Stanford

# Holly Tabor

Professor of Medicine (Primary Care & Population Health) and, by courtesy, of Pediatrics (Stanford Center of Biomedical Ethics)

Medicine - Primary Care and Population Health

### CONTACT INFORMATION

- **Administrative Contact**

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## Bio

### BIO

Holly Tabor, PhD, is the Director of the Stanford Center for Biomedical Ethics. She is Professor of Medicine at Stanford University, and by Courtesy of Pediatrics and Epidemiology and Population Health. She is also Co-Chair of the Ethics Committees at Stanford Hospital and Lucile Packard Children's Hospital. She is a globally recognized expert on the ethical issues surrounding health care and research for patients with disabilities, especially intellectual and developmental disability, and on the ethical, legal, and social issues (ELSI) in genetics. Her research has shed light on the benefits and risks of participating in genomic research, particularly of rare and undiagnosed diseases. She is Editor-in-Chief of the American Journal of Bioethical Empirical Research.

### ACADEMIC APPOINTMENTS

- Professor - University Medical Line, Medicine - Primary Care and Population Health
- Professor - University Medical Line (By courtesy), Pediatrics - Center for Biomedical Ethics

### ADMINISTRATIVE APPOINTMENTS

- Director, Stanford Center for Biomedical Ethics, (2024- present)
- Co-Chair, Lucile Packard Children's Hospital Ethics Committee, (2018- present)
- Co-Chair, Stanford Hospital Clinical Ethics Committee, (2018- present)
- Member, Stanford Hospital Clinical Ethics Committee, (2016-2018)
- Member, Lucile Packard Children's Hospital Ethics Committee, (2016-2018)

### HONORS AND AWARDS

- Henry J. Kaiser Award for Excellence in Preclinical Teaching, Stanford School of Medicine (06/2022)
- Allyship Award, Stanford Faculty Women’s Forum (06/2022)
- Outstanding Community-Engaged Faculty Award, Stanford School of Medicine (04/2023)

### PROFESSIONAL EDUCATION

- Postdoc, Stanford University School of Medicine , Bioethics (2008)
- PhD, Stanford University School of Medicine , Epidemiology (Minor Genetics) (2002)

- AB, Harvard University , History and Science (1994)

### LINKS

- Center for Biomedical Ethics: http://med.stanford.edu/bioethics.html

## Research & Scholarship

### CURRENT RESEARCH AND SCHOLARLY INTERESTS

My research focuses on ethical issues in genetics and genomics, specifically return of results and translation for exome and whole genome sequencing and translation of genomic sequencing into the clinical setting. I also conduct research on ethical issues in clinical care and research for patients and families with autism and other developmental and cognitive disabilities.

## Teaching

### COURSES

**2023-24**

- Caring for Individuals with Disabilities: MED 276 (Win)
- Modern Ethical Challenges in Neuroscience and Organ Transplantation: HUMBIO 171E (Spr)
- Shades of Grey Between Life and Death: Neuro-Ethics Across the Pacific: OSPGEN 45 (Sum)

**2022-23**

- Caring for Individuals with Disabilities: MED 276 (Win)
- Modern Ethical Challenges in Neuroscience and Organ Transplantation: HUMBIO 171E, MED 142 (Spr)
- The Responsible Conduct of Research: MED 255 (Aut, Win, Spr)

**2021-22**

- A cultural, ethical, medical and legal exploration of Japanese and American Societies: OSPGEN 29 (Sum)
- Caring for Individuals with Disabilities: MED 276 (Spr)
- Modern Ethical Challenges in Neuroscience and Organ Transplantation: HUMBIO 171E, MED 142 (Spr)

**2020-21**

- Medical Ethics I: PEDS 251A (Win)
- Modern Ethical Challenges in Neuroscience and Organ Transplantation: HUMBIO 171N, MED 142 (Spr)
- The Responsible Conduct of Research: MED 255 (Aut, Win, Spr, Sum)

