experimental uncertainty in real-world 37 settings. We explored the impact of numerical perturbations 38 through the direct comparision of structural connectomes, the 39 consistency of their features, and their eventual application in 40 a neuroscience study. We also characterized the consequences 41 of instability in these pipelines on the reliability of derived 42 datasets, and discuss how the induced variability may be har-43 nessed to increase the discriminability of datasets. Finally, 44 we make recommendations for the roles perturbation analyses 45 may play in brain imaging research and beyond.

46 Graphs Vary Widely With Perturbations

48 understanding of the induced variability was required. A sub- 85 tinct, connectomes generated with input perturbations show 49 set of the Nathan Kline Institute Rockland Sample (NKIRS) 86 considerable variability, often reaching deviations equal to 50 dataset²⁹ was randomly selected to contain 25 individuals 87 or greater than those observed across individuals or sessions 51 with two sessions of imaging data, each of which was sub- 88 (Figure 1A; right). Interpretting these results with respect to 52 sampled into two components, resulting in four samples per 89 the distinct MCA environments used suggests that the tested

18 necessarily of known or consistent quality, and it is not un- 54 tural connectomes were generated with canonical determinis-19 common that equivalent experiments may lead to diverging 55 tic and probabilistic pipelines 30,31 which were instrumented 20 conclusions^{1,5–7}. While many scientific disciplines suffer 56 with MCA, replicating computational noise either sparsely 21 from a lack of reproducibility²⁸, this was recently explored 57 or densely throughout the pipelines^{4,9}. In the sparse case, a 22 in brain imaging by a 70 team consortium which performed 58 small subset of the libraries were instrumented with MCA, al-23 equivalent analyses and found widely inconsistent results¹, 59 lowing for the evaluation of the cascading effects of numerical 60 instabilities that may arise. In the dense case, operations are The present study approached evaluating reproducibility 61 more uniformly perturbed and thus the law of large numbers 26 from a computational perspective in which a series of brain 62 suggests that perturbations will quickly offset one-another and 27 imaging studies were numerically perturbed in such a way 63 only dramatic local instabilities will have propagating effects. 28 that the plausibility of results was not affected, and the impli- 64 Importantly, the perturbations resulting from the sparse setting 65 represent a strict subset of the possible outcomes of the dense 66 implementation. The random perturbations are statistically 67 independent from one another across both settings and sim-68 ulations. Instrumenting pipelines with MCA increases their 69 computation time, in this case by multiplication factors of ₇₀ 1.2× and $7\times$ for the sparse and dense settings, respectively⁴. 71 The results obtained were compared to unperturbed (e.g. ref-72 erence) connectomes in both cases. The connectomes were 73 sampled 10 times per sample and once without perturbations, 74 resulting in a total of 4,200 connectomes. Two versions of 75 the unperturbed connectomes were generated and compared 76 such that the absence of variability aside from that induced 77 via MCA could be confirmed.

The stability of connectomes was evaluated through the 79 normalized percent deviation from reference⁴ and the num-80 ber of significant digits (Figure 1). The comparisons were 81 grouped according to differences across simulations, subsam-82 pling of data, sessions of acquisition, or subjects, and accord-83 ingly sorted from most to least similar. While the similarity 47 Prior to exploring the analytic impact of instabilities, a direct 84 of connectomes decreases as the collections become more dis-

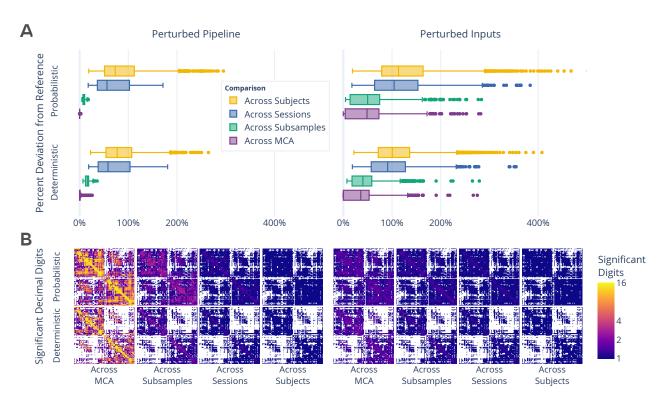


Figure 1. Exploration of perturbation-induced deviations from reference connectomes. (**A**) The absolute deviations, in the form of normalized percent deviation from reference, shown as the across MCA series relative to Across Subsample, Across Session, and Across Subject variations. (**B**) The number of significant decimal digits in each set of connectomes as obtained after evaluating the effect of perturbations. In the case of 16, values can be fully relied upon, whereas in the case of 1 only the first digit of a value can be trusted. Dense and sparse perturbations are shown on the left and right, respectively.

90 pipelines may not suffer from single dominant sources of 105 MCA comparison of connectomes generated with pipeline 91 instability, but that nevertheless there exist minor local in- 106 perturbations show nearly perfect precision for many edges 92 stabilities which may the propagate throughout the pipeline. 107 (approaching the maximum of 15.7 digits for 64-bit data), 93 Furthermore, this finding suggests that instabilities inherent 108 this evaluation uniquely shows considerable drop off in per-94 to these pipelines may mask session or individual differences, 109 formance when comparing networks across subsamplings 95 limiting the trustworthiness of derived connectomes. While 110 (average of < 4 digits). In addition, sparsely perturbed con-96 both pipelines show similar performance, the probabilistic 111 nectomes show no more than an average of 3 significant digits 97 pipeline was more stable in the face of pipeline perturbations 112 across all comparison groups, demonstrating a significant lim-98 whereas the deterministic was more stable to input perturba- 113 itation in the reliability of independent edge weights. The ₉₉ tions (p < 0.0001 for all; exploratory). As an alternative to ₁₁₄ number of significant digits across individuals did not exceed 100 the normalized percent deviation, the stability of correlations 115 a single digit per edge in any case, indicating that only the between networks can be found in Supplemental Section S1. 116 order of magnitude of edges in naively computed groupwise 117 average connectomes can be trusted. The combination of The number of significant digits per edge across connec-118 these results with those presented in Figure 1A suggests that 103 tomes (Figure 1B) similarly decreases alongside the decreaswhile specific edge weights are largely affected by instabili-104 ing similarity between comparison groups. While the cross120 ties, macro-scale network organization is stable.

122 inability

124 ing and extending a typical test-retest experiment 26 in which 125 the similarity of samples across sessions were compared to 126 distinct samples in the dataset (Table 1, with additional ex-127 periments and explanation in Supplemental Section S2). The 164 143 of perturbation methods for synthesizing robust and reliable 180 lower-bias estimates of individual connectivity. 144 individual estimates of connectivity, serving as a cost effective and context-agnostic method for dataset augmentation.

