Editorial

Introduction to the special issue on reproducibility in neuroimaging

The last decade has seen increasing attention to the problem of scientific reproducibility, across a broad range of scientific fields (Camerer et al., 2016; Morrison, 2014; Open Science Collaboration, 2015). Within the field of neuroimaging, there has been a particular focus on issues of analytic variability (Bowring et al., 2019; Carp, 2012) statistical power (Button et al., 2013; Poldrack et al., 2017), and test-retest reliability (Bennett and Miller, 2013), all of which have raised alarms regarding the potential for irreproducible results. In addition, failed replications (Boekel et al., 2015; Dinga et al., 2019) and meta-analytic null results (Müller et al., 2017) have raised particular concern about studies of group and individual differences. This special issue was developed in light of these emerging concerns, with the goal of highlighting and encouraging work that aims to both quantify and improve the reproducibility of neuroimaging research. Here we provide a brief overview of the papers within this special issue.

The terms “reproducibility”, “replicability”, and “reliability” are used in variable ways by many groups (Nichols et al., 2017). Here we use “reproducibility” as a blanket term encompassing all aspects of the ability to reproduce a result, from same data/same analysis to different data/different analysis. We use the term “replication” to refer specifically to the ability for a finding to be reproduced - qualitatively found again - in a separate dataset. By “reliability” we specifically mean the degree to which a measurement is stable across multiple repeated measurements.

1. Quantifying reliability

A number of the papers submitted for this special issue focused on quantifying the test-retest reliability of neuroimaging measurements across repeated measures. Several of these focused on functional connectivity using fMRI. Nobel et al. (this issue) performed a systematic review that combined results across 25 published studies of test-retest reliability of edge-level functional connectivity, which showed poor reliability on average and highlighted a number of features that promoted greater reliability. Badhwar et al. (this issue) took the opposite approach, assessing connectivity in a single individual across multiple sites and scanner vendors. This effort showed reasonable reliability that was moderated by both site and vendor effects, highlighting possible limitations on between-site generalizability. Baria et al. (this issue) assessed the reliability of resting state fMRI “fingerprinting” (Finn et al., 2015), finding that an optimal reconstruction method led to higher fingerprinting performance both within and across sites. Tu et al. (this issue) also assessed the ability to fingerprint individuals across multiple sessions, replicating previous successful fingerprinting results. They further showed that connectivity could be used to predict pain thresholds both within and across sessions.

Other papers assessed the reliability of structural MRI measures. Buonincontri et al. (this issue) assessed the test-retest reliability as well as the between-site reproducibility of a method for reconstructing multiple parametric maps from a single excitation (known as “MR fingerprinting”, not be confused with “connectome fingerprinting”). They report high levels of test-retest reliability as well as acceptable levels of between-site reproducibility. Drenthen et al. (this issue) assessed the retest reliability of multi-slice GRASE for use in quantitative myelin water imaging. They validated the multi-slice measures in relation to the standard whole-brain GRASE method, and further showed that this method achieved acceptable reliability even using parallel acceleration to substantially reduce scan time. Lancione et al. (this issue) assessed the specific effect of echo time (TE) on reproducibility of quantitative susceptibility mapping across 3T and 7T scanners. Their results show that it is possible to generate reproducible maps across scanners by calibrating the echo time across scanners. Lerma-Usubaig et al. (this issue) assessed the utility of fractional anisotropy in homologous pairs of tracts as a diagnostic metric across nine datasets, showing that this metric was substantially more precise compared to standard measures of diffusion anisotropy.

2. Replication

The last decade has seen increasing interest in replication studies, but it remains difficult to publish such studies. For this reason, we solicited direct replication studies as part of our call for papers, ultimately including two such studies. Geller et al. (this issue) attempted to replicate a previous finding that had shown that binarized measures of tract connection were strongly associated with post-stroke language function compared to a continuous measure of lesion load. They failed to find an over difference between the binary and continuous measures. Kampa et al. (this issue) assessed replicability between two samples within a single fMRI study of stress resilience, finding that replicability at the whole-brain level was good whereas replicability at the level of individual regions of interest was lower.

Other studies examined the nature of replication and the factors that affect it. Hong et al. (this issue) surveyed studies that had claimed to replicate previous findings, in order to assess what was meant by “replication”. They found that most studies did not provide quantitative evidence for anatomical replication, and that many of the reported peaks were distant from the original peak, suggesting that claimed replication does not always imply true replication. Xia et al. (this issue) examined the reproducibility of case-control differences in major depressive disorder using resting fMRI. They found reliable differences between groups in several measures, but these measures were only reproducible when relatively large samples were included in the analysis. Jollans et al. (this issue) assessed quantitative and qualitative reproducibility of task fMRI studies using a combination of functional and structural MRI measures. They performed three within-group analyses and two independent between-group analyses, with the most stringent protocol showing acceptable inter-site reliability in the whole brain and high within-site reliability in several regions. Finally, they showed that the combination of functional and structural MRI measures increased the reproducibility of the analysis.

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3. Confounds

A final set of papers focused on effects of confounds in neuroimaging analysis. Quax et al. focused on a specific confound in decoding of stimulus location from MEG data due to the effects of eye movements. Their results suggest that controlling for the confounding effects of eye movements is essential in any MEG decoding study. Hyatt et al. reviewed the use of covariates in structural neuroimaging studies, finding a broad range of different strategies used across the literature. In an analysis of data from the Human Connectome Project, they found that some covariates had strong effects on outcomes whereas the effects of others were minimal. They highlight the need for pre-registration to ensure that flexibility in confound modeling does not inflate error rates.

4. Conclusion

Reproducibility is a multi-faceted and ongoing concern for neuroimaging researchers. The papers in this special issue highlight the different ways that the field is tackling these issues head on. What particularly ties these articles together is a drive to efficiently and effectively utilize neuroimaging research to better understand the human mind and brain in health and disease, and across the lifespan. We hope that they inspire additional work aimed at assessing and improving reproducibility.

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