### STANFORD ADVISEES

**Med Scholar Project Advisor**

Ben Schwartz

## Publications

### PUBLICATIONS

- **"Like not having an arm": a qualitative study of the impact of visitor restrictions on cancer care during the COVID-19 pandemic.** *Supportive care in cancer : official journal of the Multinational Association of Supportive Care in Cancer*
  Holdsworth, L. M., Siden, R., Wong, B. O., Verano, M., Lessios, A. S., Tabor, H. K., Schapira, L., Aslakson, R.
  2024; 32 (5): 288

- **Genomics Research with Undiagnosed Children: Ethical Challenges at the Boundaries of Research and Clinical Care** *JOURNAL OF PEDIATRICS*

Halley, M. C., Young, J. L., Tang, C., Mintz, K. T., Lucas-Griffin, S., Maghiro, A., Ashley, E. A., Tabor, H. K., Undiagnosed Diseases Network
2023; 261

- **How are students learning to care for people with disabilities?: exploring the curriculum design of a sample of disability electives offered by US health professions schools.** *Disability and rehabilitation*
  Clarke, L., Tabor, H. K., Gisondi, M. A.
  2023: 1-11

- **Genomics Research with Undiagnosed Children: Ethical Challenges at the Boundaries of Research and Clinical Care.** *The Journal of pediatrics*
  Halley, M. C., Young, J. L., Tang, C., Mintz, K. T., Lucas-Griffin, S., Maghiro, A. S., Ashley, E. A., Tabor, H. K.
  2023: 113537

- **Rare Disease, Advocacy and Justice: Intersecting Disparities in Research and Clinical Care.** *The American journal of bioethics : AJOB*
  Halley, M. C., Halverson, C. M., Tabor, H. K., Goldenberg, A. J.
  2023: 1-10

- **The impact of inclusion: Improving medical student confidence in caring for adults with intellectual disabilities through an interactive, narrative-based session** *JOURNAL OF INTELLECTUAL & DEVELOPMENTAL DISABILITY*
  Clarke, L., Tabor, H. K.
  2023

- **Characterizing Moral Injury and Distress in US Military Surgeons Deployed to Far-Forward Combat Environments in Afghanistan and Iraq.** *JAMA network open*
  Ryu, M. Y., Martin, M. J., Jin, A. H., Tabor, H. K., Wren, S. M.
  2023; 6 (2): e230484

- **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.
  2022; 24 (12): e42084

- **Latinx attitudes, barriers, and experiences with genetic counseling and testing: A systematic review.** *Journal of genetic counseling*
  Dron, H. A., Bucio, D., Young, J. L., Tabor, H. K., Cho, M. K.
  2022

- **Supporting undiagnosed participants when clinical genomics studies end.** *Nature genetics*
  Halley, M. C., Ashley, E. A., Tabor, H. K.
  2022

- **A Tale of Two Pandemics: Cancer Coping and Care Delivery During the COVID-19 Pandemic**
  Aslakson, R., Siden, R., Verano, M., Holdsworth, L., Braun, N., Tabor, H., Wong, B., Colborn, K., Smith, S., Lira, I., Hallahan, C., Nudotor, R., Siddiqi, et al
  ELSEVIER SCIENCE INC.2022: 1082-1083

- **An Evidence-Informed Cancer Care Communication Tool to Support Patients, Family, and Cancer and Palliative Care Clinicians During the COVID-19 Pandemic**
  Siden, R., Holdsworth, L., Wong, B., Verano, M., Tabor, H., Aslakson, R.
  ELSEVIER SCIENCE INC.2022: 1075

- **"Like Not Having an Arm": Perspectives of Patients, Caregivers, and Practitioners on the Impact of Visitor Restrictions on Cancer Care During the COVID-19 Pandemic**
  Holdsworth, L., Siden, R., Wong, B., Verano, M., Tabor, H., Aslakson, R.
  ELSEVIER SCIENCE INC.2022: 1085

- **A call for an integrated approach to improve efficiency, equity and sustainability in rare disease research in the United States.** *Nature genetics*
  Halley, M. C., Smith, H. S., Ashley, E. A., Goldenberg, A. J., Tabor, H. K.
  2022

- **Perceived utility and disutility of genomic sequencing for pediatric patients: Perspectives from parents with diverse sociodemographic characteristics.** *American journal of medical genetics. Part A*
  Halley, M. C., Young, J. L., Fernandez, L., Kohler, J. N., Undiagnosed Diseases Network, Bernstein, J. A., Wheeler, M. T., Tabor, H. K.
  1800

