# **Numerical Instabilities in Analytical Pipelines Lead to** Large and Meaningful Variability in Brain Networks

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### 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 can lead to substantial differences in the results, endangering the trust in conclusions<sup>1-7</sup>. We explored the instability of results by instrumenting a connectome estimation pipeline with Monte Carlo Arithmetic<sup>8,9</sup> to introduce random noise throughout. We evaluated the reliability of the connectomes, their features<sup>10,11</sup>, and the impact on analysis<sup>12,13</sup>. 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 scientific computing, and how numerical analysis can be applied to typical analytical workflows. 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

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The modelling of brain networks, called connectomics, 10 2 has shaped our understanding of the structure and function 11 plex computational methods and software. Tools are trusted to 5 obtained in vivo through Magnetic Resonance Imaging (MRI), 14 rigorous evaluation on bespoke datasets, in the absence of 6 and show promise towards identifying biomarkers of disease. 15 ground-truth this is often evaluated through measures of re-8 topathies", such as Alzheimer's Disease and Schizophrenia, 17 existing theory. Importantly, this means that tools are not 9 but potentially pave the way for therapeutics <sup>19–23</sup>.

However, the analysis of brain imaging data relies on com-3 of the brain across a variety of organisms and scales over 12 perform everything from pre-processing tasks to downstream 4 the last decade 11, 14–18. In humans, these wiring diagrams are 13 statistical evaluation. While these tools undoubtedly undergo <sup>7</sup> This can not only improve understanding of so-called "connec-16 liability<sup>24-27</sup>, proxy outcome statistics, or agreement with 18 necessarily of known or consistent quality, and it is not un24 and it is likely that software instabilities played a role.

33 the impact of perturbations through the direct comparision 69 correlations can be found in Supplemental Section S1. 34 of structural connectomes, the consistency of their features, 70 39 reliability of brain imaging research.

### 40 Graphs Vary Widely With Perturbations

<sup>41</sup> Prior to exploring the analytic impact of instabilities, a direct 42 understanding of the induced variability was required. A sub-43 set of the Nathan Kline Institute Rockland Sample (NKIRS) 44 dataset<sup>29</sup> was randomly selected to contain 25 individuals with 45 two sessions of imaging data, each of which was subsampled 46 into two components, resulting in four collections per individ-47 ual. Structural connectomes were generated with canonical 48 deterministic and probabilistic pipelines<sup>30,31</sup> which were in-49 strumented with MCA, replicating computational noise at 50 either the inputs or throughout the pipelines<sup>4,9</sup>. The pipelines 51 were sampled 20 times per collection and once without per- 87 Subject-Specific Signal is Amplified While Off-Target 52 turbations, resulting in a total of 4,200 connectomes.

54 deviation from reference and the number of significant digits 90 ing and extending a typical test-retest experiment 26 in which

19 common that equivalent experiments may lead to diverging 55 (Figure 1). The comparisons were grouped according to dif-20 conclusions<sup>1,5–7</sup>. While many scientific disciplines suffer 56 ferences across simulations, subsampling of data, sessions of 21 from a lack of reproducibility<sup>28</sup>, this was recently explored 57 acquisition, or subjects. While the similarity of connectomes 22 in brain imaging by a 70 team consortium which performed 58 decreases as the collections become more distinct, connec-23 equivalent analyses and found widely inconsistent results<sup>1</sup>, 59 tomes generated with input perturbations show considerable 60 variability, often reaching deviations equal to or greater than The present study approached evaluating reproducibility 61 those observed across individuals or sessions (Figure 1A; 26 from a computational perspective in which a series of brain 62 right). This finding suggests that instabilities inherent to 27 imaging studies were numerically perturbed such that the 63 these pipelines may mask session or individual differences, 28 plausibility of results was not affected, and the biological 64 limiting the trustworthiness of derived connectomes. While 29 implications of the observed instabilities were quantified. We 65 both pipelines show similar performance, the probabilistic 30 accomplished this through the use of Monte Carlo Arithmetic 66 pipeline was more stable in the face of pipeline perturbations 31 (MCA)8, a technique which enables characterization of the 67 whereas the deterministic was more stable to input pertur- $_{32}$  sensitivity of a system to small perturbations. We explored  $_{68}$  bations (p < 0.0001 for all; exploratory). The stability of

The number of significant digits per edge across connec-35 and their eventual application in a neuroscience study. Finally 71 tomes (Figure 1B) similarly decreases across groups. While 36 we conclude on the consequences and opportunities afforded 72 the cross-MCA comparison of connectomes generated with 37 by the observed instabilities and make recommendations for 73 pipeline perturbations show nearly perfect precision for many 38 the roles stability analyses may play towards increasing the 74 edges (approaching the maximum of 15.7 digits for 64-bit 75 data), this evaluation uniquely shows considerable drop off  $_{76}$  in performance across data subsampling (average of < 4 dig-77 its). In addition, input perturbations show no more than an 78 average of 3 significant digits across all groups, demonstrat-79 ing a significant limitation in the reliability independent edge 80 weights. Significance across individuals did not exceed a 81 single digit per edge in any case, indicating that only the 82 magnitude of edges in naively computed groupwise average 83 connectomes can be trusted. The combination of these results 84 with those presented in Figure 1A suggests that while specific 85 edge weights are largely affected by instabilities, macro-scale 86 network topology is stable.

