Accepted Manuscript

The consequence of "doing nothing": Family caregiving for Alzheimer's disease as non-action in the US

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PII: S0277-9536(17)30731-1

DOI: 10.1016/j.socscimed.2017.12.001

Reference: SSM 11532

To appear in: Social Science & Medicine

Received Date: 23 February 2017
Revised Date: 27 November 2017
Accepted Date: 1 December 2017

Please cite this article as: Seaman, A.T., The consequence of "doing nothing": Family caregiving for Alzheimer's disease as non-action in the US, *Social Science & Medicine* (2018), doi: 10.1016/j.socscimed.2017.12.001.

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ACCEPTED MANUSCRIPT

Manuscript Number: SSM-D-17-00542R2

Title:

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Acknowledgements:

Funding for this research was made possible by the National Science Foundation and the Wenner Gren Foundation for Anthropological Research. Analysis and writing were supported by the University of Chicago's Social Sciences Division. Early versions were presented at meetings of the American Anthropological Association's and the Society for Cultural Anthropology, where Janet Carsten and Sarah Lamb provided invaluable feedback. I am indebted to Elana Buch, Kimberly Dukes, Brady G'sell, Laura Heinemann, Richard Hoffman, Julia Kowalski, Heather Reisinger, Jessica Robbins-Ruskowski, Samantha Solimeo, Kristin Yarris, and members of the Ethnographic Methods and Implementation Core (EMIC), Center for Comprehensive Access & Delivery Research and Evaluation (CADRE), Iowa City VA Health Care System (Iowa City, IA), for their extensive engagement with this work. I am grateful to the editorial staff and anonymous reviewers, who offered incredibly helpful comments on earlier versions of this paper. Finally, I am unable to acknowledge fully enough the people at the center of this research, who found time to foster yet another relational endeavor in the midst of everything else they were doing. Any remaining errors are my own.

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