While the discriminability of individuals is essential for 147 the identification of individual brain networks, it is similarly 148 reliant on network similarity across equivalent acquisitions 149 (Hypothesis 2). In this case, connectomes were grouped 150 based upon session, rather than subject, and the ability to distinguish one session from another was computed within-152 individual and aggregated. Both the unperturbed and pipeline 153 perturbation settings perfectly preserved differences between 154 cross-sectional sessions with a score of 1.0 (p < 0.005; optimal score: 0.5; chance: 0.5), indicating a dominant session-

156 dependent signal for all individuals despite no intended bio-157 logical differences. However, while still significant relative Perturbations Increase Biologically-Driven Discrim- 158 to chance (score: 0.85 and 0.88; p < 0.005 for both), sparse 159 perturbations lead to significantly lower discriminability of We assessed the reproducibility of the dataset through mimick- 160 the dataset (p < 0.005 for all). This reduction of the differ-161 ence between sessions suggests that the added variance due 162 to perturbations reduces the relative impact of non-biological 163 acquisition-dependent bias inherent in the brain graphs.

Though the previous sets of experiments inextricably eval-128 ability to discriminate connectomes across subjects (Hypothe- 165 uate the interaction between data acquisition and the pipelines, 129 sis 1) is an essential prerequisite for the application of brain 166 the use of subsampling allowed for characterizing the discrim-130 imaging towards identifying individual differences 18. In test- 167 inability of networks sampled from within a single acquisition ing hypothesis 1, we observe that the dataset is discriminable 168 (Hypothesis 3). While this experiment could not be evaluated using reference executions, the executions performed with a score of 0.64 and 0.65 (p < 0.001; optimal score: 1.0; 169 using reference executions, the executions performed with 193 chance: 0.04) for the Deterministic and Probabilistic pipelines, 170 pipeline perturbations showed near perfect discrimination be-134 respectively, in the absence of MCA. However, we can see 171 tween subsamples, with scores of 0.99 and 1.0 (p < 0.005; 195 that inducing instabilities through MCA improves the discrim- 172 optimal: 0.5; chance: 0.5). Given that there is no variability inability of the dataset to over 0.75 in each case (p < 0.001 in data acquisition or preprocessing that contributes to this 197 for all), significantly higher than without instrumentation 174 reliable identification of scans, the separability observed in $_{138}$ (p < 0.005 for all). The definition of the discriminability 175 this experiment may only be due to instability or bias inher-199 statistic is such that if all samples derived from the a given 176 ent to the pipelines. The high variability introduced through 140 session were near replicates of one another the score would be 177 input perturbations considerably lowered the discriminabilunchanging despite the increased sample size 26 . The resulting 178 ity towards chance (score: 0.71 and 0.61; p < 0.005 for all), 142 difference in discriminability impactfully suggests the utility 179 further supporting this as an effective method for obtaining

> Across all cases, the induced perturbations showed an 182 amplification of meaningful biological signal alongside a re-183 duction of off-target signal. This result appears strikingly like ¹⁸⁴ a manifestation of the well-known bias-variance tradeoff³² in machine learning, a concept which observes a decrease in 186 bias as variance is favoured by a model. In particular, this 187 highlights that numerical perturbations can be used to not 188 only evaluate the stability of pipelines, but that the induced 189 variance may be leveraged for the interpretation as a robust 190 distributions of possible results.

Table 1. The impact of instabilities as evaluated through the discriminability of the dataset based on individual (or subject) differences, session, and subsample. The performance is reported as mean discriminability. While a perfectly discriminable dataset would be represented by a score of 1.0, the chance performance, indicating minimal discriminability, is 1/the number of classes. H_3 could not be tested using the reference executions due to too few possible comparisons. The alternative hypothesis, indicating significant discrimination, was accepted for all experiments, with p < 0.005.

			Reference Execution		Perturbed Pipeline		Perturbed Inputs	
Comparison	Chance	Target	Det.	Prob.	Det.	Prob.	Det.	Prob.
H ₁ : Across Subjects	0.04	1.0	0.64	0.65	0.82	0.82	0.77	0.75
<i>H</i> ₂ : Across Sessions	0.5	0.5	1.00	1.00	1.00	1.00	0.88	0.85
<i>H</i> ₃ : Across Subsamples	0.5	0.5			0.99	1.00	0.71	0.61

Distributions of Graph Statistics Were Reliable, But 216 the feature-moments were stable with more than 10 signifi-192 Individual Statistics Were Not

193 Exploring the stability of topological features of connectomes 194 is relevant for typical analyses, as low dimensional features are 195 often more suitable than full connectomes for many analytical 196 methods in practice¹¹. A separate subset of the NKIRS dataset 197 was randomly selected to contain a single non-subsampled ses-198 sion for 100 individuals ($100 \times 1 \times 1$) using the pipelines and 199 instrumentation methods to generate connectomes as above. 200 Connectomes were generated 20 times each, resulting in a 201 dataset which also contained 4,200 connectomes with the 202 MCA simulations serving as the only source of repeated mea-203 surements.

The stability of several commonly-used multivariate graph features¹⁰ was explored in Figure 2. The cumulative den-206 sity of the features was computed within individuals and the 207 mean density and associated standard error were computed 208 for across individuals (Figures 2A and 2B). There was no sig-209 nificant difference between the distributions for each feature 210 across the two perturbation settings, suggesting that the topo-211 logical features summarized by these multivariate features are 212 robust across both perturbation modes.

215 (Figures 2C and 2D). In the face of pipeline perturbations, 240 ure 3).

217 cant digits with the exception of edge weight when using the 218 deterministic pipeline, though the probabilistic pipeline was 219 more stable for all comparisons (p < 0.0001; exploratory). 220 In stark contrast, input perturbations led to highly unstable 221 feature-moments (Figure 2D), such that none contained more 222 than 5 significant digits of information and several contained 223 less than a single significant digit, indicating a complete lack 224 of reliability. This dramatic degradation in stability for in-225 dividual measures strongly suggests that these features may 226 be unreliable as individual biomarkers when derived from a 227 single pipeline evaluation, though their reliability may be in-228 creased when studying their distributions across perturbations. 229 A similar analysis was performed for univariate statistics and 230 can be found in Supplemental Section S3.

231 Uncertainty in Brain-Phenotype Relationships

232 While the variability of connectomes and their features was 233 summarized above, networks are commonly used as inputs to 234 machine learning models tasked with learning brain-phenotype 235 relationships 18. To explore the stability of these analyses, we 236 modelled the relationship between high- or low- Body Mass 237 Index (BMI) groups and brain connectivity 12,13, using stan-In addition to the comparison of distributions, the stabil-238 dard dimensionality reduction and classification tools, and 214 ity of the first 5 moments of these features was evaluated 239 compared this to reference and random performance (Fig-

