

Original citation:

Nichols, Thomas E., Das, Samir, Eickhoff, Simon B., Evans, Alan C., Glatard, Tristan, Hanke, Michael, Kriegeskorte, Nikolaus, Milham, Michael P., Poldrack, Russell A., Poline, Jean-Baptiste, Proal, Erika, Thirion, Bertrand, Van Essen, David C., White, Tonya and Yeo, B.T. Thomas. (2017) Best practices in data analysis and sharing in neuroimaging using MRI. *Nature Neuroscience*, 20 (3). pp. 299-303.

Permanent WRAP URL:

<http://wrap.warwick.ac.uk/85042>

Copyright and reuse:

The Warwick Research Archive Portal (WRAP) makes this work by researchers of the University of Warwick available open access under the following conditions. Copyright © and all moral rights to the version of the paper presented here belong to the individual author(s) and/or other copyright owners. To the extent reasonable and practicable the material made available in WRAP has been checked for eligibility before being made available.

Copies of full items can be used for personal research or study, educational, or not-for-profit purposes without prior permission or charge. Provided that the authors, title and full bibliographic details are credited, a hyperlink and/or URL is given for the original metadata page and the content is not changed in any way.

Publisher's statement:

<https://doi.org/10.1101/054262>

A note on versions:

The version presented here may differ from the published version or, version of record, if you wish to cite this item you are advised to consult the publisher's version. Please see the 'permanent WRAP URL' above for details on accessing the published version and note that access may require a subscription.

For more information, please contact the WRAP Team at: wrap@warwick.ac.uk

Best Practices in Data Analysis and Sharing in Neuroimaging using MRI

Thomas E. Nichols^{1,*}, Samir Das^{2,15}, Simon B. Eickhoff³, Alan C. Evans^{2,15}, Tristan Glatard^{2,14}, Michael Hanke^{4,16}, Nikolaus Kriegeskorte⁵, Michael P. Milham^{6,17}, Russell A. Poldrack⁷, Jean-Baptiste Poline⁸, Erika Proal⁹, Bertrand Thirion¹⁰, David C. Van Essen¹¹, Tonya White¹², B.T. Thomas Yeo¹³

¹University of Warwick, Coventry, UK; ²McGill University, Montreal, Canada; ³Heinrich-Heine University Düsseldorf, Düsseldorf, Germany; ⁴Otto-von-Guericke-University, Magdeburg, Germany; ⁵MRC Cognition and Brain Sciences Unit, Cambridge, UK; ⁶Child Mind Institute, New York, USA; ⁷Stanford University, Stanford, USA; ⁸University of California, Berkeley, USA; ⁹NEUROingenia Clinical and Research Center, Mexico City, Mexico; ¹⁰Inria, Paris-Saclay University, Paris, France; ¹¹Washington University in St. Louis, St. Louis, USA; ¹²Erasmus University Medical Center, Rotterdam, The Netherlands; ¹³National University of Singapore, Singapore; ¹⁴Department of Computer Science and Software Engineering, Concordia University, Montreal, Canada; ¹⁵Montreal Neurological Institute, Montreal, Canada; ¹⁶Center for Behavioral Brain Sciences, Magdeburg, Germany; ¹⁷Nathan S. Kline Institute for Psychiatric Research, New York, USA;

*Correspondence: t.e.nichols@warwick.ac.uk.

Words: 3250 (excluding abstract)

Figures: 0

Tables: 1

References: 24

Abstract

Given concerns about the reproducibility of scientific findings, neuroimaging must define best practices for data analysis, results reporting, and algorithm and data sharing to promote transparency, reliability and collaboration. We describe insights from developing a set of recommendations on behalf of the Organization for Human Brain Mapping, and identify barriers that impede these practices, including how the discipline must change to fully exploit the potential of the world's neuroimaging data.

[Start of body text]

The advancement of science requires continuous examination of the principles and practices by which the research community operates. In recent years, this ongoing evaluative process has flagged concerns about the reproducibility of published research. From the early claim by John

38 Ioannidis in 2005 that “most published research findings are false”¹ to the recent work by the
39 Open Science Collaboration, which attempted to replicate 100 psychology studies and
40 succeeded in only 39 cases², there is mounting evidence that scientific results are less reliable
41 than widely assumed.

42
43 Efforts promoting open science principles across fields (e.g.³) as a means of fostering
44 transparency and reproducibility are valuable, but we also need efforts focusing specifically on
45 human neuroimaging. To address this need the Organisation for Human Brain Mapping (OHBM)
46 created the Committee on Best Practices in Data Analysis and Sharing (COBIDAS⁴,
47 <http://www.humanbrainmapping.org/cobidas>). This group was charged with creating a report
48 that would compile best practices for open science in neuroimaging and distill these principles
49 into specific research practices. The report was developed in collaboration with the OHBM
50 community, which provided feedback on a draft and ratification of the final version.

