

Virtual Mentor

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About the Contributors

Theme Issue Editor

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Contributors

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Don B. Bailey, Jr., PhD, is a fellow at RTI International. He directs several projects funded by the National Institutes of Health and the Centers for Disease Control and Prevention on various aspects of fragile X syndrome and broader issues surrounding the ethical, legal, and social consequences of genetic discoveries and the disclosure of genetic information to families, especially in the context of newborn screening.

Wylie Burke, MD, PhD, is professor and chair of the Department of Bioethics and Humanities at the University of Washington in Seattle. She is also principal investigator of the University of Washington Center for Genomics and Healthcare Equality, an NIH-funded Center of Excellence in Ethical, Legal, and Social Implications (ELSI) Research.

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Leon Dure, MD, is the division chief of pediatric neurology in the Department of Pediatrics at the University of Alabama School of Medicine at Birmingham. He is also the William Bew White Jr. Professor of Pediatrics and Neurology at the

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Robert Klitzman, MD, is the director of the Ethics and Policy Core of the HIV Center in New York City, and cofounded the Columbia University Center for Bioethics. He is the director of the master of bioethics program being established at Columbia, and a member of the Division of Psychiatry, Law and Ethics in the Department of Psychiatry. Dr. Klitzman is the author of *Being Positive: The Lives of Men and Women with HIV* and *Mortal Secrets: Truth and Lies in the Age of AIDS*, among others. He was the recipient of a Mentored Clinical Scientist Award (K08) from the National Institute of Mental Health, and received several awards for his work, including fellowships from the Russell Sage Foundation, Commonwealth Fund, and Rockefeller Foundation.

Anne-Marie Laberge, MD, PhD, is a medical geneticist at Centre Hospitalier Universitaire (CHU) Sainte-Justine and clinical assistant professor in the Department of Pediatrics at Université de Montréal. Her research addresses the use of genetic tests in clinical practice and how they influence management of patient care.

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Cynthia Powell, MD, MS, is an associate professor of pediatrics and genetics, and chief of the Division of Genetics and Metabolism in the Department of Pediatrics at the University of North Carolina at Chapel Hill. A medical geneticist and trained genetic counselor, Dr. Powell oversees the medical genetics residency program and the pediatric genetic clinic at UNC-CH.

Myra I. Roche, MS, CGC, is a certified genetic counselor and associate professor in the Division of Genetics and Metabolism in the Department of Pediatrics at the University of North Carolina at Chapel Hill. Her clinical and research interests include improving families' understanding of genetic information and the ethical implications of genetic testing.

Kristin E. Schleiter, JD, LLM, is a senior research associate for the Council on Ethical and Judicial Affairs for the American Medical Association in Chicago. She analyzes ethics policy and law and assists in the development and dissemination of ethics policy and related educational material. Ms. Schleiter received both her law degree and masters of law in health law from Loyola University Chicago School of Law, where she was a contributing writer for the *Annals of Health Law*.

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Mike Spear is the director of corporate communications for Genome Alberta, a not-for-profit organization that funds and manages genomics research. He is a former journalist with the Canadian Broadcasting Corporation and speaks extensively on the use of social media in public relations and journalism. His genome has been sequenced by four direct-to-consumer testing companies, and the results have been posted on the web.

David Wasserman, JD, is a research scholar at the University of Maryland's Institute for Philosophy and Public Policy and the director of research at the Center for Ethics at Yeshiva University. He works on ethical and policy issues in disability, genetics, and reproduction.

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