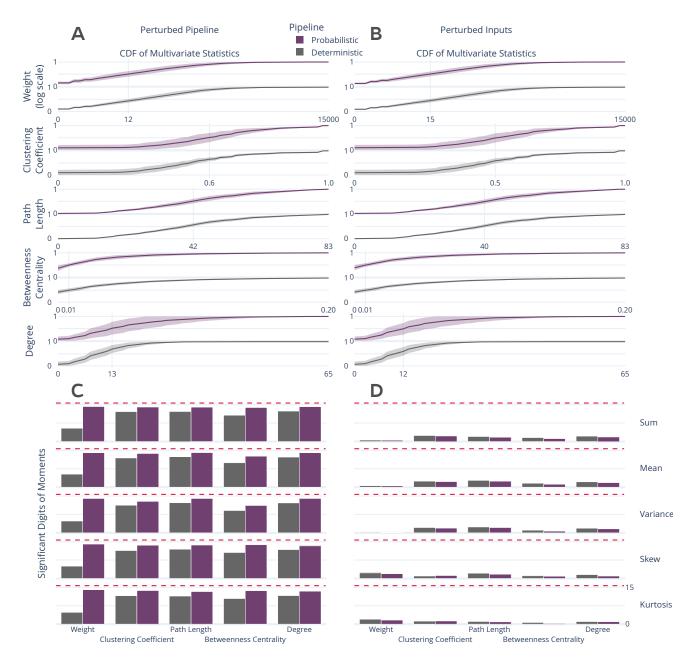


Figure 2. Distribution and stability assessment of multivariate graph statistics. (**A**, **B**) The cumulative distribution functions of multivariate statistics across all subjects and perturbation settings. There was no significant difference between the distributions in A and B. (**C**, **D**) The number of significant digits in the first 5 five moments of each statistic across perturbations. The dashed red line refers to the maximum possible number of significant digits.

The analysis was perturbed through distinct samplings of $_{246}$ performance on the reference dataset. This large variability the dataset across both pipelines and perturbation methods. $_{247}$ illustrates a previously uncharacterized margin of uncertainty The accuracy and F1 score for the perturbed models varied $_{248}$ in the modelling of this relationship, and limits confidence in from $_{244}$ from $_{0.520} - _{0.716}$ and $_{0.510} - _{0.725}$, respectively, rang- $_{249}$ reported accuracy scores on singly processed datasets. The ing from at or below random performance to outperforming $_{250}$ portion of explained variance in these samples ranged from

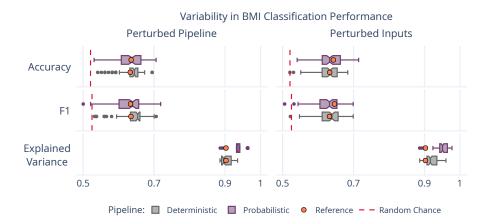


Figure 3. Variability in BMI classification across the sampling of an MCA-perturbed dataset. The dashed red lines indicate random-chance performance, and the orange dots show the performance using the reference executions.

251 88.6% — 97.8%, similar to the reference, suggesting that the 275 led to considerable variability in derived brain graphs. Across 252 range in performance was not due to a gain or loss of mean- 276 all analyses the stability of results ranged from nearly per-253 ingful signal, but rather the reduction of bias towards specific 277 feetly trustworthy (i.e. no variation) to completely unreliable 254 outcome. Importantly, this finding does not suggest that mod- 278 (i.e. containing no trustworthy information). Given that the 255 elling brain-phenotype relationships is not possible, but rather 279 magnitude of introduced numerical noise is to be expected 256 it sheds light on impactful uncertainty that must be accounted 280 in typical settings, this finding has potentially significant im-257 for in this process, and supports the use of ensemble modeling 281 plications for inferences in brain imaging as it is currently 258 techniques. 282 performed. In particular, this bounds the success of studying One distinction between the results presented here and the 283 individual differences, a central objective in brain imaging 18, 260 previous is that while networks derived from dense perturba- 284 given that the quality of relationships between phenotypic 261 tions had been shown to exhibit less dramatic instabilities in 285 data and brain networks will be limited by the stability of the 262 general, the results here show similar variability in classifica- 266 connectomes themselves. This issue was accentuated through 263 tion performance across the two methods. This consistency 287 the crucial finding that individually derived network features 284 suggests that the desired method of pipeline instrumentation 288 were unreliable despite there being no significant difference 285 may vary across experiments. While sparse perturbations 289 in their aggregated distributions. This finding is not damn-266 result in considerably more variability in networks directly, 290 ing for the study of brain networks as a whole, but rather is 267 the two techniques capture similar variability when relating 291 strong support for the aggregation of networks, either across 268 networks to this phenotypic variable. Given the dramatic 292 perturbations for an individual or across groups, over the use 269 reduction in computational overhead, a sparse instrumenta- 293 of individual estimates. 270 tion may be preferred when processing datasets for eventual 294 Underestimated False Positive Rates While the instabil-271 application in modelling brain-phenotype relationships.

272 Discussion

²⁷³ The perturbation of structural connectome estimation pipelines ²⁹⁷ context of machine learning, this limitation extends to classi-²⁷⁴ with small amounts of noise, on the order of machine error, ²⁹⁸ cal hypothesis testing, as well. Though performing individual

295 ity of brain networks was used here to demonstrate the lim296 itations of modelling brain-phenotype relationships in the
297 context of machine learning, this limitation extends to classi-

299 comparisons in a hypothesis testing framework will be accom- 336 exploring the effect of data collection paradigm or study lon-2000 panied by reported false positive rates, the accuracy of these 307 gitudinal progressions of development or disease, it could be 301 rates is critically dependent upon the reliability of the samples 338 used in conjunction with these efforts to increase the reliabil-302 used. In reality, the true false positive rate for a test would be 339 ity of each distinct sample within a dataset. In contexts where acombination of the reported confidence and the underlying 340 repeated measurements are collected to increase the fidelity of 304 variability in the results, a typically unknown quantity.

measure context, such as that afforded here through MCA, it 343 data collection. This technique also opens the door for the 307 is impossible to empirically estimate the reliability of samples. 344 characterization of reliability across axes which have been 308 This means that the reliability of accepted hypotheses is also 345 traditionally inaccessible. For instance, in the absence of a 309 unknown, regardless of the reported false positive rate. In 346 realistic noise model or simulation technique similar to MCA, 310 fact, it is a virtual certainty that the true false positive rate 347 the evaluation of network stability across data subsampling 311 for a given hypothesis exceeds the reported value simply as 348 would not have been possible. 312 a result of numerical instabilities. This uncertainty inherent 313 to derived data is compounded with traditional arguments 314 limiting the trustworthiness of claims³³, and hampers the 315 ability of researchers to evaluate the quality of results. The 316 accompaniment of brain imaging experiments with direct 317 evaluations of their stability, as was done here, would allow 318 researchers to simultaneously improve the numerical stability 319 of their analyses and accurately gauge confidence in them. 320 The induced variability in derived brain networks may be 321 leveraged to estimate aggregate connectomes with lower bias 322 than any single independent observation, leading to learned 323 relationships that are more generalizable and ultimately more 324 useful.

ability in brain imaging has historically relied upon the ex- 363 comes in brain imaging has been clearly demonstrated. The 327 pensive collection of repeated measurements choreographed 364 approach taken in these studies complement one another and by massive cross-institutional consortia^{34,35}. The finding that see explore instability at the opposite ends of the spectrum, with get perturbing experiments using MCA both increased the relia- 366 human variability in the construction of an analysis workflow 330 bility of the dataset and decreased off-target differences across 367 on one end and the unavoidable error implicit in the digital 332 Given that MCA is data-agnostic, this technique could be used 369 to combine these approaches and explore the interaction of 333 effectively in conjunction with, or in lieu of, realistic noise 370 these scientific degrees of freedom with effects from software 334 models to augment existing datasets. While this of course 371 implementations, libraries, and parametric choices. 335 would not replace the need for repeated measurements when 372

341 the dataset, MCA could potentially be employed to increase When performing these experiments outside of a repeated-342 the reliability of the dataset and save millions of dollars on

> 349 Shortcomings and Future Questions Given the complex-350 ity of recompiling complex software libraries, pre-processing was not perturbed in these experiments. Other work has shown 352 that linear registration, a core piece of many elements of preprocessing such as motion correction and alignment, is sensi-354 tive to minor perturbations⁷. It is likely that the instabilities 355 across the entire processing workflow would be compounded 356 with one another, resulting in even greater variability. While 357 the analyses performed in this paper evaluated a single dataset and set of pipelines, extending this work to other modalities and analyses is of interest for future projects.