# 88 Biases Are Reduced

The stability of connectomes was evaluated through the 89 We assessed the reproducibility of the dataset through mimick-

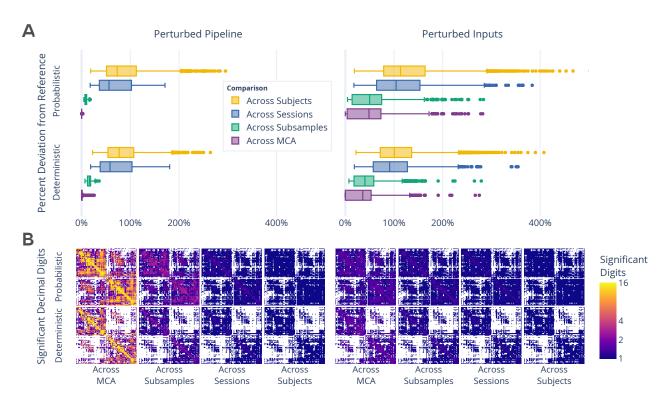


Figure 1. Exploration of perturbation-induced deviations from reference connectomes. (A) The absolute deviations, in the form of normalized percent deviation from reference, shown as the across MCA series relative to Across Subsample, Across Session, and Aross Subject variations. (B) The number of significant 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. Pipeline- and input-perturbations are shown on the left and right, respectively.

91 the similarity of samples across multiple measurements were 106 ing as a cost effective and context-agnostic method for dataset 92 compared to distinct samples in the dataset (Table 1, with 107 augmentation. 93 additional experiments and explanation in Supplemental Sec-94 tion S2). The ability to separate connectomes across subjects 95 (Hypothesis 1) is an essential prerequisite for the application 96 of brain imaging towards identifying individual differences<sup>18</sup>. 97 In testing hypothesis 1, we observe that the dataset is sep-<sub>98</sub> arable with a score of 0.64 and 0.65 (p < 0.001; optimal 99 score: 1.0; chance: 0.04) without any instrumentation. How-100 ever, we can see that inducing instabilities through MCA improves the reliability of the dataset to over 0.75 in each  $_{102}$  case (p < 0.001 for all), significantly higher than without instrumentation (p < 0.005 for all). This result impactfully 104 suggests the utility of perturbation methods for synthesizing 105 robust and reliable individual estimates of connectivity, serv-

While the separability of individuals is essential for the 109 identification of brain networks, it is similarly reliant on network similarity across equivalent acquisitions (Hypothesis 2). In this case, connectomes were grouped based upon session, 112 rather than subject, and the ability to distinguish one session 113 from another was computed within-individual and aggregated. Both the unperturbed and pipeline perturbation settings per-115 fectly preserved differences between cross-sectional sessions with a score of 1.0 (p < 0.005; optimal score: 0.5; chance: 117 0.5), indicating a dominant session-dependent signal for all 118 individuals despite no intended biological differences. However, while still significant relative to chance (score: 0.85) and 0.88; p < 0.005 for both), input perturbations lead to

**Table 1.** The impact of instabilities as evaluated through the separability of the dataset based on individual (or subject) differences, session, and subsample. The performance is reported as mean Discriminability. While a perfectly separable dataset would be represented by a score of 1.0, the chance performance, indicating minimal separability, 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 separation, was accepted for all experiments, with p < 0.005.

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

significantly lower separability of the dataset (p < 0.005 for 147 in machine learning, a concept which observes a decrease in 122 all). This reduction of the difference between sessions of data 148 bias as variance is favoured by a model. In particular, this 123 within individuals suggests that increased variance caused 149 highlights that numerical perturbations can be used to not 124 by input perturbations reduces the impact of non-biological 150 only evaluate the stability of pipelines, but that the induced acquisition-dependent bias inherent in the brain graphs.

