Research Ethics Collaborative: Member Biosketches

Melissa Abraham, PhD, MSc is Assistant Professor in the Department of Psychiatry at Massachusetts General Hospital/Harvard Medical School, Faculty Associate at the Center for Bioethics at Harvard Medical School, and a practicing Clinical Psychologist. She is the founder and Director of the Research Ethics Consultation Unit in the Division of Clinical Research at Massachusetts General Hospital. Dr. Abraham served as a Chair of the Partners HealthCare IRB for over a decade. She is interested in the ethical review of social and behavioral research methods in the biomedical setting, QI/research oversight, and improving the quality of IRB submissions and reviews.

Moji Adurogbangba, BDS, MPH, MA is a public health dentist and bioethicist who currently manages the Research and Ethics programs at The Scarborough Hospital in Ontario, Canada. She has 7 years experience serving on different research ethics boards (REB) and now serves as a research ethicist on the Network of Hospitals' REB. Her area of interest is in addressing ethical issues in consent and capacity for treatment and research.

Emily Anderson, PhD, MPH is an associate professor of bioethics and medical education at Loyola University and the director of the Regulatory Knowledge and Support Core of the Center for Clinical and Translational Science at the University of Illinois at Chicago. She serves as associate editor for JERHRE, Narrative Inquiry in Bioethics, and AJOB Empirical Bioethics. She also has over ten years of experience serving on six different IRBs. Her areas of interest and expertise include researcher and physician professionalism and misconduct; ethical issues in research with vulnerable populations; informed consent; institutional review board (IRB) policy; and the application of qualitative research techniques to the study of research ethics.

Jason Arnold, JD, MPH is a lawyer, bioethicist, and educator. He is currently a senior fellow and assistant director of the Clinical and Translational Research Ethics Program at the Medical University of South Carolina. Mr. Arnold has been working in the field of bioethics and health policy for over 20 years and has published numerous peer-review articles in medical journals. His current research focuses on clinical research ethics, global health, emerging technologies, and substance use disorders.

Wajeeh Bajwa, PhD is Director Regulatory Affairs, Clinical and Translational Sciences Institute at the University of Florida. Dr. Bajwa was a research subject advocate and regulatory consultant for the Duke General Clinical Research Center from 2002-2008. Dr. Bajwa was a founding member of the executive board of the Society of Research Subject Advocates. He was president of this organization from 2010-2012. He has extensive experience in industry and academia that includes helping to prepare investigators for pre-pivotal trial discussions with FDA, and writing INDs/IDEs. He has more than 20 years of experience on IRBs and Data and Safety Monitoring Boards/Committees. Dr. Bajwa also provides consults to investigators on ethical issues related to human subject research.

Richard Beech, MD, JD, MBA No bio available at this time.

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Jennifer A.H. Bell, PhD, MA is a Senior Bioethicist in the Department of Clinical and Organizational Ethics at the University Health Network (UHN) in Toronto, Canada, where she provides clinical, organizational, and research ethics consultation to UHN and its partners. Dr. Bell is also a Research

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Scientist within the Department of Supportive Care, Research Division at the Princess Margaret Cancer Centre, and Education Investigator 2 at The Institute for Education Research (TIER) at UHN, and holds faculty appointments in the Department of Psychiatry and Dalla Lana School of Public Health and membership in the Joint Centre for Bioethics at the University of Toronto. Dr. Bell's program of research addresses ethical issues raised by revolutionary advances in cancer care, such as leading-edge personalized genetic-oncology. She has led multiple grant-funded studies focused on ethics and clinical trials, the ethics of novel cancer therapeutics, and medical assistance in dying. Dr. Bell is an ethicist member of the Canadian Agency for Drugs and Technologies in Health pan-Canadian Oncology Drug Review Expert Review Committee and the Canadian Partnership Against Cancer - Health Economics Advisory Committee. She is Principal Investigator of the JustCan Ethics Lab.

Shelly Benjaminy, PhD is a Bioethicist at the Donnelley Ethics Program, Shirley Ryan AbilityLab and Assistant Professor of Physical Therapy and Rehabilitation at the Feinberg School of Medicine at Northwestern University. She received her PhD in Experimental Medicine with specialization in Neuroethics at the University of British Columbia, Canada, and holds an MSc in Health Policy Research from the University of Alberta, Canada. She also completed a clinical ethics fellowship at Providence Health Care and Provincial Health Services Authority in Vancouver, Canada. Her expertise in ethics is positioned at the intersection of research and medical care, and she provides both clinical and research ethics consultation services. She specializes in translational ethics, and her research is embedded within the context of active clinical trials and innovative therapy platforms.

Alina Bennett, MA, MPH, PhD is a Regional Ethicist for Kaiser Permanente, Northern California. The Department of Regional Ethics provides research, clinical and organizational ethics consultations for all Kaiser facilities in Northern California. She completed a postdoctoral research fellowship at the McGovern Center for Humanities & Ethics at the University of Texas Health Science Center at Houston (UTHealth) and her clinical ethics training took place at the University of Texas MD Anderson Cancer Center. At MD Anderson, she served as the embedded ethicist for six clinical services including the Center for Targeted Therapy or, as it is colloquially known, the Phase 1 Group. Her doctoral degree (PhD) was earned at the Institute for the Medical Humanities at the University of Texas Medical Branch at Galveston (UTMB). She also completed a Master of Public Health (MPH) while at UTMB where her work focused on correctional mental health. Dr. Bennett combines the methods of research ethics, clinical ethics, public health, and medical humanities to create a holistic approach to analyzing the moral dimensions of biomedical research and medicine in correctional and free-world clinical settings. She has taught courses on the ethics of scientific research at every institution named above and currently offers such a course through the University of California at San Francisco.

Renaud F. Boulanger, MSc is a research ethicist at the McGill University Health Centre, where he is a member of the Research Ethics Board. He is also one of the founding members of the Save the Children (UK) Research Ethics Review Committee. His scholarship focuses on humanitarian research ethics, the ethics of tuberculosis R&D, and community engagement in global health research. He helped develop the World Health Organization's "Ethics in epidemics, emergencies and disasters: Research, surveillance and patient care" training manual. He is an executive editor of the open access journal Bioéthique Online and has served on the Advisory Committee on Research Ethics of the

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International Development Research Centre, a Canadian Crown Corporation.

Alyssa Burgart, MD, MA is an Assistant Professor of Anesthesiology, Division of Pediatric Anesthesia, at Stanford University and a core faculty member at the Stanford Center for Biomedical Ethics. She is Medical Director of Clinical Ethics and Co-Chair of the Ethics Committee at Lucile Packard Children's Hospital. She provides research ethics consultations, specifically related to pediatrics. Her interests include disability rights, informed consent, organ transplantation, and excellence in ethics consultation.

Jean Cadigan, PhD is Assistant Professor of Social Medicine at the University of North Carolina-Chapel Hill School of Medicine. She is an anthropologist and member of UNC's Research Ethics Consultation Service. She serves on two observational and safety monitoring boards for the National Heart, Blood and Lung Institute. Her work focuses on ethical, social, and policy issues associated with genetic research, biobanking, and HIV cure research.