This paper does not explore methodological flexibility or 361 compare this to numerical instability. Recently, the nearly 325 **Cost-Effective Data Augmentation** The evaluation of reli- 362 boundless space of analysis pipelines and their impact on outacquisitions opens the door for a promising paradigm shift. 368 representation of data on the other. It is of extreme interest

Finally, it is important to state explicitly that the work

presented here does not invalidate analytical pipelines used in brain imaging, but merely sheds light on the fact that many studies are accompanied by an unknown degree of uncertainty due to machine-introduced errors. The presence of unknown error-bars associated with experimental findings limits the impact of results due to increased uncertainty. The desired outcome of this paper is to motivate a shift in scientific computing — both in neuroimaging and more broadly — towards a paradigm which favours the explicit evaluation of the trust-worthiness of claims alongside the claims themselves.

Methods

384 Dataset

The Nathan Kline Institute Rockland Sample (NKI-RS)²⁹ dataset contains high-fidelity imaging and phenotypic data from over 1,000 individuals spread across the lifespan. A subset of this dataset was chosen for each experiment to both match sample sizes presented in the original analyses and to minimize the computational burden of performing MCA. The selected subset comprises 100 individuals ranging in age from selected subset comprises 100 individuals ranging in age from 6 – 79 with a mean of 36.8 (original: 6 – 81, mean 37.8), 60% female (original: 60%), with 52% having a BMI over 25 (original: 54%).

Each selected individual had at least a single session of both structural T1-weighted (MPRAGE) and diffusion-weighted (DWI) MR imaging data. DWI data was acquired with 137 diffusion directions; more information regarding the acquisition of this dataset can be found in the NKI-RS data release²⁹.

In addition to the 100 sessions mentioned above, 25 individuals had a second session to be used in a test-retest analysis. Two additional copies of the data for these individuals were generated, including only the odd or even diffusion directions to (64 + 9 B0 volumes = 73 in either case). This allowed for an extra level of stability evaluation to be performed between the to levels of MCA and session-level variation.

In total, the dataset is composed of 100 downsampled sessions of data originating from 50 acquisitions and 25 in-dividuals for in depth stability analysis, and an additional 100 sessions of full-resolution data from 100 individuals for subsequent analyses.

413 Processing

414 The dataset was preprocessed using a standard FSL³⁶ work-415 flow consisting of eddy-current correction and alignment. The 416 MNI152 atlas³⁷ was aligned to each session of data, and the re-417 sulting transformation was applied to the DKT parcellation³⁸. 418 Downsampling the diffusion data took place after preprocess419 ing was performed on full-resolution sessions, ensuring that 453 while performing MCA on the output of an operation high-420 an additional confound was not introduced in this process 454 lights round-off errors that may be introduced. The former is 421 when comparing between downsampled sessions. The pre- 455 referred to as Precision Bounding (PB) and the latter is called 422 processing described here was performed once without MCA, 456 Random Rounding (RR). 423 and thus is not being evaluated.

425 data using two canonical pipelines from Dipy³⁰: deterministic 459 distribution of these results can then lead to insights on the and probabilistic. In the deterministic pipeline, a constant 460 stability of the instrumented tools or functions. To this end, 427 solid angle model was used to estimate tensors at each voxel 461 a complete software stack was instrumented with MCA and and streamlines were then generated using the EuDX algo- 462 is made available on GitHub at https://github.com/ 429 rithm³¹. In the probabilistic pipeline, a constrained spherical 463 gkiar/fuzzy. 430 deconvolution model was fit at each voxel and streamlines 431 were generated by iteratively sampling the resulting fiber ori-432 entation distributions. In both cases tracking occurred with 8 433 seeds per 3D voxel and edges were added to the graph based 434 on the location of terminal nodes with weight determined by 435 fiber count.

437 yses. Fixing this random state led to entirely deterministic 438 repeated-evaluations of the tools, and allowed for explicit at-439 tribution of observed variability to Monte Carlo simulations 440 rather than internal state of the algorithm.

441 Perturbations

442 All connectomes were generated with one reference execu-443 tion where no perturbation was introduced in the processing. 444 For all other executions, all floating point operations were 445 instrumented with Monte Carlo Arithmetic (MCA)⁸ through 446 Verificarlo⁹. MCA simulates the distribution of errors im-447 plicit to all instrumented floating point operations (flop). This 448 rounding is performed on a value x at precision t by:

$$inexact(x) = x + 2^{e_x - t}\xi \tag{1}$$

Using MCA, the execution of a pipeline may be performed Structural connectomes were generated from preprocessed 458 many times to produce a distribution of results. Studying the

Both the RR and PB variants of MCA were used indepen-465 dently for all experiments. As was presented in⁴, both the 466 degree of instrumentation (i.e. number of affected libraries) and the perturbation mode have an effect on the distribution 468 of observed results. For this work, the RR-MCA was ap-469 plied across the bulk of the relevant libraries and is referred The random state of both pipelines was fixed for all anal- 470 to as Pipeline Perturbation. In this case the bulk of numerical ⁴⁷¹ operations were affected by MCA.

> Conversely, the case in which PB-MCA was applied across 473 the operations in a small subset of libraries is here referred 474 to as Input Perturbation. In this case, the inputs to operations 475 within the instrumented libraries (namely, Python and Cython) 476 were perturbed, resulting in less frequent, data-centric pertur-477 bations. Alongside the stated theoretical differences, Input ⁴⁷⁸ Perturbation is considerably less computationally expensive 479 than Pipeline Perturbation.

> All perturbations targeted the least-significant-bit for all data (t = 24 and t = 53 in float32 and float64, respectively⁹). 482 Simulations were performed 20 times for each pipeline execu-483 tion. A detailed motivation for the number of simulations can 484 be found in³⁹.

where e_x is the exponent value of x and ξ is a uniform ran- 485 **Evaluation**

450 dom variable in the range $(-\frac{1}{2}, \frac{1}{2})$. MCA can be introduced in 486 The magnitude and importance of instabilities in pipelines 451 two places for each flop: before or after evaluation. Perform- 487 can be considered at a number of analytical levels, namely: 452 ing MCA on the inputs of an operation limits its precision, 488 the induced variability of derivatives directly, the resulting

490 ultimate change in analyses or findings. We explore the na- 524 ing a direct measure of the tool-introduced variability across 491 ture and severity of instabilities through each of these lenses. 525 perturbations. A distribution was formed by aggregating these 492 Unless otherwise stated, all p-values were computed using 526 individual results. 493 Wilcoxon signed-rank tests. To ensure avoid biasing these 494 statistics in this unique repeated-measures context, tests were 495 performed across sets of independent obversations and then 496 the results were aggregated in all cases.