51
52 In this commentary, we review the challenging issues that arose in the formation of the report,
53 and identify initial success and the key remaining shortcomings in current practice.

54 What is Reproducibility?

55 Open science comprises a number of different goals and principles. The COBIDAS was
56 specifically concerned with ‘Open Data’ and ‘Open Methodology’, both of which are in service
57 of ‘Open Reproducible Research.’ An immediate challenge was to obtain a working definition of
58 reproducibility. We considered a hierarchy of reproducibility concepts ranging from
59 measurement and analytical stability, to broader notions of generalisability (Table 1). A very
60 narrow notion of generalizability would be test-retest reliability on the same scanner, same
61 subject, within 30 minutes, while a more extended notion would be using different scanners on
62 the same subject with re-imaging occurring within 7 days. Generalization over analyses
63 corresponds to re-analysis of the same data using identical or similar tools. One variant of this is
64 “computational reproducibility”⁵, where independent researchers re-analyse the data and
65 compare their results. We also considered versions of generalizability corresponding to
66 traditional scientific notions of “replication”, such as whether a result is stable over different
67 samples of subjects or populations of subjects. The most challenging, and arguably most
68 important form of generalizability is whether a finding additionally holds under variation in the
69 stimuli and experimental methods. Underlying all of these concerns about reproducibility is how
70 theory-building requires reproducible empirical phenomena, and thus a theory will only be as
71 accurate and generalizable as the data that are used to inspire and/or test it.

72
73 Regardless of the precise scope of generalization, operationalising any of these versions of
74 reproducibility requires explicit definitions of the outcome of interest, which in itself is a
75 challenge. Previous efforts have found generally good measures of test-retest reliability of MRI
76 for both voxel-wise and region of interest measures (e.g. ⁶⁻⁸), but this is the most narrow notion
77 of reproducibility. A large scale project to measure the generalisability of MRI findings across
78 studies, akin to the Open Science Collaboration’s efforts in Psychology², has not been
79 undertaken in neuroimaging; however the one effort that set out to reproduce brain structure-

80 behavior correlations found only 1 of 17 findings were replicated⁹, though this work is limited by
81 small replication sample sizes. More work is needed in this area to better quantify the
82 generalisability of MRI findings.

83
84 In short, quantifying “reproducibility” requires precisely defining the scope of variation being
85 considered, the exact outcome that is being measured, and a metric of the stability of that
86 outcome. The COBIDAS did not set out to estimate reproducibility, but was motivated to identify
87 practices that can maximise analytical stability and generalizability of individual studies.

88 [Table 1 about here]

89 Prescribing best practice

90 Neuroimaging is a broad field, encompassing a range of approaches across a growing number
91 of modalities. We restricted the scope of the COBIDAS report to include the range of all human
92 neuroimaging using Magnetic Resonance Imaging (MRI), though most of the principles
93 discussed can be applied to other modalities. We established 7 domains of practice, from
94 experimental design and acquisition, through results reporting and data sharing. We quickly
95 realised that it is neither feasible nor desirable to prescribe exactly how any one type of
96 experiment should be conducted. For example, when looking at task fMRI, the optimal
97 experimental design to use will depend on whether one is just trying to detect the presence of
98 an effect or rather estimate the shape of the hemodynamic response function.

99
100 The one “practice” that can be universally commended is the transparent and complete
101 reporting of all facets of a study, allowing a critical reader to evaluate the work and fully
102 understand its strengths and limitations. This also facilitates subsequent research efforts by
103 other investigators, who can exactly follow (or carefully manipulate) each aspect of a study. This
104 includes conveying the “researcher degrees of freedom”, by reporting other analytical paths
105 applied unsuccessfully on the present data before arriving at the published results. Although
106 formidable, the reporting checklists provided in the COBIDAS MRI report reflects the breadth
107 and depth of information needed to ensure another researcher could replicate the work.

108
109 To further facilitate reproducibility, the COBIDAS report includes specific recommendations for
110 statistical modelling, where specific (and common) bad practices have been identified^{10,11}. We
111 have also made concrete recommendations for data sharing, where practice is still evolving.

112
113 From solicited community input, we were struck by the emphatic and diverse views on the types
114 of data to share. Some strongly felt it was essential to share the rawest form of the data from
115 the scanner (DICOM format), while others felt that preprocessed, ready-to-analyze data should
116 be shared; still others emphasized the utility of sharing extensively processed data linked to
117 published figures. We evaluated the pros and cons of each form of data sharing; for example,
118 while sharing preprocessed data can minimize the effort needed for reanalysis and speed
119 advances based on new uses of the data, it may preclude alternate preprocessing options that
120 facilitate new findings (e.g., more sophisticated image registration schemes, or changing the

121 degree of spatial smoothing used). In the end, we endorsed the sharing of data in as many
122 forms as is feasible.
123

124 Are we ready for open science in neuroimaging?

125 Brain imaging research is complicated, not only at the level of the conducting a study, but also
126 at the level of sharing its results and data. The importance of thorough reporting of results is
127 uncontroversial, and practices are improving, and the sharing of data to facilitate replication is
128 increasingly viewed as essential. However, data sharing poses new challenges. Here we
129 consider a number of concerns that investigators have with data sharing that impede adoption
130 of open practices.

