# Transparency and reproducibility in artificial intelligence

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ARISING FROM S. M. McKinney et al. Nature https://doi.org/10.1038/s41586-019-1799-6 (2020)

Breakthroughs in artificial intelligence (AI) hold enormous potential as it can automate complex tasks and go even beyond human performance. In their study, McKinney et al.<sup>1</sup> showed the high potential of AI for breast cancer screening. However, the lack of details of the methods and algorithm code undermines its scientific value. Here, we identify obstacles that hinder transparent and reproducible AI research as faced by McKinney et al.<sup>1</sup>, and provide solutions to these obstacles with implications for the broader field.

The work by McKinney et al.<sup>1</sup> demonstrates the potential of AI in medical imaging, while highlighting the challenges of making such work reproducible. The authors assert that their system improves the speed and robustness of breast cancer screening, generalizes to populations beyond those used for training, and outperforms radiologists in specific settings. Upon successful prospective clinical validation and approval by regulatory bodies, this new system holds great potential for streamlining clinical workflows, reducing false positives, and improving patient outcomes. However, the absence of sufficiently documented methods and computer code underlying the study effectively undermines its scientific value. This shortcoming limits the evidence required for others to prospectively validate and clinically implement such technologies. By identifying obstacles hindering transparent and reproducible AI research as faced by McKinney et al.<sup>1</sup>, we provide potential solutions with implications for the broader field.

Scientific progress depends on the ability of independent researchers to scrutinize the results of a research study, to reproduce the study's main results using its materials, and to build on them in future studies (https://www.nature.com/nature-research/editorial-policies/ reporting-standards). Publication of insufficiently documented research does not meet the core requirements underlying scientific discovery<sup>2,3</sup>. Merely textual descriptions of deep-learning models can hide their high level of complexity. Nuances in the computer code may have marked effects on the training and evaluation of results<sup>4</sup>, potentially leading to unintended consequences<sup>5</sup>. Therefore, transparency in the form of the actual computer code used to train a model and arrive at its final set of parameters is essential for research reproducibility. McKinney et al.<sup>1</sup> stated that the code used for training the models has "a large number of dependencies on internal tooling, infrastructure and hardware", and claimed that the release of the code was therefore not possible. Computational reproducibility is indispensable for high-quality AI applications<sup>6,7</sup>; more complex methods demand greater transparency<sup>8</sup>. In the absence of code, reproducibility falls back on replicating methods from textual description. Although, McKinney and colleagues1 claim that all experiments and implementation details were described in sufficient detail in the supplementary methods section of their Article<sup>1</sup> to "support replication with non-proprietary libraries", key details about their analysis are lacking. Even with extensive description, reproducing complex computational pipelines based purely on text is a subjective and challenging task<sup>9</sup>.

In addition to the reproducibility challenges inherent to purely textual descriptions of methods, the description by McKinney et al.<sup>1</sup> of the model development as well as data processing and training pipelines lacks crucial details. The definitions of several hyperparameters for the model's architecture (composed of three networks referred to as the breast, lesion and case models) are missing (Table 1). In their publication, McKinney et al.<sup>1</sup> did not disclose the settings for the augmentation pipeline; the transformations used are stochastic and can considerably affect model performance<sup>10</sup>. Details of the training pipeline were also missing. Without this key information, independent reproduction of the training pipeline is not possible.

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### **Matters arising**

## Table 1 | Essential hyperparameters for reproducing the study for each of the three models

	Lesion	Breast	Case
Learning rate	Missing	0.0001	Missing
Learning rate schedule	Missing	Stated	Missing
Optimizer	Stochastic gradient descent with momentum	Adam	Missing
Momentum	Missing	Not applicable	Not applicable
Batch size	4	Unclear	2
Epochs	Missing	120,000	Missing

Numerous frameworks and platforms exist to make artificial intelligence research more transparent and reproducible (Table 2). For the sharing of code, these include Bitbucket, GitHub and GitLab, among others. The many software dependencies of large-scale machine learning applications require appropriate control of the software environment, which can be achieved through package managers including Conda, as well as container and virtualization systems, including Code Ocean, Gigantum, Colaboratory and Docker. If virtualization of the McKinney et al.<sup>1</sup> internal tooling proved to be difficult, they could have released the computer code and documentation. The authors could have also created small artificial examples or used small public datasets<sup>11</sup> to show how new data must be processed to train the model and generate predictions. Sharing the fitted model (architecture along with learned parameters) should be simple aside from privacy concerns that the model may reveal sensitive information about the set of patients used to train it. Nevertheless, techniques for achieving differential privacy exist to alleviate such concerns. Many platforms allow sharing of deep learning models, including TensorFlow Hub, ModelHub. ai, ModelDepot and Model Zoo with support for several frameworks such as PyTorch and Caffe, as well as the TensorFlow library used by the authors. In addition to improving accessibility and transparency, such resources can considerably accelerate model development, validation and transition into production and clinical implementation.