- **Beyond race: Recruitment of diverse participants in clinical genomics research for rare disease.** *Frontiers in genetics*
  Young, J. L., Halley, M. C., Anguiano, B., Fernandez, L., Bernstein, J. A., Wheeler, M. T., Tabor, H. K., Undiagnosed Diseases Network Consortium
  2022; 13: 949422

- **Genetic counselor roles in the undiagnosed diseases network research study: Clinical care, collaboration, and curation.** *Journal of genetic counseling*
  Kohler, J. N., Kelley, E. G., Boyd, B. M., Sillari, C. H., Marwaha, S., Undiagnosed Diseases Network, Wheeler, M. T., Acosta, M. T., Adam, M., Adams, D. R., Agrawal, P. B., Alejandro, M. E., Alvey, J., et al
  2021

- **Opportunities and pitfalls of social media research in rare genetic diseases: a systematic review.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Miller, E. G., Woodward, A. L., Flinchum, G., Young, J. L., Tabor, H. K., Halley, M. C.
  2021

- **"Doctors can read about it, they can know about it, but they've never lived with it": How parents use social media throughout the diagnostic odyssey.** *Journal of genetic counseling*
  Deuitch, N. T., Beckman, E., Halley, M. C., Young, J. L., Reuter, C. M., Kohler, J., Bernstein, J. A., Wheeler, M. T., Undiagnosed Diseases Network, Ormond, K. E., Tabor, H. K.
  2021

- **Cancer Care during Covid-19: A multi-institutional qualitative study on physician and patient perspectives on telemedicine.**
  Wong, B. O., Aslakson, R., Holdsworth, L. M., Siden, R., Tabor, H., Verano, M., Schapira, L.
  LIPPINCOTT WILLIAMS & WILKINS.2021

- **Genetic counseling and testing for Asian Americans: a systematic review.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Young, J. L., Mak, J., Stanley, T., Bass, M., Cho, M. K., Tabor, H. K.
  2021

- **Variants in PRKAR1B cause a neurodevelopmental disorder with autism spectrum disorder, apraxia, and insensitivity to pain** *GENETICS IN MEDICINE*
  Marbach, F., Stoyanov, G., Erger, F., Stratakis, C. A., Settas, N., London, E., Rosenfeld, J. A., Torti, E., Haldeman-Englert, C., Sklirou, E., Kessler, E., Ceulemans, S., Nelson, et al
  2021

- **Beyond diagnosis: understanding the downstream impacts of genome sequencing for undiagnosed rare diseases**
  Halley, M., Young, J., Tabor, H., Undiagnosed Dis Network
  ACADEMIC PRESS INC ELSEVIER SCIENCE.2021: S290

- **Commonalities across computational workflows for uncovering explanatory variants in undiagnosed cases.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Kobren, S. N., Baldridge, D., Velinder, M., Krier, J. B., LeBlanc, K., Esteves, C., Pusey, B. N., Zuchner, S., Blue, E., Lee, H., Huang, A., Bastarache, L., Bican, et al
  2021

- **"It seems like COVID-19 now is the only disease present on Earth": living with a rare or undiagnosed disease during the COVID-19 pandemic.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Halley, M. C., Stanley, T. n., Maturi, J. n., Goldenberg, A. J., Bernstein, J. A., Wheeler, M. T., Tabor, H. K.
  2021

- **Patient and family social media use surrounding a novel treatment for a rare genetic disease: a qualitative interview study.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Iyer, A. A., Barzilay, J. R., Tabor, H. K.
  2020

- **De novo EIF2AK1 and EIF2AK2 Variants Are Associated with Developmental Delay, Leukoencephalopathy, and Neurologic Decompensation.** *American journal of human genetics*
  Mao, D. n., Reuter, C. M., Ruzhnikov, M. R., Beck, A. E., Farrow, E. G., Emrick, L. T., Rosenfeld, J. A., Mackenzie, K. M., Robak, L. n., Wheeler, M. T., Burrage, L. C., Jain, M. n., Liu, et al
  2020