Though the previous sets of experiments inextricably eval- 152 distributions of possible results. 127 uate the interaction between the dataset and tool, the use of 128 subsampling allowed for characterizing the separability of 153 Distributions of Graph Statistics Are Reliable, But 129 networks sampled from within a single acquisition (Hypoth- 154 Individual Statistics Are Not 150 esis 3). While this experiment could not be evaluated using 155 Exploring the stability of topological features of connectomes 131 reference executions, the executions performed with pipeline 156 is relevant for typical analyses, as low dimensional features are 132 perturbations showed near perfect separation between sub- 157 often more suitable than full connectomes for many analytical samples, with scores of 0.99 and 1.0 (p < 0.005; optimal: 158 methods in practice 11. A separate subset of the NKIRS dataset 134 0.5; chance: 0.5). Given that there is no variability in data 159 was randomly selected to contain a single non-subsampled acquisition or preprocessing that contributes to this reliable 160 session for 100 individuals, and connectomes were generated 136 identification of scans, the separability observed in this exper- 161 as above. 137 iment may only be due to instability or bias inherent to the 162

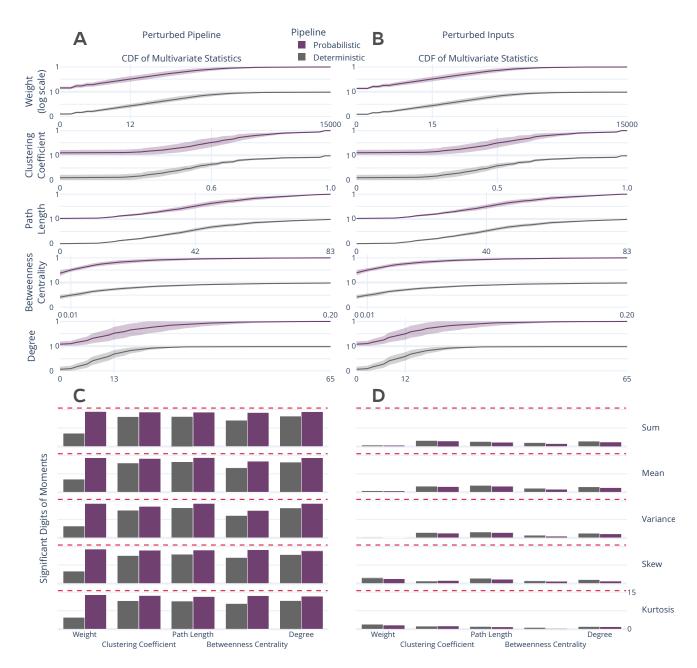
amplification of meaningful biological signal alongside a re- 169 logical features summarized by these multivariate features are duction of off-target signal. This result appears strikingly like 170 robust across both perturbation modes. 146 a manifestation of the well-known bias-variance tradeoff<sup>32</sup> 171

142 of individual connectivity.

151 variance may be leveraged for the interpretation as a robust

The stability of several commonly-used multivariate graph pipelines. The high variability introduced through input per- 163 features 10 was explored in Figure 2. The cumulative den-139 turbations considerably lowered the reliability towards chance 164 sity of the features was computed within individuals and the 140 (score: 0.71 and 0.61; p < 0.005 for all), further supporting 165 mean density and associated standard error were computed this as an effective method for obtaining lower-bias estimates 166 for across individuals (Figures 2A and 2B). There was no sig-167 nificant difference between the distributions for each feature Across all cases, the induced perturbations showed an 168 across the two perturbation settings, suggesting that the topo-

In addition to the comparison of distributions, the stabil-



**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.

172 ity of the first 5 moments of these features was evaluated 177 more stable for all comparisons (p < 0.0001; exploratory). (Figures 2C and 2D). In the face of pipeline perturbations, 178 In stark contrast, input perturbations led to highly unstable 174 the feature-moments were stable with more than 10 signifi- 179 feature-moments (Figure 2D), such that none contained more 175 cant digits with the exception of edge weight when using the 180 than 5 significant digits of information and several contained 176 deterministic pipeline, though the probabilistic pipeline was 181 less than a single significant digit, indicating a complete lack

182 of reliability. This dramatic degradation in stability for in- 217 Discussion dividual measures strongly suggests that these features may 218 The perturbation of structural connectome estimation pipelines 188 can be found in Supplemental Section S3.

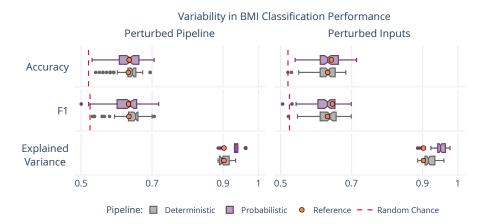
## 189 Uncertainty in Brain-Phenotype Relationships

198 ure 3).