497 Direct Evaluation of the Graphs

499 rectly through both a direct variance quantification and a 533 ranked distribution of distances between observations belong-500 comparison to other sources of variance such as individual- 504 ing to a defined set of classes. The discriminability statistic is and session-level differences.

502 Quantification of Variability Graphs, in the form of adja-503 cency matrices, were compared to one another using three 504 metrics: normalized percent deviation, Pearson correlation, 505 and edgewise significant digits. The normalized percent deviso ation measure, defined in⁴, scales the norm of the difference 507 between a simulated graph and the reference execution (that without intentional perturbation) with respect to the norm of 509 the reference graph. The purpose of this comparison is to 510 provide insight on the scale of differences in observed graphs ⁵¹¹ relative to the original signal intensity. A Pearson correlation coefficient⁴⁰ was computed in complement to normalized per-513 cent deviation to identify the consistency of structure and not 514 just intensity between observed graphs.

Finally, the estimated number of significant digits, s', for 516 each edge in the graph is calculated as:

$$s' = -log_{10} \frac{\sigma}{|\mu|} \tag{2}$$

where μ and σ are the mean and unbiased estimator of 518 standard deviation across graphs, respectively. The upper 519 bound on significant digits is 15.7 for 64-bit floating point 520 data.

The percent deviation, correlation, and number of signifi-522 cant digits were each calculated within a single session of data, 556 H_{A1} : Individuals are distinct from one another

489 downstream impact on summary statistics or features, or the 523 thereby removing any subject- and session-effects and provid-

527 Class-based Variability Evaluation To gain a concrete un-528 derstanding of the significance of observed variations we explore the separability of our results with respect to understood 530 sources of variability, such as subject-, session-, and pipeline-1 level effects. This can be probed through Discriminability 26, 498 The differences between simulated graphs was measured di- 532 a technique similar to ICC²⁴ which relies on the mean of a 535 formalized as follows:

$$Disc. = Pr(\|g_{ij} - g_{ij'}\| \le \|g_{ij} - g_{i'j'}\|)$$
(3)

where g_{ij} is a graph belonging to class i that was measured at observation j, where $i \neq i'$ and $j \neq j'$.

Discriminability can then be read as the probability that an 539 observation belonging to a given class will be more similar to 540 other observations within that class than observations of a different class. It is a measure of reproducibility, and is discussed 542 in detail in²⁶. This definition allows for the exploration of 543 deviations across arbitrarily defined classes which in practice 544 can be any of those listed above. We combine this statistic 545 with permutation testing to test hypotheses on whether differ-546 ences between classes are statistically significant in each of 547 these settings.

With this in mind, three hypotheses were defined. For 549 each setting, we state the alternate hypotheses, the variable(s) (2) 550 which were used to determine class membership, and the 551 remaining variables which may be sampled when obtaining 552 multiple observations. Each hypothesis was tested indepen-553 dently for each pipeline and perturbation mode, and in every 554 case where it was possible the hypotheses were tested using 555 the reference executions alongside using MCA.

Class definition: Subject ID

557

562

564

Comparisons: Session (1 subsample), Subsample (1

session), MCA (1 subsample, 1 session) 559

 H_{A2} : Sessions within an individual are distinct

Class definition: Session ID | Subject ID

Comparisons: **Subsample**, MCA (1 subsample)

563 H_{A3} : Subsamples are distinct

Class definition: Subsample | Subject ID, Session ID

Comparisons: MCA 565

As a result, we tested 3 hypotheses across 6 MCA ex-567 periments and 3 reference experiments on 2 pipelines and 2 568 perturbation modes, resulting in a total of 30 distinct tests. 569 While results from all tests can be found within Supplemental 570 Section S2, only the bolded comparisons in the list above have 571 been presented in the main body of this article.

572 Evaluating Graph-Theoretical Metrics

574 it is common practice to summarize them with structural mea- 610 tions. We performed the modeling task with a single sampled 575 sures, which can then be used as lower-dimensional proxies 611 connectome per individual and repeated this sampling and ₅₇₆ of connectivity in so-called graph-theoretical studies¹¹. We ₆₁₂ modelling 20 times. We report the model performance for explored the stability of several commonly-used univariate 613 each sampling of the dataset and summarize its variance. 578 (graphwise) and multivariate (nodewise or edgewise) features. 579 The features computed and subsequent methods for compari-580 son in this section were selected to closely match those com-581 puted in¹⁰.

582 Univariate Differences For each univariate statistic (edge 618 reduced the dimensionality of the connectomes through prin-583 count, mean clustering coefficient, global efficiency, modu- 619 cipal component analysis (PCA), and provided the first N-584 larity of the largest connected component, assortativity, and 620 components to a logistic regression classifier for predicting mean path length) a distribution of values across all perturba- 621 BMI class membership, similar to methods shown in 12,13. 586 tions within subjects was observed. A Z-score was computed 622 The number of components was selected as the minimum set 587 for each sample with respect to the distribution of feature 623 which explained > 90% of the variance when averaged across 588 values within an individual, and the proportion of "classically 624 the training set for each fold within the cross validation of significant" Z-scores, i.e. corresponding to p < 0.05, was 625 the original graphs; this resulted in a feature of 20 compo-590 reported and aggregated across all subjects. The number of 626 nents. We trained the model using k-fold cross validation,

592 single subject were calculated and aggregated.

593 Multivariate Differences In the case of both nodewise (de-594 gree distribution, clustering coefficient, betweenness central-595 ity) and edgewise (weight distribution, connection length) fea-596 tures, the cumulative density functions of their distributions 597 were evaluated over a fixed range and subsequently aggre-598 gated across individuals. The number of significant digits 599 for each moment of these distributions (sum, mean, variance, 600 skew, and kurtosis) were calculated across observations within 601 a sample and aggregated.

602 Evaluating A Brain-Phenotype Analysis

603 Though each of the above approaches explores the instabil-604 ity of derived connectomes and their features, many modern studies employ modeling or machine-learning approaches, for 606 instance to learn brain-phenotype relationships or identify dif-607 ferences across groups. We carried out one such study and ex-608 plored the instability of its results with respect to the upstream 573 While connectomes may be used directly for some analyses, 609 variability of connectomes characterized in the previous sec-

614 **BMI Classification** Structural changes have been linked to obesity in adolescents and adults⁴¹. We classified normal-616 weight and overweight individuals from their structural net- $_{617}$ works (using for overweight a cutoff of BMI $> 25^{13}$). We significant digits contained within an estimate derived from a 627 with k = 2, 5, 10, and N (equivalent to leave-one-out; LOO).