Alexander M. Capron, LLB is the Scott H. Bice Chair in Healthcare Law, Policy and Ethics, Professor of Law and Medicine, Keck School of Medicine, Co-Director, Pacific Center for Health Policy and Ethics at the University of Southern California. He directs the research ethics program of the Southern California CTSI and heads its consultation service. He served as the first Director of Ethics, Trade, Human Rights and Health Law at the World Health Organization in Geneva and was the Executive Director of the President's Commission for the Study of Ethical Problems in Medicine and Biomedical and Behavioral Research. He currently chairs the Board of Directors of Public Responsibility in Medicine and Research.

Vicky Cardenas, PhD, JD, MHS ~~No bio available at this time~~is a Study Responsible Scientist in the Infectious Disease and Vaccines group at Johnson & Johnson. In that role, she has had scientific oversight of HIV and RSV vaccine clinical trials, with the past 3 years dedicated to Janssen's COVID vaccine program. Vicky is a member of the Johnson & Johnson Bioethics Committee. For 30 years, Dr. Cardenas has been dedicated to running clinical trials on 6 continents. Her experience in clinical trials includes studies in cholera, malaria, tuberculosis, HIV, and COVID-19. Other work experience includes disease modeling, neonatal epidemiology, pediatric morbidity and mortality, injury epidemiology, domestic violence, and organ donation epidemiology. Dr. Cardenas is Colombian by birth and fluent in Spanish and English. Dr. Cardenas received a BS from the University of San Francisco, an MHS in International Health from the Johns Hopkins School of Hygiene and Public Health, a PhD in epidemiology from the University of Washington and a JD from the University of Washington. She maintains a license to practice law in Washington, DC and practices pro bono in the immigration and domestic violence arenas.

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Stephanie Solomon Cargill, PhD, MSPH No bio available at this time.

Michele A. Carter, PhD is the Frances C. and Courtney M. Townsend, Sr., M.D. Professor in Medical Ethics at the University of Texas Medical Branch in the Department of Preventive Medicine and Community Health. She directs the Ethics Support Program of the UTMB's Institute for Translational Sciences and the Research Ethics Consultation Service. She has served on the UTMB Institutional Review Board for more than 7 years, is the Research Subject Advocate, a member of several data and safety monitoring boards,

and a research mentor for the Post-Doctoral Fellowship in Research Ethics. Her major areas of scholarship include philosophical aspects of trust in the helping professions, ethical conduct of human subjects' research, and translational ethics.

Donna T. Chen, MD, MPH is Core Faculty in the Center for Biomedical Ethics and Humanities and Associate Professor in the Department of Public Health Sciences with a joint appointment in the Department of Psychiatry and Neurobehavioral Sciences at the University of Virginia School of Medicine. During her post-doctoral fellowship in the NIH Department of Bioethics she provided research ethics consultation with their Bioethics Consultation Service and has since served on DSMBs for NINDS, NHLBI, NIDA and provided ethics consultation for a variety of investigator-initiated clinical, epidemiologic, genetic, and translational research studies nationally and internationally. She is setting up a pilot research ethics consultation service for UVA's Translational Health Research Institute of Virginia.

Karla Childers, MS No bio available at this time.

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Mildred Cho, PhD is Professor of Pediatrics at Stanford University, Associate Director of the Stanford Center for Biomedical Ethics, and Director of the Center for Integration of Research on Genetics and Ethics (an NIH-supported Center for Excellence in Ethical, Legal and Social Implications Research). She is also Director of Stanford's Benchside Ethics Consultation Service. Dr. Cho's major areas of interest are the ethical and social impacts of genetic research and its applications, and how conflicts of interest affect the conduct of academic biomedical research. Her current research projects examine ethical and social issues in research on prenatal genetic testing, the human microbiome, and synthetic biology, and the ethics of clinical and translational research.

Ellen Clayton, MD, JD is the Craig-Weaver Chair in Pediatrics and Professor of Law at Vanderbilt University. She is currently Co-PI of Genetic Privacy and Identity in Community Settings – GetPreCiSe, a Center of Excellence in ELSI Research, and LawSeq, which will explore four major areas of law and genomics. She has served on the National Advisory Council for Human Genome Research of the NIH, as Co-Chair of the ELSI Working Group of the International HapMap Project, on IOM Committees on Genomics and the Public's Health in the 21st Century and on Assessing Interactions Among Social, Behavioral, and Genetic Factors of Health, on the American Society of Human Genetics Social Issues Committee, as Co-Chair of the Consent and Community Consultation Working Group of the eMERGE Network, and on the HUGO Committee on Ethics, Law, and Society. She is Co-Chair of the Report Review Committee of the National Academies of Sciences, Engineering, and Medicine and Chair of the Health and Medicine Division's Board on Population Health and Public Health Practice. She recently received the David P. Rall Medal for exceptional service to the IOM. Her research has focused on the ethical, legal, and social issues (ELSI) raised by genetics and genomics research and the translation of new findings into clinical care.

Elaine Collier, MD No bio available at this time.

Joshua Crites, PhD No bio available at this time.

Richard Culbertson, PhD is Professor and Director of Health Policy and Systems Management, and

previous Interim Dean, at Louisiana State University School of Public Health and Professor of Family Medicine in the LSU Medical School. He serves as Head of the Ethics Key Resource for the Louisiana Clinical & Translational Science Center. In 2018 he was appointed a formal collaborator to the Puerto Rico Clinical and Translational Research Center for development of its Ethics resource. He is a current member of the American College of Healthcare Executives and contributing Ethics columnist to its official Journal Healthcare Executive, the Medical Group Management Association, the Association of Bioethics and Humanities, Academy Health, American Association for Cancer Research, American Hospital Association Trustee Leadership Network, Kellogg Fellows Leadership Alliance, and the University of Minnesota President's Club. His primary research interests include clinical research ethics, governance, academic medical centers and managed care, organizational structure of medical schools, and physician autonomy.

Reid Cushman, PhD is Assistant Professor of Medicine at the University of Miami Miller School of Medicine and Director of Technology Development for the Collaborative Institutional Training Initiative. He is Co-Director of its Research Ethics Consultation Service and Director of UM's Responsible Conduct of Research Education Program and. As part of the Miami Clinical and Translational Science Initiative, Dr. Cushman's current projects are focused on governance issues for tissue biobanks and electronic health data collections.

Marion Danis, MD is Head of the Section on Ethics and Health Policy in the Department of Bioethics at the National Institutes of Health Clinical Center. She also serves as Chief of the Ethics Consultation Service. Dr. Danis has studied patients' treatment preferences at the end of life and the effectiveness of advance directives in promoting their preferences as well as strategies for fair rationing of limited health care resources and strategies to address the social determinants of health to reduce health disparities. As Chief of the Bioethics Consultation Service she has been the lead editor of the volume published by Oxford University Press entitled *Research Ethics Consultation: A Casebook.* She has chaired the International Society on Priorities in Health Care and has served on the board of American Society for Bioethics and Humanities.

Arlene Davis, JD is Associate Professor of Social Medicine and core faculty in the UNC Center for Bioethics. She directs the clinical ethics services for UNC Hospitals and co-chairs its Hospital Ethics Committee. She is member of the UNC TraCS research ethics consultation group and has served as an IRB member for 20 years. Arlene's practical and scholarly interests fall at the intersections of law and bioethics. They are informed by her experiences in clinical and research ethics consultation, private legal practice, and in pediatric and public health nursing. Her current research collaborations and consultations often focus upon the meanings of creating and using genetic information, the high price of vulnerability labels, especially for children, adolescents and the disabled, and the ways in which law is deployed in research and health care settings.