- **Ventilator Triage Policies During the COVID-19 Pandemic at U.S. Hospitals Associated With Members of the Association of Bioethics Program Directors.** *Annals of internal medicine*

Matheny Antommaria, A. H., Gibb, T. S., McGuire, A. L., Wolpe, P. R., Wynia, M. K., Applewhite, M. K., Caplan, A. n., Diekema, D. S., Hester, D. M., Lehmann, L. S., McLeod-Sordjan, R. n., Schiff, T. n., Tabor, et al

2020

- **Civic Engagement, Autism and Deliberative Democracy: Prioritizing the Inclusion of Marginalized Perspectives.** *The American journal of bioethics : AJOB*

  Tabor, H. K.

  2020; 20 (4): 41–43

- **Clinical sites of the Undiagnosed Diseases Network: unique contributions to genomic medicine and science.** *Genetics in medicine : official journal of the American College of Medical Genetics*

  Schoch, K. n., Esteves, C. n., Bican, A. n., Spillmann, R. n., Cope, H. n., McConkie-Rosell, A. n., Walley, N. n., Fernandez, L. n., Kohler, J. N., Bonner, D. n., Reuter, C. n., Stong, N. n., Mulvihill, et al

  2020

- **Rethinking the "open future" argument against predictive genetic testing of children** *GENETICS IN MEDICINE*

  Garrett, J. R., Lantos, J. D., Biesecker, L. G., Childerhose, J. E., Chung, W. K., Holm, I. A., Koenig, B. A., McEwen, J. E., Wilfond, B. S., Brothers, K., Berkman, B., Bernhardt, B., Caga-Anan, et al

  2019; 21 (10): 2190–98

- **De Novo Variants in WDR37 Are Associated with Epilepsy, Colobomas, Dysmorphism, Developmental Delay, Intellectual Disability, and Cerebellar Hypoplasia.** *American journal of human genetics*

  Kanca, O., Andrews, J. C., Lee, P., Patel, C., Braddock, S. R., Slavotinek, A. M., Cohen, J. S., Gubbels, C. S., Aldinger, K. A., Williams, J., Indaram, M., Fatemi, A., Yu, et al

  2019

- **"Before Facebook and before social media...we did not know anybody else that had this": parent perspectives on internet and social media use during the pediatric clinical genetic testing process** *JOURNAL OF COMMUNITY GENETICS*

  Barton, K. S., Wingerson, A., Barzilay, J. R., Tabor, H. K.

  2019; 10 (3): 375–83

- **Assessing genetic counselors' experiences with physician aid-in-dying and practice implications.** *Journal of genetic counseling*

  D'Angelo, A., Ormond, K. E., Magnus, D., Tabor, H. K.

  2019

- **Consent insufficient for data release.** *Science (New York, N.Y.)*

  Nicol, D., Eckstein, L., Bentzen, H. B., Borry, P., Burgess, M., Burke, W., Chalmers, D., Cho, M., Dove, E., Fullerton, S., Ida, R., Kato, K., Kaye, et al

  2019; 364 (6439): 445–46

- **High-Throughput Sequencing in Respiratory, Critical Care, and Sleep Medicine Research An Official American Thoracic Society Workshop Report** *ANNALS OF THE AMERICAN THORACIC SOCIETY*

  Hersh, C. P., Adcock, I. M., Celedon, J. C., Cho, M. H., Christiani, D. C., Himes, B. E., Kaminski, N., Mathias, R. A., Meyers, D. A., Quackenbush, J., Redline, S., Steiling, K. A., Tabor, et al

  2019; 16 (1): 1–16

- **Perspectives on Spinraza (Nusinersen) Treatment Study: Views of Individuals and Parents of Children Diagnosed with Spinal Muscular Atrophy.** *Journal of neuromuscular diseases*

  Pacione, M., Siskind, C. E., Day, J. W., Tabor, H. K.

  2018

- **"Before Facebook and before social mediawe did not know anybody else that had this": parent perspectives on internet and social media use during the pediatric clinical genetic testing process.** *Journal of community genetics*

  Barton, K. S., Wingerson, A., Barzilay, J. R., Tabor, H. K.

  2018

- **What Precision Medicine Can Learn from Rare Genetic Disease Research and Translation.** *AMA journal of ethics*

  Tabor, H. K., Goldenberg, A.