131
132 First, some individual researchers may assert ownership of their data and thus may not feel
133 compelled to share. Counter to this is the drive for publically funded research to produce widely
134 accessible data that can be reused and integrated into further research. Researchers may feel
135 that sharing of data will result in a loss of competitive advantage, with other researchers
136 swooping in to publish their planned studies based on the same data. The actual risk of this will
137 depend on the data and hypotheses, but it should be weighed against the opportunity of new
138 collaborations resulting from the sharing. These concerns can be alleviated by delaying the
139 sharing or using a data-sharing repository with an embargo period.

140
141 Another fear is that, upon sharing data, other researchers will discover errors in an analysis or
142 previously undiscovered problems with the data. As scientists, we are supposed to be objective
143 arbiters of evidence and theory, but we are not infallible and must be ready to accept criticism
144 and revise our claims when errors are discovered. Even when no errors are found, a re-
145 analyses may support conclusions inconsistent with the original study. For controversial topics,
146 there may also be adversarial reanalyses. We see no better way to advance understanding on a
147 contested finding than to have as many researchers as possible puzzling over the data at hand.
148 However, we need to develop a culture of constructive criticism that recognizes that errors are
149 an inevitable part of scientific progress and protects individual researchers from inappropriately
150 harsh consequences when honest mistakes are discovered.

151
152 A very practical concern, especially for junior investigators, is what is perceived as an
153 unjustifiable cost of data sharing. Current incentives do not justify spending large amounts of
154 time preparing data for sharing, as institutional promotion panels or grant reviewers currently do
155 not adequately reward such efforts. Counter to this is the greater potential impact of a work
156 when it may be cited not just for its scientific findings, but also when its data is reused in other
157 works. Data description papers can document and provide credit for high-quality data
158 acquisition efforts for the open community. We assert that if data sharing and open science
159 priorities in general are to take hold, academic institutions, journals, and granting agencies are
160 crucial for improving the incentives for open practices and developing ways to give appropriate
161 credit for efforts in data sharing.

162

163 Finally there is the very real worry of failing to comply with human ethics provisions for
164 protecting subject privacy. It can be argued that, once file headers are scrubbed of personally
165 identifiable information and structural images have facial features obscured, that the data are
166 completely anonymised and thus freely sharable. However individual ethics boards have varying
167 views on this and it is best to write ethics consent documents explicitly with data sharing in
168 mind. This topic would greatly benefit from leadership from national research organisations to
169 seek consensus and then establish exactly what comprises anonymized brain imaging data. In
170 particular, ethics boards often only try to minimize the risk to subjects when we are also obliged
171 to maximize the benefit of our research to science and society, so as to honor the contribution of
172 our subjects.¹² The future value of shared data must be considered in ethical decision making.

173
174 While studies lacking shared data and having opaque methodological detail are typical, some
175 authors have embraced the challenges of sharing data and analysis methodology. Some recent
176 examples that are particularly thorough and elegant include Waskom et al.¹³ and Whitaker et
177 al.¹⁴, that published a complete array of analysis scripts for generating all figures and results in
178 the paper (https://github.com/mwaskom/Waskom_JNeurosci_2014 and
179 https://github.com/KirstieJane/NSPN_WhitakerVertes_PNAS2016, respectively), and Pernet et
180 al.¹⁵ that likewise shared raw data and analysis scripts, as well as all results maps in electronic
181 form. From an organisational perspective, some labs are simply making open science a policy.
182 Most recently the Montreal Neurological Institute announced that their work would be open, with
183 all results and data made freely available at the time of publication¹⁶.

184
185 These few examples demonstrate that some researchers are embracing open science
186 principles, but do the tools exist to make it practical on a widespread basis?

187 Existing tools for open neuroimaging

188 There is an emerging ecosystem of open science tools for neuroimaging research. Before any
189 data is collected, there are tools to assist in creating human ethics documents that maximise the
190 ease of later data sharing, and for everything from experimental paradigm presentation,
191 preprocessing to statistical modelling, neuroimaging benefits from numerous, free and well-
192 supported software tools (see Supplementary Table 1 for an incomplete list). This constellation
193 of tools could be seen as fuel for limitless researcher degrees-of-freedom, and indeed there is a
194 need for the community to identify a set of ‘reference pipelines’ for common analyses. However,
195 since each tool makes particular assumptions about neuroanatomical and neurophysiological
196 processes, it is not possible to recommend the optimal analyses for every possible type of data
197 and analysis objective. Only with user experience and reproducibility comparisons, will the field
198 be able to identify what are the preferred analytical approaches.