Liza Dawson, PhD No bio available at this time.

Sabrina Derrington, MD, MA is the Director of the Center for Pediatric Bioethics at Children's Hospital Los Angeles, and Associate Professor of Clinical Pediatrics at the Keck School of Medicine at the University

of Southern California. Dr. Derrington received her bachelor's degree in Chemistry from Hillsdale College (Summa Cum Laude), her medical degree from the University of California, Davis (AOA), and a master's degree in Bioethics and Health Policy from Loyola University Chicago. She completed her residency in pediatrics and a fellowship in pediatric critical care at Children's Hospital Los Angeles. She is a certified healthcare ethics consultant and a member of the board of directors for the American Society for Bioethics and Humanities. After helping to build the ethics program at Ann & Robert H. Lurie Children's Hospital of Chicago, where she was the Associate Director for Clinical Ethics and Education, Dr. Derrington was recruited back to CHLA as the inaugural director for the bioethics center. Dr. Derrington's academic interests include interdisciplinary ethics education and the power of narrative medicine and feminist ethics/standpoint theory to breakdown intercultural barriers. Her research brings a health equity/social justice lens to issues including genomic research and biorepositories, the developmental origins of health and disease, shared decision-making in pediatrics, and the impact of social determinants of health on child/family outcomes after critical illness.

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Arthur R. Derse, MD, JD is Julia and David Uihlein Professor of Medical Humanities and Professor of Bioethics and Emergency Medicine and Director of the Center for Bioethics and Medical Humanities at the Medical College of Wisconsin. He serves as a clinical ethics consultant for the Milwaukee VA Hospital and Children's Hospital of Wisconsin. He is the Ethics Committee Chair and a clinical ethicist at Froedtert Hospital, consultant in emergency medical research at MCW, and former member of the Research Ethics Consultation Service for MCW's Clinical and Translational Science Institute. He has served on the IRB of the University of Wisconsin-Milwaukee. His publications and research have focused on emergency medicine and ethics, emergency medical research, informed consent, confidentiality, end-of-life decision making, and the doctor-patient relationship. He is a member and past chair of the Ethics Committee of the American College of Emergency Physicians, past president of the American Society for Bioethics and Humanities, and former chair of the National Ethics Committee of the Veterans Health Administration. He served on the NIH Working Group on Informed Consent in Clinical Research Conducted under Emergent Circumstances.

Raymond De Vries, PhD No bio available at this time.

Adelaide Doussau, MD, PhD is a postdoctoral research fellow in the Biomedical Ethics Unit of McGill University School of Medicine. She previously studied Public Health and Epidemiology, more specifically on Clinical Trials Methodology in France at the Curie Institute, Paris and Bordeaux School of Public Health. She worked four years as a fellow/assistant professor in the Clinical Trial Unit of Bordeaux University Hospital / School of Public Health (ISPED), and subsequently completed a postdoctoral fellowship in the Department of Bioethics of the National Institutes of Health. She is a member of the Research Ethics Committee of MSF. Her research interests are concentrated on research ethics and she recently focused her research on stepped-wedge design for experimental vaccines in the setting of Ebola outbreak, placebo-controlled clinical trials in cancer, and researchers' judgement in drug development and precision medicine.

Devan Duenas, MA is a Research Associate at the Treuman Katz Center for Pediatric Bioethics at Seattle Children's Research Institute. He currently serves as the Managing Editor for the Challenging Cases in

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Research Ethics series in the American Journal of Bioethics, as well as the Coordinator for the Clinical Research Ethics Consultation Collaborative (CRECC). His research interests include the attitudes, perceptions, and decision-making processes of research participants and the ethical issues related to data and privacy.

Jacob Earl, PhD No bio available at this time.

Lisa Eckstein, SJD is Faculty of Law at the University of Tasmania. She previously completed a post-doctoral fellowship at the NIH Department of Bioethics. Her current research focuses on the governance of medical research, especially in relation to genomics and other emerging technologies. Particular research interests include strategies for gaining and assessing participant consent, the disclosure of genetic research findings, clinical trial monitoring, and racially targeted biomedical research. She has previously held positions at the Australian Law Reform Commission and state and federal Departments of Health.

Leah Eisenberg, JD, MA is Assistant Professor of Medical Humanities at the University of Arkansas for Medical Sciences (UAMS). She serves on two IRBs and is a clinical and research ethicist at UAMS and Arkansas Children's Hospital. She has a special interest in health literacy and improving patient understanding of informed consent, assent, and HIPAA documents.

Carole Federico, PhD, MSc No bio available at this time.

Stuart G. Finder, PhD is Associate Professor in the Department of Medicine at the UCLA David Geffen School of Medicine. He is the Director of the Center for Healthcare Ethics at Cedars-Sinai Medical Center and Chief of the Clinical Ethics Consultation Service. He is also co-chair of the Bioethics Committee and on the Stem Cell Research Oversight Committee/IRB. Dr. Finder is the leader for the Research Ethics Consortium organized under the UCLA Clinical and Translational Science Institute. Dr. Finder is interested in exploring the complexity and implications of moral experiences as actualized in healthcare contexts, including the entire spectrum of the healthcare arena, from patient care to clinical and basic sciences research.

Robin N. Fiore, PhD is Associate Professor of Medicine at the University of Miami Miller School of Medicine and core faculty in the University of Miami Ethics Programs. She is Co-Director of UM's Research Ethics Consultation Service and serves on the Embryonic Stem Cell Research Oversight Committee. As part of the Miami Clinical and Translational Science Initiative, Dr. Fiore's current projects are focused on ethically robust practices in connection with translational research and research involving biobanks and electronic health data.

Celia B. Fisher, PhD is the Marie Ward Doty University Chair in Ethics, Professor of Psychology and the founding Director of the Fordham University Center for Ethics Education and the NIDA funded HIV and Drug Abuse Prevention Research Ethics Institute. She has chaired research and professional ethics committees including for the American Psychological Association, the Society for Research in Child Development, and the American Public Health Association and served as a member of numerous federal and NIH committees including SACHRP, the NIH ABCD study and the HEALing Communities studies, and

chaired the Environmental Protection Agency's Human Studies Review Board. She is the author of Decoding the Ethics Code: A Practical Guide for Psychologists now in its 5th edition, has over 300 publications and 8 edited volumes on the rights and welfare of racial and sexual and gender minority children and adults. Her research has been supported by NIDA, NICHD, NIAID, NIAAA, NIMHD, and NSF. She is a Fellow of the American Association for the Advancement of Science. Her awards include the Lifetime Achievement Award for Excellence in Human Research Protections and the America Psychological Association Award for Outstanding Contributions to Ethics Education.

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Linda Furlini, PhD No bio available at this time.

Cory Goldstein, MA is a Doctoral Candidate in the Department of Philosophy at Western University. His philosophical interests are primarily in applied ethics, particularly the ethical design and conduct of health research involving human participants. He is currently working with two CIHR-funded research teams whose aim is to produce guidance for the ethical design and conduct of pragmatic randomized controlled trials. His doctoral project aims to provide a robust ethical framework to help facilitate the research teams' overlapping goals.