 $_{\rm 202}$  from 0.520-0.716 and 0.510-0.725, respectively, rang-  $^{\rm 238}$  of individual estimates. 203 ing from at or below random performance to outperforming 239 Underestimated False Positive Rates While the instabilperformance on the reference dataset. This large variability 240 ity of brain networks was used here to demonstrate the lim-205 illustrates a previously uncharacterized margin of uncertainty 241 itations of modelling brain-phenotype relationships in the 206 in the modelling of this relationship, and limits confidence in 242 context of machine learning, this limitation extends to classi-207 reported accuracy scores on singly processed datasets. The 243 cal hypothesis testing, as well. Though performing individual 208 portion of explained variance in these samples ranged from 244 comparisons in a hypothesis testing framework will be accom-88.6% - 97.8%, similar to the reference, suggesting that the 245 panied by reported false positive rates, the accuracy of these 210 range in performance was not due to a gain or loss of mean- 246 rates is critically dependent upon the reliability of the samples 211 ingful signal, but rather the reduction of bias towards specific 247 used. In reality, the true false positive rate for a test would be outcome. Importantly, this finding does not suggest that mod- 248 a combination of the reported confidence and the underlying elling brain-phenotype relationships is not possible, but rather 249 variability in the results, a typically unknown quantity. 214 it sheds light on impactful uncertainty that must be accounted 250 216 techniques.

be unreliable as individual biomarkers when derived from a 219 with small amounts of noise, on the order of machine error, single pipeline evaluation, though their reliability may be in- 220 led to considerable variability in derived brain graphs. Across 186 creased when studying their distributions across perturbations. 221 all analyses the stability of results ranged from nearly per-187 A similar analysis was performed for univariate statistics and 222 fectly trustworthy (i.e. no variation) to completely unreliable 223 (i.e. containing no trustworthy information). Given that the 224 magnitude of introduced numerical noise is to be expected 225 in typical settings, this finding has potentially significant im-190 While the variability of connectomes and their features was 226 plications for inferences in brain imaging as it is currently summarized above, networks are commonly used as inputs to 227 performed. In particular, this bounds the success of studying machine learning models tasked with learning brain-phenotype 228 individual differences, a central objective in brain imaging 18, 193 relationships 18. To explore the stability of these analyses, we 229 given that the quality of relationships between phenotypic modelled the relationship between high- or low- Body Mass 230 data and brain networks will be limited by the stability of the 195 Index (BMI) groups and brain connectivity 12,13, using stan-231 connectomes themselves. This issue was accentuated through dard dimensionality reduction and classification tools, and 232 the crucial finding that individually derived network features 197 compared this to reference and random performance (Fig- 233 were unreliable despite there being no significant difference 234 in their aggregated distributions. This finding is not damn-The analysis was perturbed through distinct samplings of 235 ing for the study of brain networks as a whole, but rather is 200 the dataset across both pipelines and perturbation methods. 236 strong support for the aggregation of networks, either across 201 The accuracy and F1 score for the perturbed models varied 237 perturbations for an individual or across groups, over the use

When performing these experiments outside of a repeatedfor in this process, and supports the use of ensemble modeling 251 measure context, such as that afforded here through MCA, it 252 is impossible to empirically estimate the reliability of samples. 253 This means that the reliability of accepted hypotheses is also



**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.

254 unknown, regardless of the reported false positive rate. In 278 effectively in conjunction with, or in lieu of, realistic noise 255 fact, it is a virtual certainty that the true false positive rate 279 models to augment existing datasets. While this of course 256 for a given hypothesis exceeds the reported value simply as 280 would not replace the need for repeated measurements when 257 a result of numerical instabilities. This uncertainty inherent 281 exploring the effect of data collection paradigm or study lon-258 to derived data is compounded with traditional arguments 282 gitudinal progressions of development or disease, it could be 259 limiting the trustworthiness of claims<sup>33</sup>, and hampers the 283 used in conjunction with these efforts to increase the reliabil-260 ability of researchers to evaluate the quality of results. The 284 ity of each distinct sample within a dataset. In contexts where 261 accompaniment of brain imaging experiments with direct 285 repeated measurements are collected to increase the fidelity of 262 evaluations of their stability, as was done here, would allow 286 the dataset, MCA could potentially be employed to increase 263 researchers to simultaneously improve the numerical stability 287 the reliability of the dataset and save millions of dollars on 264 of their analyses and accurately gauge confidence in them. 288 data collection. This technique also opens the door for the 265 The induced variability in derived brain networks may be 289 characterization of reliability across axes which have been 266 leveraged to estimate aggregate connectomes with lower bias 290 traditionally inaccessible. For instance, in the absence of a 267 than any single independent observation, leading to learned 291 realistic noise model or simulation technique similar to MCA, 268 relationships that are more generalizable and ultimately more 292 the evaluation of network stability across data subsampling 269 useful. 293 would not have been possible.

Cost-Effective Data Augmentation The evaluation of reli- 294 Shortcomings and Future Questions Given the complex- ability in brain imaging has historically relied upon the ex- 295 ity of recompiling complex software libraries, pre-processing pensive collection of repeated measurements choreographed 296 was not perturbed in these experiments. Other work has shown by massive cross-institutional consortia 4,35. The finding that 297 that linear registration, a core piece of many elements of pre- 274 perturbing experiments using MCA both increased the relia- 298 processing such as motion correction and alignment, is sensi-275 bility of the dataset and decreased off-target differences across 299 tive to minor perturbations 7. It is likely that the instabilities acquisitions opens the door for a promising paradigm shift. 300 across the entire processing workflow would be compounded Given that MCA is data-agnostic, this technique could be used 301 with one another, resulting in even greater variability. While

302 the analyses performed in this paper evaluated a single dataset and set of pipelines, extending this work to other modalities 304 and analyses is of interest for future projects.