199
200 There is a particular embrace of data sharing in the resting-state fMRI community. Since
201 resting-state analyses methods remain in flux, sharing of this data has particular value as it
202 allows future improvements in methods to be assessed and benchmarked relative to previous
203 analyses. For resting and task fMRI and structural MRI, there are a number of projects that have
204 led the way in this area, including the sibling projects FCON1000 and INDI¹⁷, and the

205 Alzheimer's Disease Neuroimaging Initiative (<http://www.adni-info.org>). These have become
206 invaluable tools for methodologists to apply novel image processing algorithms, not to mention
207 the primary scientific outputs from these projects.

208
209 One promising new standard is the Brain Imaging Data Structure (BIDS)¹⁸, a simple system for
210 organising MRI data after conversion to the NIFTI format. BIDS provides a common, consistent
211 directory hierarchy and naming system for files, as well as supporting 'side car' files for key
212 associated data (like stimulus timing information for task fMRI). With a fixed standard for
213 representing data, this has supported the creation of a number of "BIDS Apps", self-contained
214 programs that can automatically process data arranged according to BIDS. Simple, widely used
215 standards such as this have the potential to dramatically reduce the effort required to exchange
216 and share data.

217
218 New tools are set to dramatically advance computational reproducibility. A challenge to even
219 something as simple as re-running the same data with the same code is the ever-changing
220 versions of software and libraries that software depends on. The last five years has seen the
221 growth of virtual machines and containers to share not just data but a complete environment for
222 processing data. A virtual machine (VM) is an emulator of a computer, including its hardware,
223 operating system and file system. It can be shared as a single file and when run, an entire
224 computer system comes into existence based on a snapshot of the libraries and software
225 interdependencies of one particular system. From within this VM, data can be run through a
226 complete processing pipeline; with the original data of a study this will reproduce the results
227 exactly, while new data can also be imported to evaluate the unique aspects of a pipeline. A
228 downside to VMs is their gross size, as they are as large as any operating system. Containers
229 are miniature VMs, lacking the full operating system but providing the specialised software and
230 libraries required to execute a given task. The BIDS Apps mentioned above rely on such
231 containers, encapsulating software packages large and small that alleviate installation of a
232 myriad of software dependencies.

233
234 Open science tools are gaining traction. For example, the CBRAIN web-based analysis service
235 supports over 260 collaborators in 20 countries; the COINS service currently hosts data on over
236 40,000 subjects for 643 studies; the LONI Pipeline has an average of 100,000 daily jobs from
237 200 different analysis workflows; the Neurovault repository hosts 450 public collections; and the
238 FCP/INDI is openly sharing over 15,000 resting fMRI and structural MRI datasets.

239 Continuous improvement of research practices

240 Despite a seeming wealth of tools, there remain specific areas in the field of neuroimaging that
241 need to be embraced to increase reproducibility. Aside from the importance of carefully
242 reporting the study design, methods, and results mentioned above, we also identified priorities
243 including archiving of statistical results, software engineering for reproducibility, and optimizing
244 projects for generalizability.

245

246 In genetics, the routine sharing of “summary data” (SNP-level statistical results) has facilitated
247 meta-analyses and methodological developments. For example, LD-score regression is a tool
248 that can estimate genetic correlation using just Z-score summary data, and has had dramatic
249 impact in a short timespan due to the availability of such results¹⁹. In brain imaging, we have no
250 tradition of sharing summary statistics (i.e. images of T- or Z-scores, or images of percent
251 change effect and standard errors). As a result the quality of meta-analyses are currently limited
252 by their reliance on reported tables of maximum location coordinates, for which there is a
253 substantial loss of information relative to the original statistic images²⁰. In the current age, the
254 costs of sharing such images of summary statistics (~1MB compressed), either through generic
255 or dedicated repositories (e.g., NeuroVault.org, or BALSAs, <http://balsa.wustl.edu>), are relatively
256 minimal. As such, COBIDAS recommends the deposition of unthresholded statistical images
257 into archival resources for all studies. Widespread adoption of this practice will dramatically
258 increase our capacity for more precise meta-analyses, and allow more critical assessment of
259 study results through exploration of the complete 3D image.

260
261 One foundation of computational reproducibility is modern software engineering practice.
262 Whether a small set of scripts or a comprehensive end-to-end pipeline, neuroimaging data
263 analysis depends on coding. Modern software engineering includes practices like version
264 control and unit testing. Version control ensures that revisions of the code are identifiable and
265 archived, and ideally is based on an open platform that allows wide inspection and input; unit
266 tests verify the correctness of individual facets of the code, and can be set to automatically run
267 each time the code is updated. This is not to say that every group should hire a programmer,
268 but rather that every researcher writing scripts or code should obtain proficiency with basic
269 software engineering skills and practices²¹ (see Software Carpentry for basics instruction for
270 non-programmers, <http://software-carpentry.org/>). With routine research grounded in well-
271 written, less fragile code, it will be much easier to establish analysis pipelines that can both be
272 reused within a lab and facilitate computational reproducibility verified by others.