Sidney Golub, PhD is Professor Emeritus of Microbiology & Molecular Genetics at the University of California Irvine (UCI). He currently directs the clinical research ethics unit of the UCI Institute for Clinical and Translational Research and chaired the UCI Human Stem Cell Research Oversight Committee from its inception in 2005 until 2013.He served on the UCI IRB for 10 years and continues to serve on a special IRB responsible for compliance and oversight problems. He also served on the founding Board of Directors of AAHRPP, the accreditation body for IRBs. His major interests are in stem cell ethics and public policy.

Hank Greely, JD is the Deane F. and Kate Edelman Johnson Professor of Law and Professor at Stanford University. He provides research ethics consults as a member of Stanford's Benchside Ethics Consultation Service. He specializes in ethical, legal, and social issues arising from advances in the biosciences, particularly from genetics, neuroscience, and human stem cell research. He chairs the California Advisory Committee on Human Stem Cell Research and directs the Stanford Center for Law and the Biosciences and the Stanford Program in Neuroscience and Society. He serves as a member of the National Academy of Sciences Committee on Science, Technology, and Law.

Colin Halverson, PhD No bio available at this time.

Joyce C. Havstad is Associate Professor with joint appointments in the Department of Philosophy, the Office of Research Integrity and Compliance, and the Center for Clinical and Translational Science. She has written on animal ethics, avian origins, chemical classification, climate science and policy, homology, human reproductive cloning, macroevolution, natural selection, the new mechanistic philosophy, nuclear receptors, philanthropic pedagogy, and voucher specimen collection. She is an editor at Philosophy, Theory, and Practice in Biology, has written informally for The OUP Blog, has appeared on The Brain Scoop, and is a founder and contributor at Extinct, the philosophy of paleontology blog. She teaches ethical theory, history, and practice to undergraduate and graduate students at the University of

Utah in Salt Lake City.

Ann Heesters is the Director of Bioethics at Toronto's University Health Network and Chair of the UHN Rehabilitation Science and Medicine Research Ethics Board. She has been practicing in the field for approximately fifteen years and was the Director of Ethics at The Ottawa Hospital before coming to Toronto in 2009. She has an abiding interest in the professionalization of practicing health care ethicists and, with her colleagues at the American Society of Bioethics and Humanities, helped to author a code of ethics for ethicists. She is also a founding member of Practicing Healthcare Ethicists Exploring Professionalization (PHEEP) and a director of the newly established non-profit Board called Canadian Association of Practicing Healthcare Ethicists (CAPHE).

Saskia Hendriks, MD, PhD is a faculty member in the NIH Department of Bioethics. She conducts empirical and conceptual research on the ethical, social and legal implications of emerging medical technologies in reproduction and neuroscience. Dr. Hendriks has a joint appointment in the Neuroethics Program of the National Institute of Neurological Disorders and Stroke, where she serves as a faculty neuroethics consultant. Dr. Hendriks received her undergraduate degree from Amsterdam University College and her MD-PhD at the University of Amsterdam. She completed a post-doctoral fellowship at the NIH Department of Bioethics before taking up her present position.

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Elizabeth Heitman, PhD is Professor in the Program in Ethics in Science and Medicine at University of Texas Southwestern Medical Center in Dallas, Texas. Her work focuses on cultural dimensions of ethics, international standards for research oversight, and education for responsible conduct of collaborative research. In addition to her work with UT Southwestern's Center for Translational Medicine, she is Co-Director of a Fogarty-sponsored research ethics education and capacity building program in Mozambique, co-Investigator in the NHLB-sponsored Obesity Health Disparities PRIDE research training program, a member of the National Academy of Science's Standing Committee on Educational Institutes for Teaching Responsible Science.

Laurie Herraiz, BS No bio available at this time.

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D. Micah Hester, PhD is Professor of Medical Humanities and Pediatrics and Division Chief of the Division of Medical Humanities at the University of Arkansas Medical School (UAMS), where he directs the research ethics consult service for the UAMS Translational Research Institute and provides clinical ethics consultations at both UAMS and the Arkansas Children's Hospital (ACH), where he is also a clinical ethicist. Dr. Hester serves on the UAMS IRB and has been a member of two Institutional Animal Care and Use Committees. He also coordinates the Pediatric Ethics Consortium, which is a national professional initiative to promote pediatric ethics scholarship and education.

Dana Howard, PhD is a Philosophy faculty member with a primary appointment in the The Ohio State University Center for Bioethics, College of Medicine. Prior to coming to OSU, Dana was a post-doctoral fellow in the Department of Bioethics at the National Institutes of Health. At OSU, she serves as a steering committee member for the Center for Ethics and Human Values, and directs its Conversations About Research Ethics (CARE) program. She received her Ph.D. in Philosophy in 2013 from Brown

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University.

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Raymond Hutchinson, MD, MS No bio available at this time.

Iekuni Ichikawa, MD is Emeritus Professor of Pediatrics at Vanderbilt University. After pediatric residency in Japan, and fellowship at UCSF and Harvard, he served at Vanderbilt University in 1985-2015, and then at Tokai University, Japan in 1998-2012. His research focuses on kidney diseases under the grant support by NIH for nearly 30 years, which led to a discovery of new disease entity, "congenital anomalies of the kidney and urinary tract or "CAKUT", a concept now globally accepted to include a wide spectrum of anomalies found in newborns. Those genetic studies ignited his research and teaching interest in the promotion of research ethics and integrity. He became involved as a member of the Collaborative Institutional Training Initiative (CITI) program, which creates and distributes the Internet teaching materials for biomedical research. In 2016, he co-founded and assumed the position of the Executive Director of, the Association for the Promotion of Research Integrity (APRIN) with the goal of promoting ethics and integrity in research via various ways including providing internet teaching materials, hosting training sessions, organizing discussion groups for proposal to the Government and to the world-wide-research arena. APRIN hosts annual Asia Pacific Rim Research Integrity Network (APRI) meeting in Tokyo in 2023. He is currently chairing the Research Ethics Committee, the Japanese Medical Science Federation and teaching at Shinshu University School of Medicine as a visiting professor.

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Nicholas Jabre, MD, MS is a pediatric pulmonologist and bioethicist at Johns Hopkins All Children's Hospital in St. Petersburg, FL. There, he is a member of the IRB and clinical ethics consult service. He earned his medical degree from the University of Cincinnati College of Medicine and completed his training in pediatrics at Johns Hopkins All Children's Hospital. Dr. Jabre then completed a clinical fellowship in pediatric pulmonology at Johns Hopkins University School of Medicine and a postdoctoral fellowship in bioethics at the Johns Hopkins Berman Institute. His clinical interests include pediatric asthma, aerodigestive disorders, chronic mechanical ventilation, and bronchoscopy. His research has focused on parent-physician communication and how it affects decision-making around tracheostomy and home ventilation.

Brian Jackson, MD No bio available at this time.

Liza-Marie Johnson, MD, MPH, MSB is a pediatric oncology hospitalist and bioethicist at St. Jude Children's Research Hospital in Memphis, TN. She is Chair of the Hospital Ethics Committee and a member of the St. Jude Institutional Review Board. Dr. Johnson conducts clinical and research ethics consultations at St. Jude and is actively engaged in clinical research. Her research interests are focused in pediatric ethics as well as quality-of-life concerns in the context of pediatric cancer. Dr. Johnson is particularly interested in improving communication and decision-making in the context of early phase clinical research trials or in research involving advanced genomic sequencing technologies.