628 Data & Code Provenance

of Reliability and Reproducibility (http://fcon_1000.666 ship with Health Canada, for the Canadian Open Neuroscience 631 projects.nitrc.org/indi/enhanced/), including 667 Platform initiative. 632 both the imaging data as well as phenotypic data which may be obtained upon submission and compliance with a Data Us- 668 References age Agreement. The connectomes generated through simula- 669 [1] R. Botvinik-Nezer, F. Holzmeister, C. F. Camerer, A. Dreber, J. Huber, $_{635}$ tions have been bundled and stored permanently (https:// 670 636 doi.org/10.5281/zenodo.4041549), and are made 637 available through The Canadian Open Neuroscience Platform 638 (https://portal.conp.ca/search, search term "Kiar,"). All software developed for processing or evaluation is 675 640 publicly available on GitHub at https://github.com/ 676 $_{\rm 641}$ gkpapers/2020ImpactOfInstability. Experiments $^{\rm 677}$ $_{642}$ were launched using Boutiques 42 and Clowdr 43 in Compute 643 Canada's HPC cluster environment. MCA instrumentation 680 644 was achieved through Verificarlo⁹ available on Github at 681 [4] 645 https://github.com/verificarlo/verificarl89?

648 Author Contributions

649 GK was responsible for the experimental design, data pro-650 cessing, analysis, interpretation, and the majority of writing. 689 651 All authors contributed to the revision of the manuscript. YC, 690 652 POC, and EP were responsible for MCA tool development and 653 software testing. AR, GV, and BM contributed to experimen-654 tal design and interpretation. TG contributed to experimental 655 design, analysis, and interpretation. TG and ACE were re- 695 656 sponsible for supervising and supporting all contributions 696 made by GK. The authors declare no competing interests for 697 [8] 658 this work. Correspondence and requests for materials should 659 be addressed to Tristan Glatard at tristan.glatard@ 660 concordia.ca.

646 A set of MCA instrumented software containers is available

647 on Github at https://github.com/gkiar/fuzzy.

661 Acknowledgments

662 This research was financially supported by the Natural Sci-663 ences and Engineering Research Council of Canada (NSERC) 705

664 (award no. CGSD3-519497-2018). This work was also sup-The unprocessed dataset is available through The Consortium 665 ported in part by funding provided by Brain Canada, in partner-

686

687

702

- M. Johannesson, M. Kirchler, R. Iwanir, J. A. Mumford, R. A. Adcock et al., "Variability in the analysis of a single neuroimaging dataset by many teams," Nature, pp. 1-7, 2020.
- C. M. Bennett, M. B. Miller, and G. L. Wolford, "Neural correlates of interspecies perspective taking in the post-mortem Atlantic salmon: An argument for multiple comparisons correction," Neuroimage, vol. 47, no. Suppl 1, p. S125, 2009.
- A. Eklund, T. E. Nichols, and H. Knutsson, "Cluster failure: Why fMRI inferences for spatial extent have inflated false-positive rates," Proceedings of the national academy of sciences, vol. 113, no. 28, pp. 7900-7905, 2016.
- G. Kiar, P. de Oliveira Castro, P. Rioux, E. Petit, S. T. Brown, A. C. Evans, and T. Glatard, "Comparing perturbation models for evaluating stability of neuroimaging pipelines," The International Journal of High Performance Computing Applications, 2020.
- A. Salari, G. Kiar, L. Lewis, A. C. Evans, and T. Glatard, "File-based localization of numerical perturbations in data analysis pipelines," arXiv preprint arXiv:2006.04684, 2020.
- L. B. Lewis, C. Y. Lepage, N. Khalili-Mahani, M. Omidyeganeh, S. Jeon, P. Bermudez, A. Zijdenbos, R. Vincent, R. Adalat, and A. C. Evans, "Robustness and reliability of cortical surface reconstruction in CIVET and FreeSurfer," Annual Meeting of the Organization for Human Brain Mapping, 2017.
- T. Glatard, L. B. Lewis, R. Ferreira da Silva, R. Adalat, N. Beck, C. Lepage, P. Rioux, M.-E. Rousseau, T. Sherif, E. Deelman, N. Khalili-Mahani, and A. C. Evans, "Reproducibility of neuroimaging analyses across operating systems," Front. Neuroinform., vol. 9, p. 12, Apr. 2015.
- D. S. Parker, Monte Carlo Arithmetic: exploiting randomness in floatingpoint arithmetic. University of California (Los Angeles). Computer Science Department, 1997.
- C. Denis, P. de Oliveira Castro, and E. Petit, "Verificarlo: Checking floating point accuracy through monte carlo arithmetic," 2016 IEEE 23nd Symposium on Computer Arithmetic (ARITH), 2016.
- R. F. Betzel, A. Griffa, P. Hagmann, and B. Mišić, "Distance-dependent consensus thresholds for generating group-representative structural brain networks," Network neuroscience, vol. 3, no. 2, pp. 475-496, 2019.

- M. Rubinov and O. Sporns, "Complex network measures of brain con-751 [27] G. Kiar, E. Bridgeford, W. G. Roncal, V. Chandrashekhar, and othnectivity: uses and interpretations," Neuroimage, vol. 52, no. 3, pp. 752 707 1059-1069, Sep. 2010.
- connectivity of people with obesity and prediction of body mass index 755 [29] using connectivity," PLoS One, vol. 10, no. 11, p. e0141376, Nov. 2015. 756 711
- ₇₁₂ [13] A. Gupta, E. A. Mayer, C. P. Sanmiguel, J. D. Van Horn, D. Woodworth, 757 B. M. Ellingson, C. Fling, A. Love, K. Tillisch, and J. S. Labus, "Pat-758 [30] 713 terns of brain structural connectivity differentiate normal weight from 759 714 overweight subjects," Neuroimage Clin, vol. 7, pp. 506-517, Jan. 2015. 760
- T. E. Behrens and O. Sporns, "Human connectomics," Current opinion 761 in neurobiology, vol. 22, no. 1, pp. 144-153, 2012.
- ₇₁₈ [15] M. Xia, Q. Lin, Y. Bi, and Y. He, "Connectomic insights into topologi- 763 cally centralized network edges and relevant motifs in the human brain," 764 719 Frontiers in human neuroscience, vol. 10, p. 158, 2016. 720
- ₇₂₁ [16] J. L. Morgan and J. W. Lichtman, "Why not connectomics?" Nature 766 methods, vol. 10, no. 6, p. 494, 2013.
- ₇₂₃ [17] M. P. Van den Heuvel, E. T. Bullmore, and O. Sporns, "Comparative 768 [33] connectomics," Trends in cognitive sciences, vol. 20, no. 5, pp. 345–361, 769 724
- ₇₂₆ [18] J. Dubois and R. Adolphs, "Building a science of individual differences 771 from fMRI," Trends Cogn. Sci., vol. 20, no. 6, pp. 425-443, Jun. 2016. 772
- ₇₂₈ [19] A. Fornito and E. T. Bullmore, "Connectomics: a new paradigm for 773 [35] understanding brain disease," European Neuropsychopharmacology, 774 vol. 25, no. 5, pp. 733-748, 2015. 730
- G. Deco and M. L. Kringelbach, "Great expectations: using whole- 776 brain computational connectomics for understanding neuropsychiatric 777 disorders," Neuron, vol. 84, no. 5, pp. 892-905, 2014. 733
- T. Xie and Y. He, "Mapping the alzheimer's brain with connectomics," 779 Frontiers in psychiatry, vol. 2, p. 77, 2012.
- ₇₃₆ [22] M. Filippi, M. P. van den Heuvel, A. Fornito, Y. He, H. E. H. Pol, 781 [37] F. Agosta, G. Comi, and M. A. Rocca, "Assessment of system dys-782 function in the brain through mri-based connectomics," The Lancet 783 Neurology, vol. 12, no. 12, pp. 1189-1199, 2013.
- ₇₄₀ [23] M. P. Van Den Heuvel and A. Fornito, "Brain networks in schizophrenia," 785 [38] Neuropsychology review, vol. 24, no. 1, pp. 32-48, 2014. 741
- 742 [24] J. J. Bartko, "The intraclass correlation coefficient as a measure of 787 reliability," Psychol. Rep., vol. 19, no. 1, pp. 3-11, Aug. 1966.
- ₇₄₄ [25] A. M. Brandmaier, E. Wenger, N. C. Bodammer, S. Kühn, N. Raz, 789 and U. Lindenberger, "Assessing reliability in neuroimaging research 790 [40] 745 through intra-class effect decomposition (ICED)," Elife, vol. 7, Jul. 2018. 791
- ₇₄₇ [26] E. W. Bridgeford, S. Wang, Z. Yang, Z. Wang, T. Xu, C. Craddock, 792 J. Dey, G. Kiar, W. Gray-Roncal, C. Coulantoni et al., "Eliminating 793 748 accidental deviations to minimize generalization error: applications in 794 [41] C. A. Raji, A. J. Ho, N. N. Parikshak, J. T. Becker, O. L. Lopez, L. H. 749 connectomics and genomics," bioRxiv, p. 802629, 2020. 750