273
274 Study designs have traditionally been optimised to maximise statistical power to detect
275 differences between groups. With a growing emphasis on prediction, whether (e.g.) identifying
276 early risk for psychosis or progression of a neurodegenerative disease, studies should be
277 optimised for building predictive models that will generalise to the population of interest in yet-
278 unseen data. Large multi-site studies that capture wide variation in human populations, as well
279 as site-specific technical idiosyncrasies, are essential to build classifiers with good performance
280 on new data. Whether obtained with prospectively optimized homogeneous acquisition and
281 preprocessing strategies (e.g. Human Connectome Project and its successors²²) or via larger
282 but more heterogeneous, aggregate multisite approaches (e.g., FCON1000/INDI; ADNI, PING,
283 and the upcoming ABCD Study) that have optimized image processing strategies determined
284 retrospectively²³, generalisability of predictive models will be a key design objective and
285 performance indicator going forward.

286 Beyond the investigator

287 Many of the practices advocated here and in the full COBIDAS MRI report require individuals to
288 change the way they conduct research. Almost every such change requires an investment of
289 time and resources. While we argue these have implicit rewards (e.g. shared data will never be
290 lost when the post doc moves on), the advance of open science will require leadership at the
291 institutional level. To provide appropriate incentives, universities and research centers need to
292 explicitly consider the value of sharing of data and code as an unique research output in
293 promotion and review exercises. Journals should require that papers' statistic images are
294 archived, and promote papers with shared data, provide full analytical detail, and ideally share
295 code or even executable containers or VMs. Foundations and granting agencies need to make
296 data sharing a priority, recognizing and funding the explicit costs of data management required
297 to make this happen. And finally professional organisations like OHBM should prioritize efforts in
298 education to make open science practices accessible to all.

299
300 With the coordinated efforts of individual researchers, academic institutions, journals, granting
301 agencies, and professional organisations, we can accelerate the drive towards open science
302 and maximise the reproducibility of neuroimaging findings going forward.

303
304
305
306

307 Acknowledgements.

308 TEN is supported by the Wellcome Trust (100309/Z/12/Z) and NIH (R01 NS075066-01A1, R01
309 EB015611-01). BTTY is supported by Singapore MOE Tier 2 (MOE2014-T2-2-016), NUS
310 (DPRT/944/09/14, R185000271720), NMRC (CBRG14nov007) and the NUS YIA. SBE is
311 supported by the Deutsche Forschungsgemeinschaft (DFG, EI 816/4-1, LA 3071/3-1; EI 816/6-
312 1), the National Institute of Mental Health (R01-MH074457), the Helmholtz Portfolio Theme
313 "Supercomputing and Modeling for the Human Brain" and the European Union Seventh
314 Framework Programme (FP7/2007-2013) under grant agreement no. 604102). MH was
315 supported by funds from the German federal state of Saxony-Anhalt and the European Regional
316 Development Fund (ERDF), project: Center for Behavioral Brain Sciences, and CRCNS
317 BMBF/NSF (01GQ1411/1429999). NK was supported by the UK Medical Research Council
318 and a European Research Council Starting Grant (261352). The authors declare no competing
319 financial interests. ACE, SD and TG are supported by the Irving Ludmer Family Foundation and
320 the Ludmer Centre for Neuroinformatics and Mental Health. TW was supported by a ZonMw
321 TOP grant (91211021). RAP is supported by the Laura and John Arnold Foundation. MPM is a
322 Phyllis Green and Randolph Cowen Scholar and is supported in part by the NIH (U01
323 MH099059; R01 AG047596)