T.J. Kasperbauer, PhD is a Postdoctoral Fellow in the Center for Bioethics at the Indiana University School of Medicine. He conducts research on privacy for genetic and health data, biobanking, and informed consent.

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Jenny Kingsley, MD, MA No bio available at this time

Barbara Koenig, RN, PhD is Professor of Bioethics and Medical Anthropology at the Institute for Health & Aging, University of California, San Francisco, where she leads the research ethics consultation service for UCSF's Clinical and Translational Science Institute. She co-directs an NHGRI "Center of Excellence in ELSI Research" that focuses on translational genomics, co-leads an NCI project on return of results in genomic biobanks, and directs the ELSI component of an NICHD award on newborn screening in an era of whole genome analysis. Dr. Koenig pioneered the use of empirical methods in the study of ethical questions in science, medicine, and health. She was the founding executive director of the Center for Biomedical Ethics at Stanford University; she created and led the Biomedical Ethics Research Program at the Mayo Clinic in Rochester, MN.

Robert H. Kolb, RN, CCRC No bio available at this time.

Tracy Koogler, MD is an Associate Professor of Pediatrics in Pediatric Critical Care Medicine at the University of Chicago. She is also Assistant Director and Co-Director of the Clinical Ethics Consultation Service at the MacLean Center for Clinical Medical Ethics. She has been on the University of Chicago IRB for 9 years and Vice Chair for 5 years. She is on the Ethics and Regulatory Committee for the Chicago PCORI program and is a member on its central IRB Chair B. Her research involves organ donation, decision making for neurologically disabled children, and death and dying.

Stanley Korenman, MD is a Distinguished Professor of Medicine and Associate Dean for Ethics at the David Geffen School of Medicine at UCLA. He is the Regulatory and Ethics Program Director of the UCLA CTSI. He has been director of the Medical Scientist Training Program for MD-PhDs for more than 20 years. He has conducted empirical investigations on the ethical beliefs of scientists in comparison to their administrative overseers regarding the hierarchy of research misconduct and means of punishment. He has authored a book providing instructors methods and materials to teach research ethics. He initiated the Ethics Advisory Committee of the Endocrine Society and led the writing of its Code of Ethics. He is consulted on questions of research integrity as they arise before and during the course of research.

Stephanie Alessi Kraft, JD is an Assistant Professor at the Treuman Katz Center for Pediatric Bioethics at Seattle Children's Research Institute and the Division of Bioethics in the Department of Pediatrics at the University of Washington School of Medicine. Her research addresses the ethics of research in integrated clinical-research settings, the ethical, legal, and social implications of genetic testing, and issues in clinical communication and quality of life for patients with serious illnesses.

Benjamin Krohmal, JD No bio available at this time.

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Lisa M. Lee, PhD, MA, MS is Associate Vice President for Research and Innovation and Director of the Division of Scholarly Integrity and Research Compliance at Virginia Tech. She also holds a faculty appointment in the Department of Population Health Sciences. Previously, she served as Chief of Bioethics & Human Subjects Research at the Walter Reed Army Institute of Research where she served as the IRB Chair, the Research Integrity Officer, and Chair of the Research Ethics Consultation Service. From 2012-

2017, she served as Executive Director of President Obama's bioethics commission. For 14 years she served in numerous positions at the Centers for Disease Control and Prevention, including the agency's Assistant Science Officer and Chief of the Office of Scientific Integrity. She is an epidemiologist, bioethicist, and ethics educator.

Sandra Soo-Jin Lee, PhD is Chief of the Division of Ethics and Associate Professor in the Department of Medical Humanities and Ethics at Columbia University. She is a medical anthropologist with extensive experience leading empirical bioethics research that focuses on the sociocultural and ethical dimensions of emerging genomic technologies. Dr. Lee has served as Chairperson of the Cancer Prevention Institute of California IRB and on the NIH Coriell Consultation and Oversight Committee of the International Haplotype Map. She currently serves on both the Scientific and Bioethics Advisory Boards of the Kaiser Permanente National Research Biobank and the NIH/NHGRI Genomics and Society Working Group. Dr. Lee has expertise on the ethics of precision medicine research, the learning health system, recruitment of diverse and historically under-represented groups in biomedical research and in qualitative research methodologies.

Jason D. Lesandrini, PhD(c) is Assistant Vice President of Ethics, Advance Care Planning and Spiritual Health for WellStar Health System and an adjunct Faculty member at Mercer University, Atlanta Campus. His research interests include methods of ethics consultation, ethics program development and research ethics consultation services.

Anna Lewis, PhD No bio available at this time.

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Walter Limehouse, MD, MA is an Associate Professor of Emergency Medicine at Medical University of South Carolina in Charleston. He directs the healthcare ethics consultation service, chairs the Medical University Hospital Ethics Committee, and founded the research ethics consultation service. Dr. Limehouse is faculty of South Carolina Translational Research Institute's Clinical Research Ethics Fellowship and teaches clinical ethics to MUSC students & residents; his current research involves implementing and evaluating clinicians' respect for patients' choices regarding end-of-life treatment options.

Katherine E. MacDuffie, PhD is a Licensed Clinical Psychologist and Postdoctoral Research Associate at the University of Washington Autism Center. She is currently working on an F32 training grant in neuroethics through the NIH BRAIN Initiative. Her research is focused on ethical issues in neuroimaging research, including results disclosure to participants with psychiatric or neurodevelopment disorders.

David Magnus, PhD is Thomas A. Raffin Professor of Medicine and Biomedical Ethics, and Professor of Pediatrics at Stanford University. He directs the Stanford Center for Biomedical Ethics and co-chairs Stanford Hospital and Clinics' Ethics Committee. Dr. Magnus is co-director of the research ethics program for Stanford's CTSA, is a member of Stanford's IRB and Stem Cell Research Oversight Committee, and has extensive experience as a research ethics consultant. His research focuses on a wide range of topics in bioethics, including research ethics, the ethics of comparative effectiveness research, transplant ethics,

genetics/genomics, and issues in patient/physician communication. Other leadership responsibilities include past President of the Association of Bioethics Program Directors and Editor in Chief of the American Journal of Bioethics.

Daphne Martschenko, PhD is a Postdoctoral Research Fellow at the Stanford Center for Biomedical Ethics, a BioFutures Fellow in the Stanford Department of BioEngineering, and co-organizer of the international Race, Empire, and Education Research Collective (REE). Dr. Martschenko holds an MPhil from the University of Cambridge in Politics, Development, and Democratic Education and a Ph.D. in Education, also from the University of Cambridge. Her work advocates for and facilitates research efforts that promote socially and ethically responsible research, research communication, and community engagement with social and behavioral genomics.

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Zubin Master, PhD is an Associate Consultant II in the Biomedical Ethics Research Program at Mayo Clinic. He completed post-doctoral fellowships in bioethics and health policy at Dalhousie University and the University of British Columbia and previously held the position of Associate Professor at the Alden March Bioethics Institute of Albany Medical College, Research Associate for University of Alberta's Health Law Institute. Dr. Master also worked in public service as a Senior Policy Advisor at Health Canada where he led the development of Health Canada's Scientific Integrity Framework and developed regulations under the Assisted Human Reproduction Act. His research interests broadly cover the ethical and policy issues related to stem cells and regenerative medicine, genetics, research ethics, and the responsible conduct of research. Dr. Master is part of the Clinical and Translational Research Ethics Consultation service at Mayo Clinic and serves on several other committees and journal editorial boards.