This paper does not explore methodological flexibility or 306 compare this to numerical instability. Recently, the nearly 307 boundless space of analysis pipelines and their impact on out-308 comes in brain imaging has been clearly demonstrated<sup>1</sup>. The 309 approach taken in these studies complement one another and 310 explore instability at the opposite ends of the spectrum, with 311 human variability in the construction of an analysis workflow 312 on one end and the unavoidable error implicit in the digital 313 representation of data on the other. It is of extreme interest 314 to combine these approaches and explore the interaction of 315 these scientific degrees of freedom with effects from software 316 implementations, libraries, and parametric choices.

318 presented here does not invalidate analytical pipelines used in 319 brain imaging, but merely sheds light on the fact that many 320 studies are accompanied by an unknown degree of uncertainty due to machine-introduced errors. The presence of unknown 322 error-bars associated with experimental findings limits the 323 impact of results due to increased uncertainty. The desired 324 outcome of this paper is to motivate a shift in scientific com-325 puting – both in neuroimaging and more broadly – towards a paradigm which favours the explicit evaluation of the trust- 351 extra level of stability evaluation to be performed between the 327 worthiness of claims alongside the claims themselves.

## **Methods**

#### 329 Dataset

330 The Nathan Kline Institute Rockland Sample (NKI-RS)<sup>29</sup> 331 dataset contains high-fidelity imaging and phenotypic data 332 from over 1,000 individuals spread across the lifespan. A 333 subset of this dataset was chosen for each experiment to both 334 match sample sizes presented in the original analyses and to 335 minimize the computational burden of performing MCA. The 336 selected subset comprises 100 individuals ranging in age from 3376 - 79 with a mean of 36.8 (original: 6 - 81, mean 37.8), 338 60% female (original: 60%), with 52% having a BMI over 25 339 (original: 54%).

Each selected individual had at least a single session 341 of both structural T1-weighted (MPRAGE) and diffusion-Finally, it is important to state explicitly that the work 342 weighted (DWI) MR imaging data. DWI data was acquired 343 with 137 diffusion directions; more information regarding the 344 acquisition of this dataset can be found in the NKI-RS data з45 release<sup>29</sup>.

> In addition to the 100 sessions mentioned above, 25 indi-347 viduals had a second session to be used in a test-retest analysis. 348 Two additional copies of the data for these individuals were 349 generated, including only the odd or even diffusion directions  $_{350}$  (64 + 9 B0 volumes = 73 in either case). This allowed for an 352 levels of MCA and session-level variation.

> In total, the dataset is composed of 100 downsampled 354 sessions of data originating from 50 acquisitions and 25 in-355 dividuals for in depth stability analysis, and an additional 356 100 sessions of full-resolution data from 100 individuals for 357 subsequent analyses.

#### 358 Processing

359 The dataset was preprocessed using a standard FSL36 work-360 flow consisting of eddy-current correction and alignment. The 361 MNI152 atlas<sup>37</sup> was aligned to each session of data, and the re-362 sulting transformation was applied to the DKT parcellation<sup>38</sup>. 363 Downsampling the diffusion data took place after preprocess365 an additional confound was not introduced in this process 399 referred to as Precision Bounding (PB) and the latter is called 366 when comparing between downsampled sessions. The pre-400 Random Rounding (RR). 367 processing described here was performed once without MCA, 401 368 and thus is not being evaluated.

370 data using two canonical pipelines from Dipy<sup>30</sup>: deterministic 404 stability of the instrumented tools or functions. To this end, and probabilistic. In the deterministic pipeline, a constant 405 a complete software stack was instrumented with MCA and 372 solid angle model was used to estimate tensors at each voxel 406 is made available on GitHub at https://github.com/ and streamlines were then generated using the EuDX algo- qkiar/fuzzy. 374 rithm<sup>31</sup>. In the probabilistic pipeline, a constrained spherical 375 deconvolution model was fit at each voxel and streamlines 376 were generated by iteratively sampling the resulting fiber ori-377 entation distributions. In both cases tracking occurred with 8 378 seeds per 3D voxel and edges were added to the graph based on the location of terminal nodes with weight determined by 413 plied across the bulk of the relevant libraries and is referred 380 fiber count.

The random state of the probabilistic pipeline was fixed 382 for all analyses. Fixing this random seed allowed for explicit 383 attribution of observed variability to Monte Carlo simulations 384 rather than internal state of the algorithm.

