



Meghan Halley

Sr Research Scholar, School of Medicine - Biomedical Ethics

Bio

BIO

Meghan Halley, PhD, MPH, is a Senior Research Scholar in the Center for Biomedical Ethics (SCBE) at Stanford University. She completed her doctorate in medical anthropology from Case Western Reserve University in 2012, and additional training in health services research at the Palo Alto Medical Foundation Research Institute from 2012 through 2016. Her current research focuses at the intersection of the ethics and economics of new genomic technologies. Her current projects include examining ethical issues related to sustainability and governance of patient data and relationships when large clinical genomic studies transition to new models of funding; ethnographic work exploring how diverse stakeholders perceive value in the use of genome sequencing for diagnosis of rare diseases; and the development of new measures for assessing patient-centered outcomes in pediatric rare diseases.

INSTITUTE AFFILIATIONS

- Member, Maternal & Child Health Research Institute (MCHRI)

SERVICE, VOLUNTEER, AND COMMUNITY WORK

- Co-Chair, Patient Education and Empowerment Resource (January 1, 2021 - 12/31/2022)
- Member, Board of Directors (11/1/2021)

Publications

PUBLICATIONS

- **Not in my AI: Moral engagement and disengagement in health care AI development.** *Pacific Symposium on Biocomputing. Pacific Symposium on Biocomputing*
Nichol, A. A., Halley, M. C., Federico, C. A., Cho, M. K., Sankar, P. L.
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- **Developer Perspectives on Potential Harms of Machine Learning Predictive Analytics in Health Care: Qualitative Analysis.** *Journal of medical Internet research*
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- **Blurred Boundaries: Toward an Expanded Ethics of Research and Clinical Care** *AMERICAN JOURNAL OF BIOETHICS*
Halley, M. C., Olson, N. W.
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- **Genomics Research with Undiagnosed Children: Ethical Challenges at the Boundaries of Research and Clinical Care.** *The Journal of pediatrics*

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- **Rare Disease, Advocacy and Justice: Intersecting Disparities in Research and Clinical Care.** *The American journal of bioethics : AJOB*
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Miner, A. S., Stewart, S. A., Halley, M. C., Nelson, L. K., Linos, E.
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- **What is "Personal" About Personal Experience? A Call to Reflexivity for All.** *The American journal of bioethics : AJOB*
Halverson, C., Halley, M.
2023; 23 (1): 39-41
- **Perspectives of Rare Disease Social Media Group Participants on Engaging With Genetic Counselors: Mixed Methods Study.** *Journal of medical Internet research*
Yabumoto, M., Miller, E., Rao, A., Tabor, H. K., Ormond, K. E., Halley, M. C.
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- **Beyond "Ensuring Understanding": Toward a Patient-Partnered Neuroethics of Brain Device Research.** *AJOB neuroscience*
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- **Supporting undiagnosed participants when clinical genomics studies end.** *Nature genetics*
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- **Perceived utility and disutility of genomic sequencing for pediatric patients: Perspectives from parents with diverse sociodemographic characteristics.** *American journal of medical genetics. Part A*
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- **Beyond race: Recruitment of diverse participants in clinical genomics research for rare disease.** *Frontiers in genetics*
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- **From "Ought" to "Is": Surfacing Values in Patient and Family Advocacy in Rare Diseases.** *The American journal of bioethics : AJOB*
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- **Beyond Consent: Building Trusting Relationships With Diverse Populations in Precision Medicine Research** *AMERICAN JOURNAL OF BIOETHICS*
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