Jennifer B. McCormick, PhD, MPP is an Assistant Professor of Biomedical Ethics in the Division of Health Care Policy and Research at the Mayo Clinic Rochester. She is one of the core faculty members of the Biomedical Ethics Research Program and is the Associate Director of the Clinical and Translational Research Ethics Program. She also directs the Mayo Clinic's Responsible Conduct in Research course. Dr. McCormick's major areas of interest are the ethical, policy, and social impacts of translational genetic and genomic research and its applications; the challenges of navigating the blurriness between translational research and clinical care; and 'big data'.

Ross McKinney, Jr, MD is a Professor of Pediatrics and Director of the Trent Center for Bioethics, Humanities, and History of Medicine at the Duke University School of Medicine. He is also Director of the Research Ethics Core for Duke's Clinical and Translational Science Award program. Previously, Dr. McKinney was the Vice Dean for Research for the Duke School of Medicine. Dr. McKinney's ethics research has focused on conflict of interest, but he has also published articles related to the ethics of sports medicine and issues related to informed consent. He has chaired several studies of antiretroviral treatment strategies for children with HIV infection.

Lindsay McNair, MD, MPH, MSB is the Principal Consultant at Equipoise Consulting, LLC. From 2013 to 2023, Dr. McNair was the Chief Medical Officer for the WIRB-Copernicus Group (WCG). As part of the WCG IRB Executive Committee, she oversaw IRB member selection and training, IRB policy development, and helped to maintain regulatory and accreditation compliance. She also provided consultation to

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institutions and pharma/biotech companies on a wide range of issues related to protocol design, regulatory compliance, human subject protection, and ethical policy development (pre-approval access, subject compensation). Dr. McNair is adjunct faculty at Boston University and teaches graduate courses on the scientific design of clinical research studies. Dr. McNair is an associate editor for the Journal of Empirical Research on Human Research Ethics, and is in the NYU Compassionate Use Pre-Approval Access (CUPA) Working Group. She has previously been part of the Human Subjects Review Board of the US Environmental Protection Agency (EPA HSRB), various working groups within Harvard's Multi-Regional Clinical Trials (MRCT) program and in the Advancing Effective Research Ethics Oversight (AEREO) consortium.

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Michelle N. Meyer, PhD, JD is an Assistant Professor and Associate Director of Research Ethics in the Center for Translational Bioethics and Health Care Policy at Geisinger Health System, where she serves as a faculty advisor to the IRB, chair of the IRB Leadership Committee, director of the Research Ethics and Advice Consultation Service, and a member of a task force that seeks to advance Geisinger as a learning healthcare system and develop oversight systems for learning healthcare activities. In her own research, she focuses on ethical, legal, and policy issues that arise in biospecimens and genetic/genomic research; social science research; research with big data; corporate research; research on medical practice, standard of care research, and comparative effectiveness research; and randomized evaluations, QI/QA, innovation, and other learning activities that may not meet the federal regulatory definition of human subjects research.

Skye A. Miner, PhD is an Assistant Professor in the Department of Medical Humanities and Bioethics at University of Arkansas for Medical Sciences. She received her PhD in Sociology at McGill University and completed a postdoctoral fellowship in the Department of Bioethics at the National Institutes of Health. Her research focuses on ethical issues in emerging technologies at the beginning of life and in pediatric populations. More specifically, she is interested in how genetic information is used in the context of family-building and making health care decisions about children. She is also passionate about research ethics consultations and is looking for ways to bolster the research ethics consultation service at UAMS.

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Kevin Mintz, PhD, received his Ph.D. from the Department of Political Science here at Stanford University in 2019. He also holds an AB in Government from Harvard College, an MSc in Political Theory from the London School of Economics and Political Science, and a Doctorate of Human Sexuality from the Institute for Advanced Study of Human Sexuality. Prior to returning to Stanford, Kevin was a Postdoctoral Fellow in The Department of Bioethics at the National Institutes of Health. His research focuses on disability bioethics, research ethics, business ethics, and the degree to which genetics should be used to construct social or political identities. His work has appeared in a variety of academic journals and newspapers, including Pediatrics, The Hastings Center Report, and the Los Angeles Times.

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Stephanie Morain, PhD, MPH is a core faculty member at the Berman Institute of Bioethics, and an assistant professor in the Department of Health Policy and Management at the Bloomberg School of Public Health. She conducts both empirical and normative research into issues at the intersection of ethics, law, and health policy. Dr. Morain's work examines political and ethical issues concerning the scope of government authority in public health and the role of stakeholder opinion in shaping decision-

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making in public health policy. Specific research interests include the ethics and politics of disease control and injury prevention; public health law; and ethical and policy challenges presented by the transition to learning health care systems. She was most recently an assistant professor in the Center for Medical Ethics and Health Policy at Baylor College of Medicine.

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Daniel Nelson, MSc No bio available at this time.

Pilar Ossorio, PhD, JD is a Professor of Law and Bioethics at the University of Wisconsin (UW). She is co-director of the Research Ethics Consultation Service, co-director of the Law and Neuroscience Program, and leader of the ethics core for Center for Predictive Computational Phenotyping. She served for 11 years on the health sciences IRB. She is also director of the Ethics Program at the UW-affiliated Morgridge Institute for Research. Dr. Ossorio's research interests include ethical and social issues in genome research and clinical genomics; human subjects research; uses of race in research and medicine; governance of data sharing in research; ethical and social issues in data science; and regulation of medical devices.

Rebecca D. Pentz, PhD is Professor of Research Ethics at Emory University School of Medicine in Atlanta. She directs the research ethics consultation service jointly sponsored by Winship Cancer Institute and Atlanta Clinical and Translational Science Institute. Her empirical ethics research focuses on genetic testing, confidentiality, biobanking, return of results, duty to warn and informed consent ethical issues in early drug development. She has a special interest in pediatric bone marrow transplant and the effect on the family. Before coming to Emory, she designed and directed the clinical ethics program at The University of Texas MD Anderson Cancer Center. She represents Emory on various national data safety monitoring boards and scientific advisory committees, including the Bone Marrow Transplant Clinical Trials Network, ALS repository, St. Jude, and the National Disease Research Interchange.

Kathryn M. Porter, JD, MPH is a Research Scientist for the Treuman Katz Center for Pediatric Bioethics at Seattle Children's Research Institute. She serves as Director of the Research Bioethics Consult Service for the Institute of Translational Health Sciences. She is the Co-Chair for the CREC Collaborative and the Chair of the American Society for Bioethics and Humanities' Clinical Research Ethics Consultation Affinity Group. Her interests include research ethics, the informed consent process, and the ethical and legal issues related to genetics.

Vasiliki (Vaso) Rahimzadeh, PhD, MSc is a Postdoctoral Fellow at the Stanford Center for Biomedical Ethics. Her research centers on the ethical, legal and social implications of emerging health information technologies with special interest in international protections of genetic/genomic data involving children. She serves on the Regulatory and Ethics Work Stream of the Global Alliance for Genomics and Health, where she develops guidance on data access committee review standards and research ethics approval procedures for international, multi-site studies in the data-intensive sciences.

Alan Regenberg, MBe No bio available at this time.

Elisa Reverman, MA No bio available at this time.