## 385 Perturbations

386 All connectomes were generated with one reference execu-387 tion where no perturbation was introduced in the processing. 388 For all other executions, all floating point operations were 389 instrumented with Monte Carlo Arithmetic (MCA)<sup>8</sup> through <sup>390</sup> Verificarlo<sup>9</sup>. MCA simulates the distribution of errors im-391 plicit to all instrumented floating point operations (flop). This 392 rounding is performed on a value x at precision t by:

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

where  $e_x$  is the exponent value of x and  $\xi$  is a uniform ran- 429 **Evaluation** dom variable in the range  $(-\frac{1}{2}, \frac{1}{2})$ . MCA can be introduced in 430 The magnitude and importance of instabilities in pipelines 395 two places for each flop: before or after evaluation. Perform- 431 can be considered at a number of analytical levels, namely: 396 ing MCA on the inputs of an operation limits its precision, 432 the induced variability of derivatives directly, the resulting

364 ing was performed on full-resolution sessions, ensuring that 398 lights round-off errors that may be introduced. The former is

Using MCA, the execution of a pipeline may be performed 402 many times to produce a distribution of results. Studying the Structural connectomes were generated from preprocessed 403 distribution of these results can then lead to insights on the

> Both the RR and PB variants of MCA were used indepen-409 dently for all experiments. As was presented in<sup>4</sup>, both the 410 degree of instrumentation (i.e. number of affected libraries) and the perturbation mode have an effect on the distribution 412 of observed results. For this work, the RR-MCA was ap-414 to as Pipeline Perturbation. In this case the bulk of numerical 415 operations were affected by MCA.

> Conversely, the case in which PB-MCA was applied across 417 the operations in a small subset of libraries is here referred 418 to as Input Perturbation. In this case, the inputs to operations 419 within the instrumented libraries (namely, Python and Cython) 420 were perturbed, resulting in less frequent, data-centric pertur-421 bations. Alongside the stated theoretical differences, Input 422 Perturbation is considerably less computationally expensive 423 than Pipeline Perturbation.

> All perturbations targeted the least-significant-bit for all 425 data (t = 24 and t = 53 in float32 and float64, respectively<sup>9</sup>). 426 Simulations were performed 20 times for each pipeline execu-427 tion. A detailed motivation for the number of simulations can 428 be found in<sup>39</sup>.

397 while performing MCA on the output of an operation high- 493 downstream impact on summary statistics or features, or the

434 ultimate change in analyses or findings. We explore the na- 468 Class-based Variability Evaluation To gain a concrete un-495 ture and severity of instabilities through each of these lenses. 469 derstanding of the significance of observed variations we ex-496 Unless otherwise stated, all p-values were computed using 470 plore the separability of our results with respect to understood 437 Wilcoxon signed-rank tests.

# 438 Direct Evaluation of the Graphs

439 The differences between simulated graphs was measured di-440 rectly through both a direct variance quantification and a 441 comparison to other sources of variance such as individual-442 and session-level differences.

443 Quantification of Variability Graphs, in the form of adja-444 cency matrices, were compared to one another using three 445 metrics: normalized percent deviation, Pearson correlation, 446 and edgewise significant digits. The normalized percent devi- 477 ation measure, defined in  $^4$ , scales the norm of the difference  $^{478}$  at observation j, where  $i \neq i'$  and  $j \neq j'$ . 448 between a simulated graph and the reference execution (that 479 without intentional perturbation) with respect to the norm of 480 observation belonging to a given class will be more similar to 450 the reference graph. The purpose of this comparison is to 481 other observations within that class than observations of a difprovide insight on the scale of differences in observed graphs 482 ferent class. It is a measure of reproducibility, and is discussed 452 relative to the original signal intensity. A Pearson correlation 483 in detail in<sup>26</sup>. This definition allows for the exploration of 453 coefficient 40 was computed in complement to normalized per-484 deviations across arbitrarily defined classes which in practice 454 cent deviation to identify the consistency of structure and not 485 can be any of those listed above. We combine this statistic 455 just intensity between observed graphs.

457 each edge in the graph is calculated as:

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

461 data. The percent deviation, correlation, and number of signifi- 496 the reference executions alongside using MCA. 463 cant digits were each calculated within a single session of data, 464 thereby removing any subject- and session-effects and provid- 497  $H_{A1}$ : Individuals are distinct from one another 465 ing a direct measure of the tool-introduced variability across 498 466 perturbations. A distribution was formed by aggregating these 499 467 individual results.

471 sources of variability, such as subject-, session-, and pipeline-472 level effects. This can be probed through Discriminability<sup>26</sup>. <sup>473</sup> a technique similar to ICC<sup>24</sup> which relies on the mean of a 474 ranked distribution of distances between observations belong-475 ing to a defined set of classes. The discriminability statistic is 476 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

Discriminability can then be read as the probability that an 486 with permutation testing to test hypotheses on whether differ-Finally, the estimated number of significant digits, s', for 487 ences between classes are statistically significant in each of 488 these settings.