- ers, "A High-Throughput pipeline identifies robust connectomes but troublesome variability," bioRxiv, 2018.
- B.-Y. Park, J. Seo, J. Yi, and H. Park, "Structural and functional brain 754 [28] M. Baker, "1,500 scientists lift the lid on reproducibility," Nature, 2016.
 - K. B. Nooner, S. J. Colcombe, R. H. Tobe, M. Mennes et al., "The NKI-Rockland sample: A model for accelerating the pace of discovery science in psychiatry," Front. Neurosci., vol. 6, p. 152, Oct. 2012.
 - E. Garyfallidis, M. Brett, B. Amirbekian, A. Rokem, S. van der Walt, M. Descoteaux, I. Nimmo-Smith, and Dipy Contributors, "Dipy, a library for the analysis of diffusion MRI data," Front. Neuroinform., vol. 8, p. 8, Feb. 2014.
 - E. Garyfallidis, M. Brett, M. M. Correia, G. B. Williams, and I. Nimmo-Smith, "QuickBundles, a method for tractography simplification," Front. Neurosci., vol. 6, p. 175, Dec. 2012.
 - 765 [32] S. Geman, E. Bienenstock, and R. Doursat, "Neural networks and the bias/variance dilemma," Neural computation, vol. 4, no. 1, pp. 1-58,
 - J. P. Ioannidis, "Why most published research findings are false," PLoS medicine, vol. 2, no. 8, p. e124, 2005.
 - ₇₇₀ **[34]** D. C. Van Essen, S. M. Smith, D. M. Barch, T. E. Behrens, E. Yacoub, K. Ugurbil, W.-M. H. Consortium et al., "The WU-Minn human connectome project: an overview," Neuroimage, vol. 80, pp. 62-79, 2013.
 - X.-N. Zuo, J. S. Anderson, P. Bellec, R. M. Birn, B. B. Biswal, J. Blautzik, J. C. Breitner, R. L. Buckner, V. D. Calhoun, F. X. Castellanos et al., "An open science resource for establishing reliability and reproducibility in functional connectomics," Scientific data, vol. 1, no. 1, pp. 1-13, 2014.
 - M. Jenkinson, C. F. Beckmann, T. E. J. Behrens, M. W. Woolrich, and S. M. Smith, "FSL," Neuroimage, vol. 62, no. 2, pp. 782-790, Aug. 2012.
 - J. L. Lancaster, D. Tordesillas-Gutiérrez, M. Martinez, F. Salinas, A. Evans, K. Zilles, J. C. Mazziotta, and P. T. Fox, "Bias between mni and talairach coordinates analyzed using the icbm-152 brain template," Human brain mapping, vol. 28, no. 11, pp. 1194-1205, 2007.
 - A. Klein and J. Tourville, "101 labeled brain images and a consistent human cortical labeling protocol," Front. Neurosci., vol. 6, p. 171, Dec. 2012
 - D. Sohier, P. De Oliveira Castro, F. Févotte, B. Lathuilière, E. Petit, and O. Jamond, "Confidence intervals for stochastic arithmetic," Jul. 2018.
 - J. Benesty, J. Chen, Y. Huang, and I. Cohen, "Pearson correlation coefficient," in Noise Reduction in Speech Processing, I. Cohen, Y. Huang, J. Chen, and J. Benesty, Eds. Berlin, Heidelberg: Springer Berlin Heidelberg, 2009, pp. 1-4.
 - Kuller, X. Hua, A. D. Leow, A. W. Toga, and P. M. Thompson, "Brain

- structure and obesity," *Hum. Brain Mapp.*, vol. 31, no. 3, pp. 353–364,
 Mar. 2010.
- 798 [42] T. Glatard, G. Kiar, T. Aumentado-Armstrong, N. Beck, P. Bellec,
- 799 R. Bernard, A. Bonnet, S. T. Brown, S. Camarasu-Pop, F. Cervenansky,
- 800 S. Das, R. Ferreira da Silva, G. Flandin, P. Girard, K. J. Gorgolewski,
- C. R. G. Guttmann, V. Hayot-Sasson, P.-O. Quirion, P. Rioux, M.-É.
- Rousseau, and A. C. Evans, "Boutiques: a flexible framework to inte-
- $\,$ grate command-line applications in computing platforms," Gigascience,
- vol. 7, no. 5, May 2018.
- $_{\mbox{\scriptsize 805}}$ G. Kiar, S. T. Brown, T. Glatard, and A. C. Evans, "A serverless tool
- 806 for platform agnostic computational experiment management," Front.
- 807 Neuroinform., vol. 13, p. 12, Mar. 2019.
- 808 [44] H. Huang and M. Ding, "Linking functional connectivity and structural
- sog connectivity quantitatively: a comparison of methods," Brain connectiv-
- ity, vol. 6, no. 2, pp. 99–108, 2016.

S1. Graph Correlation

The correlations between observed graphs (Figure S1) across each grouping follow the same trend to as percent deviation, as shown in Figure 1. However, notably different from percent deviation, there is no significant difference in the correlations between pipeline or input instrumentations. By this measure, the probabilistic pipeline is more stable in all cross-MCA and cross-directions except for the combination of input perturbation and cross-MCA (p < 0.0001 for all; exploratory).