  2018; 20 (9): E834–840

- **Approaches to carrier testing and results disclosure in translational genomics research: The clinical sequencing exploratory research consortium experience.** *Molecular genetics & genomic medicine*

  Porter, K. M., Kauffman, T. L., Koenig, B. A., Lewis, K. L., Rehm, H. L., Richards, C. S., Strande, N. T., Tabor, H. K., Wolf, S. M., Yang, Y., Amendola, L. M., Azzariti, D. R., Berg, et al

2018

- **Ethical Challenges Confronted When Providing Nusinersen Treatment for Spinal Muscular Atrophy.** *JAMA pediatrics*
  Burgart, A. M., Magnus, D. n., Tabor, H. K., Paquette, E. D., Frader, J. n., Glover, J. J., Jackson, B. M., Harrison, C. H., Urion, D. K., Graham, R. J., Brandsema, J. F., Feudtner, C. n.
  2018; 172 (2): 188–92

- **Pathways from autism spectrum disorder diagnosis to genetic testing.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Barton, K. S., Tabor, H. K., Starks, H., Garrison, N. A., Laurino, M., Burke, W.
  2017

- **My46: a Web-based tool for self-guided management of genomic test results in research and clinical settings.** *Genetics in medicine*
  Tabor, H. K., Jamal, S. M., Yu, J., Crouch, J. M., Shankar, A. G., Dent, K. M., Anderson, N., Miller, D. A., Futral, B. T., Bamshad, M. J.
  2016

- **Gene discovery for Mendelian conditions via social networking: de novo variants in KDM1A cause developmental delay and distinctive facial features.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Chong, J. X., Yu, J. H., Lorentzen, P., Park, K. M., Jamal, S. M., Tabor, H. K., Rauch, A., Saenz, M. S., Boltshauser, E., Patterson, K. E., Nickerson, D. A., Bamshad, M. J.
  2016; 18 (8): 788-95

- **Clinical Sequencing Exploratory Research Consortium: Accelerating Evidence-Based Practice of Genomic Medicine** *AMERICAN JOURNAL OF HUMAN GENETICS*
  Green, R. C., Goddard, K. B., Jarvik, G. P., Amendola, L. M., Appelbaum, P. S., Berg, J. S., Bernhardt, B. A., Biesecker, L. G., Biswas, S., Blout, C. L., Bowling, K. M., Brothers, K. B., Burke, et al
  2016; 98 (6): 1051–66

- **Use of metaphors about exome and whole genome sequencing.** *American journal of medical genetics. Part A*
  Nelson, S. C., Crouch, J. M., Bamshad, M. J., Tabor, H. K., Yu, J.
  2016; 170A (5): 1127-1133

- **When Participants in Genomic Research Grow Up: Contact and Consent at the Age of Majority** *JOURNAL OF PEDIATRICS*
  Brothers, K. B., Holm, I. A., Childerhose, J. E., Antommaria, A. M., Bernhardt, B. A., Clayton, E., Gelb, B. D., Joffe, S., Lynch, J. A., McCormick, J. B., McCullough, L. B., Parsons, D., Sundaresan, et al
  2016; 168: 226-+

- **The Genetic Basis of Mendelian Phenotypes: Discoveries, Challenges, and Opportunities.** *American journal of human genetics*
  Chong, J. X., Buckingham, K. J., Jhangiani, S. N., Boehm, C., Sobreira, N., Smith, J. D., Harrell, T. M., McMillin, M. J., Wiszniewski, W., Gambin, T., Coban Akdemir, Z. H., Doheny, K., Scott, et al
  2015; 97 (2): 199-215

- **Exome Sequencing of Phenotypic Extremes Identifies CAV2 and TMC6 as Interacting Modifiers of Chronic Pseudomonas aeruginosa Infection in Cystic Fibrosis.** *PLoS genetics*
  Emond, M. J., Louie, T., Emerson, J., Chong, J. X., Mathias, R. A., Knowles, M. R., Rieder, M. J., Tabor, H. K., Nickerson, D. A., Barnes, K. C., Go, L., Gibson, R. L., Bamshad, et al
  2015; 11 (6): e1005273