With this in mind, three hypotheses were defined. For 490 each setting, we state the alternate hypotheses, the variable(s) 491 which were used to determine class membership, and the where  $\mu$  and  $\sigma$  are the mean and unbiased estimator of 492 remaining variables which may be sampled when obtaining 459 standard deviation across graphs, respectively. The upper 493 multiple observations. Each hypothesis was tested indepenbound on significant digits is 15.7 for 64-bit floating point 494 dently for each pipeline and perturbation mode, and in every 495 case where it was possible the hypotheses were tested using

Class definition: Subject ID

Comparisons: Session (1 subsample), Subsample (1 session), MCA (1 subsample, 1 session)

501  $H_{A2}$ : Sessions within an individual are distinct Class definition: Session ID | Subject ID Comparisons: Subsample, MCA (1 subsample) 503

504  $H_{A3}$ : Subsamples are distinct

Class definition: Subsample | Subject ID, Session ID

Comparisons: MCA 506

As a result, we tested 3 hypotheses across 6 MCA ex-508 periments and 3 reference experiments on 2 pipelines and 2 509 perturbation modes, resulting in a total of 30 distinct tests.

#### **510 Evaluating Graph-Theoretical Metrics**

512 it is common practice to summarize them with structural mea- 548 tions. We performed the modeling task with a single sampled 513 sures, which can then be used as lower-dimensional proxies 549 connectome per individual and repeated this sampling and of connectivity in so-called graph-theoretical studies<sup>11</sup>. We 550 modelling 20 times. We report the model performance for explored the stability of several commonly-used univariate 551 each sampling of the dataset and summarize its variance. 516 (graphwise) and multivariate (nodewise or edgewise) features. 517 The features computed and subsequent methods for compari-518 son in this section were selected to closely match those com-519 puted in<sup>10</sup>.

521 count, mean clustering coefficient, global efficiency, modu- 557 cipal component analysis (PCA), and provided the first N-<sup>522</sup> larity of the largest connected component, assortativity, and <sup>558</sup> components to a logistic regression classifier for predicting mean path length) a distribution of values across all perturba- 559 BMI class membership, similar to methods shown in 12,13. 524 tions within subjects was observed. A Z-score was computed 560 The number of components was selected as the minimum set 525 for each sample with respect to the distribution of feature 561 which explained > 90% of the variance when averaged across 526 values within an individual, and the proportion of "classically 562 the training set for each fold within the cross validation of significant" Z-scores, i.e. corresponding to p < 0.05, was 563 the original graphs; this resulted in a feature of 20 compo-528 reported and aggregated across all subjects. The number of 564 nents. We trained the model using k-fold cross validation, significant digits contained within an estimate derived from a set with k = 2, 5, 10, and N (equivalent to leave-one-out; LOO). 530 single subject were calculated and aggregated.

531 Multivariate Differences In the case of both nodewise (de- 567 The unprocessed dataset is available through The Consortium 532 gree distribution, clustering coefficient, betweenness central- 568 of Reliability and Reproducibility (http://fcon\_1000. 553 ity) and edgewise (weight distribution, connection length) fea- 569 projects.nitrc.org/indi/enhanced/), including 534 tures, the cumulative density functions of their distributions 570 both the imaging data as well as phenotypic data which may

536 gated across individuals. The number of significant digits 537 for each moment of these distributions (sum, mean, variance, 538 skew, and kurtosis) were calculated across observations within 539 a sample and aggregated.

### 540 Evaluating A Brain-Phenotype Analysis

Though each of the above approaches explores the instabil-542 ity of derived connectomes and their features, many modern 543 studies employ modeling or machine-learning approaches, for 544 instance to learn brain-phenotype relationships or identify dif-545 ferences across groups. We carried out one such study and ex-546 plored the instability of its results with respect to the upstream While connectomes may be used directly for some analyses, 547 variability of connectomes characterized in the previous sec-

552 **BMI Classification** Structural changes have been linked to obesity in adolescents and adults<sup>41</sup>. We classified normal-554 weight and overweight individuals from their structural net- $_{555}$  works (using for overweight a cutoff of BMI  $> 25^{13}$ ). We 520 Univariate Differences For each univariate statistic (edge 556 reduced the dimensionality of the connectomes through prin-

#### 566 Data & Code Provenance

595 were evaluated over a fixed range and subsequently aggre- 571 be obtained upon submission and compliance with a Data Us-

age Agreement. The connectomes generated through simula- 606 References 573 tions have been bundled and stored permanently (https:// 607 [1] 574 doi.org/10.5281/zenodo.4041549), and are made 608 575 available through The Canadian Open Neuroscience Platform 576 (https://portal.conp.ca/search, search term "Kiar").[2] All software developed for processing or evaluation is 612 578 publicly available on GitHub at https://github.com/ 579 gkpapers/2020ImpactOfInstability. Experiments [3] 580 were launched using Boutiques<sup>42</sup> and Clowdr<sup>43</sup> in Compute 616 581 Canada's HPC cluster environment. MCA instrumentation 617 582 was achieved through Verificarlo<sup>9</sup> available on Github at 583 https://github.com/verificarlo/verificarlo. 584 A set of MCA instrumented software containers is available 621 585 **on Github at** https://github.com/gkiar/fuzzy.