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Research Ethics Collaborative: Member Biosketches

Habib Rizk, MD No bio available at this time.

Rosamond Rhodes, PhD is Professor of Medical Education and Director of Bioethics Education at Icahn School of Medicine at Mount Sinai, Professor of Philosophy at The Graduate Center, CUNY, and Professor of Bioethics and Associate Director of the Union-Mount Sinai Bioethics Program. For the past 27 years she has served in numerous roles at Mount Sinai including member of the Institutional Animal Care and Use Committee (IACUC), Secretary to the Ethics Committee, and Director of the Research ethics Consult Service. Dr. Rhodes collaborates on a variety studies related to genetics, emergency medicine, research without consent, biobanks, and controlled substance research.

Lainie Friedman Ross, MD, PhD is the Carolyn and Matthew Bucksbaum Professor of Clinical Medical Ethics at the University of Chicago and a Professor in the Departments of Pediatrics, Medicine, Surgery, and the College. She is also Director of the Research Ethics Consultation Service, Associate Director of the MacLean Center for Clinical Medical Ethics, and Co-Director of Translational Medicine (ITM) at the University of Chicago. Dr. Ross' research focuses on ethical and policy issues in pediatrics, organ and tissue transplantation, research ethics, and genetic testing and screening.

Matthew D. Rotelli, PhD is the Senior Advisor for the Bioethics Program at Eli Lilly and Company in Indianapolis, Indiana where he leads the company's evaluation of bioethical considerations across the continuum of its research, development, and commercialization activities. Dr. Rotelli is a graduate of the Lilly Bioethics Leadership Academy (BELA) and a member of the American Statistical Association (ASA), the American Society for Clinical Pharmacology and Therapeutics, the International Society of Pharmacometrics (ISoP), the American Society for Bioethics and Humanites (ASBH), and Public Responsibility in Medicine and Research (PRIM&R). His research interests include the intersection of science and bioethics.

Erin Rothwell, PhD is the Associate Vice President for Research Integrity and a professor in the Department of Ob/Gyn in the School of Medicine at the University of Utah. She has extensive experience in bioethics and human subjects protections having served on the IRB and hospital ethics committees, and completed a competitive bioethics fellowship from the Medical College of Wisconsin. Dr. Rothwell also brings a wealth of experience from her successful program of research on informed patient decision making and the ethical implications of emerging technologies within the context of genomics, population screening and public health across the reproductive continuum of care. Currently, she is a member of the leadership teams for the Center for Clinical and Translational Science and the NIH Center in Excellence for Ethical, Legal and Social Implications in Genetics. Her current research focuses on the consent process for the storage and research use of residual newborn screening blood spots.

Alham Saadat, MS No bio available at this time.

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John Z. Sadler, MD is the Daniel W. Foster, M.D. Professor of Medical Ethics and a Professor of Psychiatry and Clinical Sciences at the University of Texas Southwestern Medical Center. He has 25 years of clinical ethics consultation experience and developed his research ethics consultation knowledge and skills in the

context of an NIH CTSA award to his institution in 2006. His own research in this area has focused on the ethics of research funding priorities, relationships between IRB and research ethics consultation, and most recently ethical issues in perinatal research populations.

Donya Sarrafian No bio available at this time.

Toby Schonfeld, PhD No bio available at this time.

Peter Schwartz, MD, PhD is Associate Professor of Medicine at the Indiana University School of Medicine and Faculty Investigator at the Indiana University Center for Bioethics. He is also Associate Professor of Philosophy at Indiana University – Purdue University, Indianapolis (IUPUI). He directs the Translational Research Ethics Consultation Service of the Indiana Clinical and Translational Sciences Institute and is a research subject advocate in the Bioethics and Subject Advocacy Program (BSAP) and. Dr. Schwartz's current research focuses on ethical issues and patient behavior in preventive medicine, personalized (or "precision") medicine, and in the design and use of electronic health records.

J. Jina Shah, MD, MPH is Senior Medical Director, Bioethics at Genentech, a member of the Roche Group. Board certified in Family Medicine and General Preventive Medicine, she leads the Ethics Consultation service for drug development teams globally at Roche. Areas of focus include study design and conduct, informed consent in pediatric trials, return of genomic results to individual patients, and global health.

Seema K. Shah, JD is Associate Professor in the Division of Bioethics in the Department of Pediatrics at the University of Washington School of Medicine and a faculty member in the Treuman Katz Center for Pediatric Bioethics at Seattle Children's Research Institute. She serves as an attending for the Bioethics Consultation Service and the Research Bioethics Consult Service for the Institute of Translational Health Sciences. She has 9 years of experience on IRBs and DSMBs. Her current scholarship focuses on ethical and policy issues related to international and pediatric research and the determination of death.

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Richard Sharp, PhD is Professor of Medicine and Director of the Biomedical Ethics Research Program at Mayo Clinic and the Clinical & Translational Research Ethics Program. He has studied the integration of genetic technologies into patient care, best practices for clinical ethics consultation, financial conflicts of interest in research, and ethical dimensions of patient advocacy. His current research examines how patients and healthcare providers view new forms of personalized medicine. Dr. Sharp advises healthcare organizations on ethical issues and has served on advisory committees for the National Institutes of Health, Institute of Medicine, and the Environmental Protection Agency.

Emily Shulkin, BA No bio available at this time.

Anne R. Simpson, MD is the Rust Professor of Ethics and Professor of Medicine, Division of Geriatrics at the University of New Mexico Health Sciences Center School of Medicine, where she is also Associate Vice Chancellor for African American Health She is director of the Institute for Ethics, executive director for the Black Health Resource Center, and chair of the Ethics Consult Service. Her clinical practice is focused in geriatric medicine and end-of-life care with an additional focus on the social determinants of health.

Research Ethics Collaborative: Member Biosketches

~~Dr.~~ **Elise Smith, PhD, MA** is an Assistant Professor in the Department of Preventive Medicine and Population Health (PMPH) with a membership in The Institute of Translational Science and The Institute for Medical Humanities. As a bioethics scholar with a background in philosophy, law, and the social sciences, she works on projects in research ethics, research integrity and public health ethics. As a research ethics consultant at the University of Texas Medical Branch, Dr Smith is assisting in developing, applying, and evaluating relational research ethics approaches in multi-institutional multidisciplinary research teams.

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Kayte Spector-Bagdady, JD, MBE is an Assistant Professor in the Department of Obstetrics and Gynecology at the University of Michigan Medical School. She is also Chief of the Research Ethics Service and Chair of the Research Ethics Committee under the Center for Bioethics and Social Sciences in Medicine (CBSSM). She is a former drug and device attorney and Associate Director of the Presidential Commission for the Study of Bioethical Issues. She is also a clinical ethics consultant for the Adult and Pediatric Ethics Committees and Clinical Ethics Service and is a member of the UM IRB Council. Her current research explores informed consent to emerging technologies with a focus on reproduction and genetics.

Ryan Spellecy, PhD is Associate Professor of Bioethics and Medical Humanities in the Center for Bioethics and Medical Humanities at the Medical College of Wisconsin. His scholarship has focused on research ethics, informed consent, ethical issues in psychiatry, and community involvement in research. He chairs an IRB at the Medical College of Wisconsin and is faculty advisor for the Regulatory and Ethics Module of the Clinical and Translational Science Institute of Southeastern Wisconsin.

Jeffrey Spike, PhD No bio available at this time.