324
325 REFERENCES

326

- 327 1. Ioannidis, J. P. A. (2005). Why most published research findings are false. *PLoS*
328 *Medicine*, 2(8), e124. doi:10.1371/journal.pmed.0020124
- 329 2. Open Science Collaboration. (2015). PSYCHOLOGY. Estimating the reproducibility of
330 psychological science. *Science (New York, N.Y.)*, 349(6251), aac4716.
331 doi:10.1126/science.aac4716
- 332 3. Journals unite for reproducibility. (2014). *Nature*, 515(7525), 7. doi:10.1038/515007a
- 333 4. Nichols, T. E., Das, S., Eickhoff, S. B., Evans, A. C., Glatard, T., Hanke, M., Yeo, B.
334 T. T. (2016). Best Practices in Data Analysis and Sharing in Neuroimaging using MRI.
335 bioRxiv, 54262. <http://doi.org/10.1101/054262>
- 336 5. Peng, R. D. (2011). Reproducible research in computational science. *Science*,
337 334(6060), 1226–7. doi:10.1126/science.1213847
- 338 6. Bennett, C. M., & Miller, M. B. (2013). fMRI reliability: influences of task and
339 experimental design. *Cognitive, Affective & Behavioral Neuroscience*, 13(4), 690–702.
340 <http://doi.org/10.3758/s13415-013-0195-1>
- 341 7. Schnack, H. G., van Haren, N. E. M., Brouwer, R. M., van Baal, G. C. M., Picchioni, M.,
342 Weisbrod, M., Hulshoff Pol, H. E. (2010). Mapping reliability in multicenter MRI: voxel-
343 based morphometry and cortical thickness. *Human Brain Mapping*, 31(12), 1967–82.
344 <http://doi.org/10.1002/hbm.20991>
- 345 8. Noble, S., Scheinost, D., Finn, E. S., Shen, X., Papademetris, X., McEwen, S. C.,
346 Constable, R. T. (2016). Multisite Reliability of MR-Based Functional Connectivity.
347 *NeuroImage*. <http://doi.org/10.1016/j.neuroimage.2016.10.020>
- 348 9. Boebel, W., Wagenmakers, E.-J., Belay, L., Verhagen, J., Brown, S., & Forstmann, B. U.
349 (2015). A purely confirmatory replication study of structural brain-behavior correlations.
350 *Cortex*, 1–19. <http://doi.org/10.1016/j.cortex.2014.11.019>
- 351 10. Kriegeskorte, N., Simmons, W. K., Bellgowan, P. S. F., & Baker, C. I. (2009). Circular
352 analysis in systems neuroscience: the dangers of double dipping. *Nature Neuroscience*,
353 12(5), 535–40. doi:10.1038/nn.2303
- 354 11. Poldrack, R. A., Fletcher, P. C., Henson, R. N., Worsley, K. J., Brett, M., & Nichols, T. E.
355 (2008). Guidelines for reporting an fMRI study. *NeuroImage*, 40(2), 409–14.
356 <http://doi.org/10.1016/j.neuroimage.2007.11.048>
- 357 12. Brakewood, B., & Poldrack, R. A. (2013). The ethics of secondary data analysis:
358 Considering the application of Belmont principles to the sharing of neuroimaging data.
359 *NeuroImage*, 82, 671–676. <http://doi.org/10.1016/j.neuroimage.2013.02.040>
- 360 13. Waskom, M. L., Kumaran, D., Gordon, A. M., Rissman, J., & Wagner, A. D. (2014).
361 Frontoparietal Representations of Task Context Support the Flexible Control of Goal-
362 Directed Cognition. *Journal of Neuroscience*, 34(32), 10743–10755. JOUR.
363 <http://doi.org/10.1523/JNEUROSCI.5282-13.2014>
- 364 14. Whitaker, K. J., Vértes, P. E., Romero-Garcia, R., Váša, F., Moutoussis, M., Prabhu, G.,
365 Bullmore, E. T. (2016). Adolescence is associated with genomically patterned
366 consolidation of the hubs of the human brain connectome. *Proceedings of the National*
367 *Academy of Sciences*, 113(32), 201601745. <http://doi.org/10.1073/pnas.1601745113>
- 368 15. Pernet, C. R., McAleer, P., Latinus, M., Gorgolewski, K. J., Charest, I., Bestelmeyer, P.
369 E. G., Belin, P. (2015). The human voice areas: Spatial organization and inter-

- 370 individual variability in temporal and extra-temporal cortices. *NeuroImage*, 119, 164–74.
371 <http://doi.org/10.1016/j.neuroimage.2015.06.050>
- 372 16. Owens, B. (2016). Montreal institute going “open” to accelerate science. *Science*.
373 <http://doi.org/10.1126/science.aae0265>
- 374 17. Mennes, M., Biswal, B. B., Castellanos, F. X., & Milham, M. P. (2013). Making data
375 sharing work: The FCP/INDI experience. *NeuroImage*, 82, 683–691.
376 <http://doi.org/10.1016/j.neuroimage.2012.10.064>
- 377 18. Gorgolewski, K. J., Auer, T., Calhoun, V. D., Craddock, R. C., Das, S., Duff, E. P.,
378 Poldrack, R. A. (2016). The brain imaging data structure, a format for organizing and
379 describing outputs of neuroimaging experiments. *Scientific Data*, 3, 160044.
380 <http://doi.org/10.1038/sdata.2016.44>
- 381 19. Bulik-Sullivan, B. K., Loh, P.-R., Finucane, H. K., Ripke, S., Yang, J., Patterson, N.,
382 Neale, B. M. (2015). LD Score regression distinguishes confounding from polygenicity in
383 genome-wide association studies. *Nature Genetics*, 47(3), 291–295.
384 <http://doi.org/10.1038/ng.3211>
- 385 20. Salimi-Khorshidi, G., Smith, S. M., Keltner, J. R., Wager, T. D., & Nichols, T. E. (2009).
386 Meta-analysis of neuroimaging data: A comparison of image-based and coordinate-
387 based pooling of studies. *NeuroImage*, 45(3), 810–823.
388 <http://doi.org/10.1016/j.neuroimage.2008.12.039>
- 389 21. Goble, C. (2014). Better software, better research. *IEEE Internet Computing*, 18(5), 4–8.
390 <http://doi.org/10.1109/MIC.2014.88>
- 391 22. Glasser, M. F., Smith, S. M., Marcus, D. S., Andersson, J. L. R., Auerbach, E. J.,
392 Behrens, T. E. J., Van Essen, D. C. (2016). The Human Connectome Project’s
393 neuroimaging approach. *Nature Neuroscience*, 19(9), 1175–87.
394 <http://doi.org/10.1038/nn.4361>
- 395 23. Abraham, A., Milham, M., Martino, A. D., Craddock, R. C., Samaras, D., Thirion, B., &
396 Varoquaux, G. (2016). Deriving reproducible biomarkers from multi-site resting-state
397 data: An Autism-based example. *Neuroimage*.
398 <http://doi.org/10.1016/j.neuroimage.2016.10.045>
- 399 24. ISO. (2006). *Statistics - Vocabulary and symbols. Part 2: Applied statistics. ISO 3534–2*
400 (Second.). Geneva: ISO.
401
402