#### 586 Author Contributions

587 GK was responsible for the experimental design, data pro- 626 588 cessing, analysis, interpretation, and the majority of writing. 627 589 All authors contributed to the revision of the manuscript. YC, 590 POC, and EP were responsible for MCA tool development and 591 software testing. AR, GV, and BM contributed to experimen- 631 [7] 592 tal design and interpretation. TG contributed to experimental 632 593 design, analysis, and interpretation. TG and ACE were re-594 sponsible for supervising and supporting all contributions 595 made by GK. The authors declare no competing interests for 636 596 this work. Correspondence and requests for materials should 637 597 be addressed to Tristan Glatard at tristan.glatard@ 598 concordia.ca.

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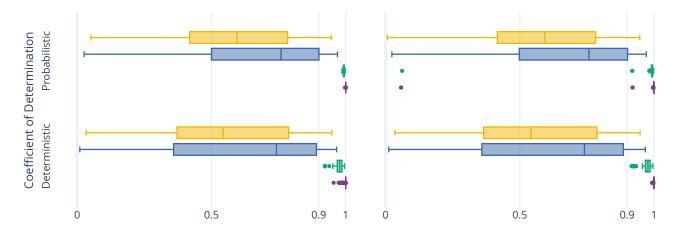
# $Numerical\ Instabilities\ in\ Analytical\ Pipelines\ Lead\ to\ Large\ and\ Meaningful\ Variability\ in\ Brain\ Networks\ --\ 14/18$

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# 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 others<sup>44</sup>, correlation is not a commonly used measure in the context of structural connectivity.



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

# 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	$0.64 \pm 0.00$	$0.65 \pm 0.00$	$0.82 \pm 0.00$	$0.82\pm0.00$	$0.77 \pm 0.00$	$0.75 \pm 0.00$	
1.2	All	1	All	$1.00 \pm 0.00$	$1.00\pm0.00$	$1.00 \pm 0.00$	$1.00\pm0.00$	$0.93 \pm 0.02$	$0.90\pm0.02$	
1.3	All	1	1			$1.00 \pm 0.00$	$1.00\pm0.00$	$0.94 \pm 0.02$	$0.90\pm0.02$	
2.4	1	All	All	$1.00 \pm 0.00$	$1.00\pm0.00$	$1.00 \pm 0.00$	$1.00\pm0.00$	$0.88 \pm 0.12$	$0.85\pm0.12$	
2.5	1	All	1			$1.00 \pm 0.00$	$1.00\pm0.00$	$0.89 \pm 0.11$	$0.84 \pm 0.12$	
3.6	1	1	All			$0.99 \pm 0.03$	$1.00\pm0.00$	$0.71\pm0.07$	$0.61\pm0.05$	

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 rould 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.

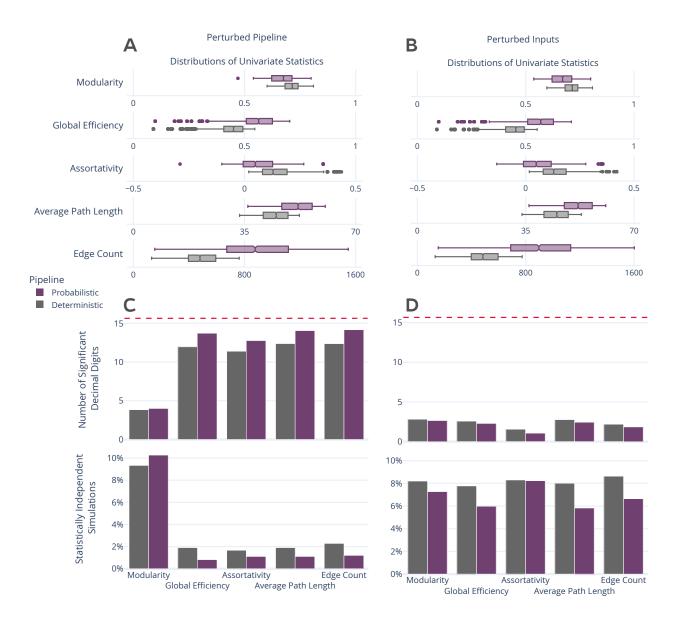
767 **Session Variation** Similar to subject variation, the session variation was also modelled across either both or a single subsample. In both of these cases the performance was similar, and the finding that input perturbation reduced the off-target signal was consistent.

# 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 more than 3 significant digits or a false positive rate lower than nearly 6% (Figure S2D). The deterministic pipeline was more more than 4 significant digits or a false positive rate lower than nearly 6% (Figure S2D). The deterministic pipeline was more than 5 significant digits or a false positive rate lower than nearly 6% (Figure S2D).

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.