Mark A. Stein, PhD is Professor of Psychiatry and Behavioral Sciences and Pediatrics at the University of Washington. He completed a fellowship in Clinical and Medical Ethics at the MacLean Center at The University of Chicago. Areas of interest include neuroethics, clinical trials, incidental findings in research, and performance enhancement.

Eric S. Swirsky, JD is Clinical Assistant Professor in the Department of Biomedical and Health Information Management at the University of Illinois at Chicago. He serves on the ethics committee of the University of Illinois Hospital and the Research Ethics Core of the university's Center for Clinical and Translational Science. His current research interests focus on the impacts of electronic medical records and digital media upon heath care economics, clinical research and decision-making, and provider-patient relationships.

Holly A. Taylor, MPH, PhD is Research Bioethicist in the Department of Bioethics, Clinical Center, National Institutes of Health (NIH). She will be an attending faculty member of the Clinical Center Bioethics Consultation Service. Prior to her move to the NIH she directed the Research Ethics Consulting Service at Johns Hopkins University, serving faculty in the Schools of Public Health, Medicine, and Nursing. Dr. Taylor has expertise in public health research, researchers' obligation to their research subjects, informed consent, recruitment into clinical trials, research oversight, and has experience with quantitative and

qualitative research methodology.

Kory Trott, JD is Director of Research Integrity and Consultation within the Division of Scholarly Integrity and Research Compliance at Virginia Polytechnic Institute and State University. Before joining Virginia Tech, he worked for the Centers for Disease Control and Prevention (CDC) in Atlanta, where he started as a Presidential Management Fellow. Upon completion of his postgraduate fellowship, he joined the Office of the Associate Director for Science, where he served as a member of the Public Health Ethics and Strategy Unit. He provided ethics training to and consultation for the agency's scientists. He served on the Zika Virus Policy Team in the CDC's Emergency Operations Center. Prior to joining CDC, he worked as a reviewer for the University of Wisconsin Health Sciences IRB.

Emma Tumilty, PhD is the Translational Ethics Postdoctoral Fellow in the Institute for Translational Sciences (ITS) at University of Texas Medical Branch Galveston (UTMB) and a Clinical Ethics Fellow within the UTMB Health System. She sits on both the Institutional Review Board and the Institutional Ethics Committee at UTMB and provides research consultation services within ITS and to the broader UTMB research community. Her interests include the effectiveness of research ethics review and research ethics education strategies, as well as the ethical issues that arise in translational science specifically around the intersection of research and clinical practice. Her background includes health service & system research and empirical research approaches to bioethical issues.

Leigh Turner, PhD is a Professor in the Department of Health, Society, and Behavior, Program in Public Health at the University of California, Irvine. He is the Executive Director of the UCI Bioethics Program and leads the UCI Clinical Research Ethics Consultation Service. Turner's current research addresses ethical, legal, and social issues related to stem cells and regenerative medicine products. In particular, he uses approaches from bioethics and the social sciences to examine clinics engaged in direct-to-consumer marketing of unproven and unlicensed cell-based interventions. He also studies ethical issues related to crowdfunding for medical care, cross-border health-related travel, and other topics. Turner is a co-editor of *Risks and Challenges in Medical Tourism: Understanding the Global Market for Health Services and The View from Here: Bioethics and the Social Sciences*.

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Sarah B. Vittone DBe, MSN, MA, RN is an assistant professor at Georgetown University. She joined the School of Nursing and Health Studies in 2007. Dr. Vittone's teaching interests are in pediatrics and clinical/research ethics. She has 25+ years experience in clinical ethics consultation and since 2007 has been a primary consultant with the Ethics Consultation Service of the Pellegrino Center for Clinical Bioethics here at GUMC and MGUH. Her clinical ethics interests are in complex decision making, surrogate decision makers, and issues at the interface of vulnerable populations and health systems. Since 2011, Dr. Vittone has received grant support for her work in human protection and research ethics for the Clinical and Translational Sciences Award with Georgetown-Howard Universities-Veterans Administration- Oak Ridge National Lab and Medstar Health Research Institute as a Research Subject Advocate. She teaches Research Ethics for the Masters in Clinical and Translational Research program. She teaches Advanced Ethical Reasoning and the Ethics Consultation Intensive for the Catholic Clinical Ethics Masters program. She also currently serves on IRB-E, and the Adverse Events/DSMB Committee (CAESM) at Georgetown University. She is the course director for the Ethics,

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Research Ethics Collaborative: Member Biosketches

Regulatory and Implementation of Clinical Research Studies with Human Subjects on SiTel provided by the GHUCCTS CTSA.

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Charles Weijer, MD, PhD is Professor and Canada Research Chair in Bioethics at the Rotman Institute of Philosophy at Western University in London, Canada. He is a leading expert on the ethics of randomized controlled trials and co-led a collaboration that produced the first international ethics guidelines for cluster randomized trials. Dr. Weijer founded the Rotman Institute of Philosophy, which is dedicated to fostering collaboration between the humanities and the sciences, and served as the Institute's first director. In 2014, he received Western's Hellmuth Prize for Achievement in Research and in 2016 was elected to the Royal Society of Canada. Dr. Weijer's current work explores ethical issues in pragmatic randomized controlled trials that evaluate health interventions in real-world conditions to better inform patients, health providers and health systems managers.

Elliott Weiss, MD No bio available at this time.

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Benjamin S. Wilfond, MD is Professor and Chief of the Division of Bioethics in the Department of Pediatrics at the University of Washington School of Medicine and the Director of the Treuman Katz Center for Pediatric Bioethics at Seattle Children's Research Institute. Dr. Wilfond is the Chief of the Bioethics Consultation Service and a pediatric pulmonologist at Seattle Children's Hospital. He serves as a consultant for the Research Bioethics Consult Service for the Institute of Translational Health Sciences. Dr. Wilfond is the former chair of the intramural NHGRI IRB and has 25 years of experience on IRBs and DMCs. His current scholarship focuses on ethical and policy issues related to genetic testing, pediatrics, and clinical research.

Susan Wootton, MD is Associate Professor in the Pediatric Infectious Disease Division at the University of Texas Health Science Center at Houston and member of the Center for Clinical Research & Evidence Based Medicine. Since 2009 she has been actively involved in issues related to research ethics through her participation in a local ethics working group at UT and joined the IRB in September 2016. Her current research interests include vaccine policy and she is developing a collaborative project among multiple institutions to address vaccine delinquency rates within the largest school district in Texas.

Daniel Wyzynski is the Clinical and Research Ethicist for the London Health Sciences Centre in London, Ontario. He is actively involved in research ethics consultations within LHSC and is also an associate member for the Canadian Association of Practicing Healthcare Ethicists. Daniel has completed his Master's in bioethics at the University of Toronto's Joint Centre of Bioethics, and is interested in the distinctions between research ethics oversight, quality improvement, and quality assurance.

Mark Yarborough, PhD is the Dean's Professor of Bioethics at the University of California Davis and Director of the Clinical Research Ethics Program for the Clinical and Translational Science Center. He established the research ethics consultation service at UC Davis. His major area of scholarly focus has been on topics concerning the ethical conduct of human subjects' research and he is particularly interested in discovering what practices contribute to making research trustworthy, as well as ethical issues related to stem cell research for neurological disorders. He has served on or consulted with IRBs

since 1984.