Table 1. A partial taxonomy of reproducibility in neuroimaging. For each type of reproducibility (row), the variable (column) that is held constant (•, bullet) or allowed to vary (D=different) is indicated; minus (-) indicates not applicable. Variations in the participant studied can be described in terms of the population they belong to (e.g. different patient groups or people from different cultures), or whether the same sample or a distinct sample of individuals is used. The MRI scanner used can be the same or not, and if the same participant sample is considered, the very same data can be used or new data can be acquired on the same or different days (visits) to the scanner. Experimental variation has many forms including the particular experimental design, but here we only consider stimuli. The type of stimulus used (stimulus population) may change, for example in a working memory experiment, letter stimuli might be replaced with shape stimuli; a more subtle change would be to use a different sample of stimuli of the same type, e.g. different particular shapes. The analysis method may vary; for example, with structural MRI for prediction of patient disease status, a linear discriminant might be used instead of a nonlinear support vector machine. Analysis code more narrowly reflects the particular implementation of a given method. Personnel conducting the research is another important source of variation, whether this is the experimenter or data analyst. Finally, note that the International Standards Organisation (ISO) has precise definitions of reproducibility²⁴ as indicated in the first three rows, but these capture only the minimal levels of generalizability.

Levels of generalization	Participants		MRI Acquisition			Experiment		Analysis		Personnel	
	Population	Sample	Scanner	Visit	Data	Stimulus Population	Stimulus Sample	Method	Code	Experimenter	Data Analyst
Generalization over measurements											
ISO Repeatability e.g. 30-minute intra-scanner reliability	•	•	•	•	D	•	•	•	•	•	•
ISO Intermediate Reproducibility e.g. 7-day intra-scanner reliability	•	•	•	D	D	•	•	•	•	•	•
ISO Reproducibility e.g. 7-day inter-scanner reliability	•	•	D	D	D	•	•	•	•	•	•
Generalization over analyses											
Analysis Replicability	•	•	•	•	•	•	•	•	•	•	•
Collegial Analysis Replicability	•	•	•	•	•	•	•	•	•	•	D
Peng ⁵ Reproducibility	•	•	•	•	•	•	•	•	D	D	D
Generalization over materials and methods											
Near Replicability (different subjects)	•	D	•	-	-	•	•	•	•	•	•
Intermediate Replicability (different labs)	•	D	D	-	-	•	•	•	•	D	D
Far Replicability (different experimental & analytical methods)	•	D	D	-	-	•	D	D	D	D	D
Hypothesis Generalisability (different subject populations & types of stimuli)	D	D	D	-	-	D	D	D	D	D	D

Supplementary Table 1. An incomplete but illustrative list of free and well-supported tools for open science tools for neuroimaging. This table highlights analysis tools that can be scripted, allowing replicable analyses, as well as pipeline environments that bind together different software for replicable analyses, across heterogeneous software tools. The items under Data Sharing focus on tools to facilitate sharing and repositories that accept data. As repositories can have varying cost structures depending on the scale of data to be shared, we did not attempt classify as “free” or not; likewise, repositories generally do not comprise software that need to be downloaded, and we likewise did not attempt to classify by open source nature of the project. Results sharing tools either facilitate sharing or serve as repositories for shared results data. The Reproducibility Tools are a loose collection of resources that facilitate research using open science methods.