Another crucial aspect of ensuring reproducibility lies in access to the data the models were derived from. In their study, McKinney et al.<sup>1</sup> used two large datasets under license, properly disclosing this limitation in their publication. The sharing of patient health information is highly regulated owing to privacy concerns. Despite these challenges, the sharing of raw data has become more common in biomedical literature, increasing from under 1% in the early 2000s to 20% today<sup>12</sup>. However, if the data cannot be shared, the model predictions and data labels themselves should be released, allowing further statistical analyses. Above all, concerns about data privacy should not be used as a way to distract from the requirement to release code.

Although sharing of code and data are widely seen as a crucial part of scientific research, the adoption varies across fields. In fields such as genomics, complex computational pipelines and sensitive datasets have been shared for decades<sup>13</sup>. Guidelines related to genomic data are clear, detailed and, most importantly, enforced. It is generally accepted that all code and data are released alongside a publication. In other fields of medicine and science as a whole, this is much less common, and data and code are rarely made available. For scientific efforts in which a clinical application is envisioned and human lives would be at stake, we argue that the bar of transparency should be set even higher. If a dataset cannot be shared with the entire scientific community, because of licensing or other insurmountable issues, at a minimum a mechanism should be set so that some highly-trained, independent investigators can access the data and verify the analyses.

The lack of access to code and data in prominent scientific publications may lead to unwarranted and even potentially harmful clinical trials<sup>14</sup>. These unfortunate lessons have not been lost on journal editors

### Table 2 | Frameworks to share code, software dependencies and deep-learning models

Resource	URL	
Code		
BitBucket	https://bitbucket.org	
GitHub	https://github.com	
GitLab	https://about.gitlab.com	
Software dependencies		
Conda	https://conda.io	
Code Ocean	https://codeocean.com	
Gigantum	https://gigantum.com	
Colaboratory	https://colab.research.google.com	
Deep-learning models		
TensorFlow Hub	https://www.tensorflow.org/hub	
ModelHub	http://modelhub.ai	
ModelDepot	https://modeldepot.io	
Model Zoo	https://modelzoo.co	
Deep-learning frameworks		
TensorFlow	https://www.tensorflow.org/	
Caffe	https://caffe.berkeleyvision.org/	
PyTorch	https://pytorch.org/	

and their readers. Journals have an obligation to hold authors to the standards of reproducibility that benefit not only other researchers, but also the authors themselves. Making one's methods reproducible may surface biases or shortcomings to authors before publication<sup>5</sup>. Preventing external validation of a model will likely reduce its impact, as it also prevents other researchers from using and building upon it in future studies. The failure of McKinney et al. to share key materials and information transforms their work from a scientific publication open to verification and adoption by the scientific community into a promotion of a closed technology.

We have high hopes for the utility of AI methods in medicine. Ensuring that these methods meet their potential, however, requires that these studies be scientifically reproducible. The recent advances in computational virtualization and AI frameworks are greatly facilitating the implementations of complex deep neural networks in a more structured, transparent, and reproducible way. Adoption of these technologies will increase the impact of published deep-learning algorithms and accelerate the translation of these methods into clinical settings.

### **Reporting summary**

Further information on research design is available in the Nature Research Reporting Summary linked to this paper.

### Data availability

No data have been generated as part of this manuscript.

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Competing interests A.H. is a shareholder of and receives consulting fees from Altis Labs. M.M.H. received a GPU Grant from Nvidia. H.J.W.L.A. is a shareholder of and receives consulting fees from Onc.Al. B.H.K. is a scientific advisor for Altis Labs. C.M. holds an equity position in Bridge7Oncology and receives royalties from RaySearch Laboratories. A.K. is on the SAB of ImmuneAl Inc, a consultant for Biogen Inc., a scientific co-founder of RavelBio Inc. and a shareholder of Freenome Inc. G.A.A., F.K., L.W., B.W., C.S.G., J.T.L., W.H., A.B., J.P., R.T., T.H., J.P.A.I. and J.Q. declare no other competing interests related to the manuscript.

### Additional information

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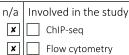
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