

LETTER RESPONSE

Open Access

Response to “Avoiding Methodological Bias in Studies of Amyloid Imaging Results Disclosure”



Joshua D. Grill^{1,2,3,4*} , Chelsea G. Cox¹, Kristin Harkins^{5,6} and Jason Karlawish^{5,6,7,8}

The goal of “Reactions to learning a ‘not elevated’ amyloid PET result in a preclinical Alzheimer’s disease trial” was to study how learning one is not eligible for a trial based on an Alzheimer’s disease (AD) biomarker result affects willingness to be in subsequent trials, as well as how it affects other behaviors [1]. Answering this question fills a critical gap in the literature, as preclinical AD trials are increasingly common but the ideal criteria for participant inclusion remains an area of active research. Thus, a person ineligible for one trial may be eligible for another.

Taswell and colleagues correctly observe that our study did not include a comparison group, which would have necessarily been individuals who demonstrated elevated amyloid and therefore were eligible for randomization in the preclinical AD trial [2]. The primary question under study—whether a subject who screen-fails for one AD trial is willing to participate in subsequent trials—was not applicable to this population. Their exclusion is not a bias. It was not necessary.

Looking more broadly, the gist of Taswell and colleagues’ commentary is a sensible summary of social science research. Multiple methods, and indeed multiple studies, are needed to arrive at a common understanding of the way the world is and how it can be changed. Only then can high confidence be achieved. In the nascent space of preclinical AD trials, much remains to be learned and a breadth of research designs and methods will be essential to advancing the field.

Acknowledgements

There are no acknowledgements.

Funding

There was no funding for this letter.

* Correspondence: jgrill@uci.edu

¹Institute for Memory Impairments and Neurological Disorders, Irvine, CA, USA

²Institute for Clinical and Translational Science, Irvine, CA, USA

Full list of author information is available at the end of the article

Availability of data and materials

No data were involved in this letter.

Authors’ contributions

Each author contributed to the writing of this letter. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Ethics approval was not necessary for this letter.

Consent for publication

We provide consent for publication.

Competing interests

The authors declare that they have no competing interests.

Publisher’s Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

¹Institute for Memory Impairments and Neurological Disorders, Irvine, CA, USA. ²Institute for Clinical and Translational Science, Irvine, CA, USA.

³Department of Psychiatry and Human Behavior, Irvine, CA, USA.

⁴Department of Neurobiology and Behavior, University of California, Irvine, CA, USA.

⁵Penn Memory Center, Philadelphia, PA, USA. ⁶Department of Medicine, Philadelphia, PA, USA.

⁷Department of Medical Ethics and Health Policy, Philadelphia, PA, USA. ⁸Department of Neurology, University of Pennsylvania, Philadelphia, PA, USA.

⁸Department of Neurology, University of Pennsylvania, Philadelphia, PA, USA.

Received: 30 April 2019 Accepted: 13 May 2019

Published online: 04 June 2019

References

1. Grill JD et al. Reactions to learning a “not elevated” amyloid PET result in a preclinical Alzheimer’s disease trial. *Alzheimers Res Ther* 2018;10:125
2. Taswell et al. Avoiding methodological bias in studies of amyloid imaging results disclosure. *Alzheimers Res Ther* 2019. <https://doi.org/10.1186/s13195-019-0495-y>