Resource	Type	Short Description	Free	Open Source	Link
Open Brain Consent	Consent	Ethics template oriented for neuroimaging data sharing	x	x	http://open-brain-consent.readthedocs.io
OpenSesame	Paradigm software	Graphical experiment builder	x	x	http://osdoc.cogsci.nl
PsychoPy	Paradigm software	Psychophysics software in Python	x	x	http://www.psychopy.org
Psychtoolbox	Paradigm software	Psychophysics Toolbox	x	x	http://psychtoolbox.org/
aa	Pipeline	Automatic Analysis, Matlab-based workflow tool	x (Matlab)	x	http://automaticanalysis.org
C-BRAIN	Pipeline	Web-based software for computationally intensive analyses	x	x	http://cbrain.mcgill.ca
CCS	Pipeline	Connectome Computation System, a pipeline primarily for resting data	x	x	http://github.com/zuoxinian/CCS
C-PAC	Pipeline	Configurable Pipeline for the Analysis of Connectomes	x	x	http://fcp-indi.github.io
DPARSF/DPABI	Pipeline	Data Processing & Analysis for Brain Imaging, including resting-state fMRI	x	x	http://rfmri.org/dpabi
DTIPrep	Pipeline	Pipeline for diffusion weighted / diffusion tensor image data	x	x	http://www.nitrc.org/projects/dtiprep/
HCP Pipeline	Pipeline	Human Connectome Project Pipeline	x	x	http://github.com/Washington-University/Pipelines
LONI Pipeline	Pipeline	Cross-platform workflow tool for neuroimaging, genomics, bioinformatics	NC		http://pipeline.loni.usc.edu
LORIS	Pipeline	Web-based data and project management software for neuroimaging	x	x	http://loris.ca
NIAC	Pipeline	Library of modules and pipelines for fMRI processing in Matlab/Octave	x	x	http://www.nitrc.org/projects/niac
NiDB	Pipeline	Neuroimaging database software that includes pipeline tools	x	x	http://github.com/gbook/nidb
NiPype	Pipeline	Neuroimaging in Python Pipelines and Interfaces	x	x	http://nipy.org/nipype
PANDA	Pipeline	Pipeline for Analyzing brain Diffusion images	x	x	http://www.nitrc.org/projects/panda
SimNIBS	Pipeline	Simulation of Non-invasive Brain Stimulation	x	x	http://simnibs.de
AFNI	Scriptable Analysis	Neuroimaging analysis software for functional MRI	x	x	http://afni.nimh.nih.gov/afni
CONN	Scriptable Analysis	Functional connectivity toolbox, Matlab-based pipeline tool	x (Matlab)	x	http://www.nitrc.org/projects/conn
Connectir	Scriptable Analysis	Analysis software for Connectome-Wide Association Studies, based in R	x	x	http://czarrar.github.io/connectir
DiPy	Scriptable Analysis	Diffusion analysis pipeline using Python	x	x	http://nipy.org/dipy
Freesurfer	Scriptable Analysis	Neuroimaging analysis software for MRI, emphasis on surface-based analysis	x	x	http://surfer.nmr.mgh.harvard.edu
FSL	Scriptable Analysis	Neuroimaging analysis software for MRI	NC	x	http://www.fmrib.ox.ac.uk/fsl
MindBoggle	Scriptable Analysis	Automated labeling and shape analysis of brain images	x	x	http://www.mindboggle.info
SPM	Scriptable Analysis	Neuroimaging analysis software based in Matlab, for MRI, M/EEG, PET.	x (Matlab)	x	http://www.fil.ion.ucl.ac.uk/spm
Voxel	Scriptable Analysis	Mass-Univariate Voxelwise Analysis of Medical Imaging Data, based in R	x	x	http://cran.r-project.org/web/packages/voxel
BIDS	Data Sharing	Standard for organising MRI data and associated supporting data			http://bids.neuroimaging.io
COINS	Data Sharing	Web-based data management and analysis tool			http://coins.mrn.org
FCP/INDI	Data Sharing	Repository for resting state fMRI data			http://fcon_1000.projects.nitrc.org
Figshare	Data Sharing	Generic data sharing repository			http://figshare.com
LONI IDA	Data Sharing	Image data archive, repository for primarily neuroimaging data			http://ida.loni.usc.edu
LORIS	Data Sharing	Database for longitudinal imaging studies			http://bigbrain.loris.ca
NDA	Data Sharing	NIMH Data Archive, repository for data from NIMH-funded studies			http://data-archive.nimh.nih.gov
NITRC-IR	Data Sharing	Image repository for neuroimaging data			http://www.nitrc.org/ir
OpenfMRI	Data Sharing	Repository for task fMRI data, including all image and task paradigm data			https://openfMRI.org
PCP	Data Sharing	Preprocessed connectome project - pipelines for resting state data			http://preprocessed-connectomes-project.org
XNAT-Central	Data Sharing	Repository for raw MRI data			http://central.xnat.org
BALSA	Results Sharing	Sharing of surface-based statistical results	x		http://balsa.wustl.edu
NeuroVault	Results Sharing	Sharing tool for statistical maps	x	x	http://neurovault.org
NIDM	Results Sharing	Standard for exporting statistical results independent of the analysis tool	x	x	http://nidm.nidash.org
Docker	Reproducibility tool	Containerisation tool	x	x	http://www.docker.com
GitHub	Reproducibility tool	Version and issue tracking for software projects	x	x	http://github.org

Resource	Type	Short Description	Free	Open Source	Link
NeuroDebian	Reproducibility tool	Archive of research software packages for use on workstations & VMs	x	x	http://neuro.debian.net