The marked lack in drop-off of performance across these settings, inconsistent with the measures show in Figure 1 is due to the nature of the measure and the graphs. Given that structural graphs are sparse and contain considerable numbers of zero-weighted edges, the presence or absense of an edge dominated the correlation measure where it was less impactful for the others. For this reason and others⁴⁴, correlation is not a commonly used measure in the context of structural connectivity.

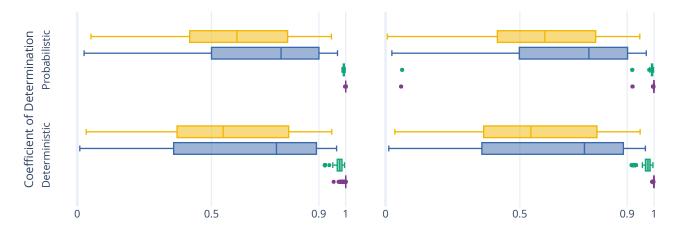


Figure S1. The correlation between perturbed connectomes and their reference.

811

S2. Complete Discriminability Analysis

Table S1. The complete results from the Discriminability analysis, with results reported as mean \pm standard deviation Discriminability. As was the case in the condensed table, the alternative hypothesis, indicating significant separation across groups, was accepted for all experiments, with p < 0.005.

				Reference Execution		Perturbed P	ipeline	Perturbed Inputs		
Exp.	Subj.	Sess.	Samp.	Det.	Prob.	Det.	Prob.	Det.	Prob.	
1.1	All	All	1	0.64 ± 0.00	0.65 ± 0.00	0.82 ± 0.00	0.82 ± 0.00	0.77 ± 0.00	0.75 ± 0.00	
1.2	All	1	All	1.00 ± 0.00	1.00 ± 0.00	1.00 ± 0.00	1.00 ± 0.00	0.93 ± 0.02	0.90 ± 0.02	
1.3	All	1	1			1.00 ± 0.00	1.00 ± 0.00	0.94 ± 0.02	0.90 ± 0.02	
2.4	1	All	All	1.00 ± 0.00	1.00 ± 0.00	1.00 ± 0.00	1.00 ± 0.00	0.88 ± 0.12	0.85 ± 0.12	
2.5	1	All	1			1.00 ± 0.00	1.00 ± 0.00	0.89 ± 0.11	0.84 ± 0.12	
3.6	1	1	All			0.99 ± 0.03	1.00 ± 0.00	0.71 ± 0.07	0.61 ± 0.05	

The complete discriminability analysis includes comparisons across more axes of variability than the condensed version.
The reduction in the main body was such that only axes which would be relevant for a typical analysis were presented. Here,
each of Hypothesis 1, testing the difference across subjects, and 2, testing the difference across sessions, were accompanied
with additional comparisons to those shown in the main body.

Subject Variation Alongside experiment 1.1, that which mimicked a typical test-retest scenario, experiments 1.2 and 1.3 could be considered a test-retest with a handicap, given a single acquisition per individual was compared either across subsamples or simulations, respectively. For this reason, it is unsurprising that the dataset achieved considerably higher discriminability scores.

Session Variation Similar to subject variation, the session variation was also modelled across either both or a single subsample. In both of these cases the performance was similar, and the finding that input perturbation reduced the off-target signal was consistent.

S3. Univariate Graph Statistics

Figure S2 explores the stability of univariate graph-theoretical metrics computed from the perturbed graphs, including modularity, global efficiency, assortativity, average path length, and edge count. When aggregated across individuals and perturbations, the distributions of these statistics (Figures S2A and S22B) showed no significant differences between perturbation methods for either deterministic or probabilistic pipelines.

However, when quantifying the stability of these measures across connectomes derived from a single session of data, the two perturbation methods show considerable differences. The number of significant digits in univariate statistics for Pipeline Perturbation instrumented connectome generation exceeded 11 digits for all measures except modularity, which contained more than 4 significant digits of information (Figure S2C). When detecting outliers from the distributions of observed statistics for a given session, the false positive rate (using a threshold of p = 0.05) was approximately 2% for all statistics with the exception of modularity which again was less stable with an approximately 10% false positive rate. The probabilistic pipeline is significantly more stable than the deterministic pipeline (p < 0.0001; exploratory) for all features except modularity. When similarly evaluating these features from connectomes generated in the input perturbation setting, no statistic was stable with more than 3 significant digits or a false positive rate lower than nearly 6% (Figure S2D). The deterministic pipeline was more

Two notable differences between the two perturbation methods are, first, the uniformity in the stability of the statistics, and second, the dramatic decline in stability of individual statistics in the input perturbation setting despite the consistency in the overall distribution of values. It is unclear at present if the discrepancy between the stability of modularity in the pipeline perturbation context versus the other statistics suggests the implementation of this measure is the source of instability or if it is implicit to the measure itself. The dramatic decline in the stability of features derived from input perturbed graphs despite no difference in their overall distribution both shows that while individual estimates may be unstable the comparison between aggregates or groups may be considered much more reliable; this finding is consistent with that presented for multivariate statistics.

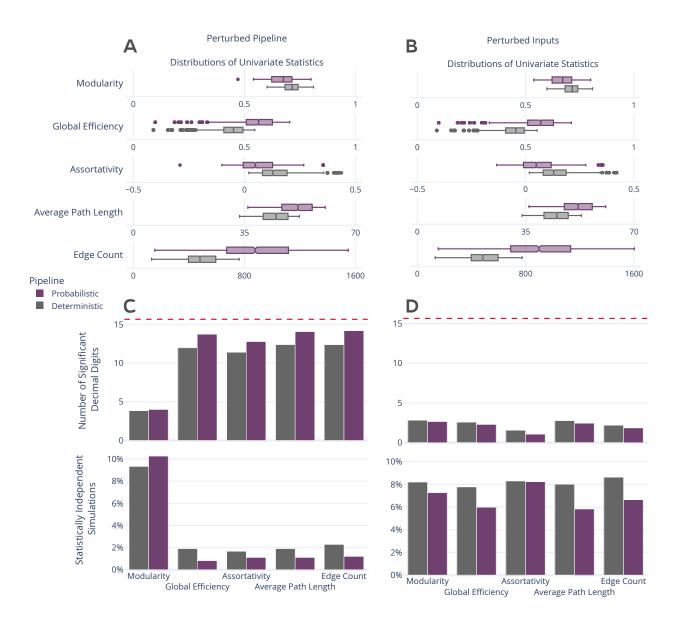


Figure S2. Distribution and stability assessment of univariate graph statistics. (**A**, **B**) The distributions of each computed univariate statistic across all subjects and perturbations for Pipeline and Input settings, respectively. There was no significant difference between the distributions in A and B. (**C**, **D**; top) The number of significant decimal digits in each statistic across perturbations, averaged across individuals. The dashed red line refers to the maximum possible number of significant digits. (**C**, **D**; bottom) The percentage of connectomes which were deemed significantly different (p < 0.05) from the others obtained for an individual.