- **Single-nucleotide polymorphism arrays and unexpected consanguinity: considerations for clinicians when returning results to families.** *Genetics in medicine : official journal of the American College of Medical Genetics*
  Delgado, F., Tabor, H. K., Chow, P. M., Conta, J. H., Feldman, K. W., Tsuchiya, K. D., Beck, A. E.
  2015; 17 (5): 400-4

- **Actionable exomic incidental findings in 6503 participants: challenges of variant classification** *GENOME RESEARCH*
  Amendola, L. M., Dorschner, M. O., Robertson, P. D., Salama, J. S., Hart, R., Shirts, B. H., Murray, M. L., Tokita, M. J., Gallego, C. J., Kim, D. S., Bennett, J. T., Crosslin, D. R., Ranchalis, et al
  2015; 25 (3): 305-315

- **"We Don't Know Her History, Her Background": Adoptive Parents' Perspectives on Whole Genome Sequencing Results** *JOURNAL OF GENETIC COUNSELING*
  Crouch, J., Yu, J., Shankar, A. G., Tabor, H. K.
  2015; 24 (1): 67-77

- **De novo mutations in NALCN cause a syndrome characterized by congenital contractures of the limbs and face, hypotonia, and developmental delay.** *American journal of human genetics*
  Chong, J. X., McMillin, M. J., Shively, K. M., Beck, A. E., Marvin, C. T., Armenteros, J. R., Buckingham, K. J., Nkinsi, N. T., Boyle, E. A., Berry, M. N., Bocian, M. n., Foulds, N. n., Uzielli, et al
  2015; 96 (3): 462–73

- **Pathogenic Variants for Mendelian and Complex Traits in Exomes of 6,517 European and African Americans: Implications for the Return of Incidental Results** *AMERICAN JOURNAL OF HUMAN GENETICS*
  Tabor, H. K., Auer, P. L., Jamal, S. M., Chong, J. X., Yu, J., Gordon, A. S., Graubert, T. A., O'Donnell, C. J., Rich, S. S., Nickerson, D. A., Bamshad, M. J.
  2014; 95 (2): 183-193

- **Attitudes of genetics professionals toward the return of incidental results from exome and whole-genome sequencing.** *American journal of human genetics*
  Yu, J. H., Harrell, T. M., Jamal, S. M., Tabor, H. K., Bamshad, M. J.
  2014; 95 (1): 77-84

- **Quantifying rare, deleterious variation in 12 human cytochrome P450 drug-metabolism genes in a large-scale exome dataset** *HUMAN MOLECULAR GENETICS*
  Gordon, A. S., Tabor, H. K., Johnson, A. D., Snively, B. M., Assimes, T. L., Auer, P. L., Ioannidis, J. P., Peters, U., Robinson, J. G., Sucheston, L. E., Wang, D., Sotoodehnia, N., Rotter, et al
  2014; 23 (8): 1957-1963

- **Whole-Exome Sequencing Identifies Rare and Low-Frequency Coding Variants Associated with LDL Cholesterol.** *American journal of human genetics*
  Lange, L. A., Hu, Y., Zhang, H., Xue, C., Schmidt, E. M., Tang, Z., Bizon, C., Lange, E. M., Smith, J. D., Turner, E. H., Jun, G., Kang, H. M., Peloso, et al
  2014; 94 (2): 233-245

- **Mutations in PIEZO2 cause Gordon syndrome, Marden-Walker syndrome, and distal arthrogryposis type 5.** *American journal of human genetics*
  McMillin, M. J., Beck, A. E., Chong, J. X., Shively, K. M., Buckingham, K. J., Gildersleeve, H. I., Aracena, M. I., Aylsworth, A. S., Bitoun, P. n., Carey, J. C., Clericuzio, C. L., Crow, Y. J., Curry, et al
  2014; 94 (5): 734–44

- **Parent perspectives on pediatric genetic research and implications for genotype-driven research recruitment.** *Journal of empirical research on human research ethics : JERHRE*
  Tabor, H. K., Brazg, T., Crouch, J., Namey, E. E., Fullerton, S. M., Beskow, L. M., Wilfond, B. S.
  2011; 6 (4): 41-52

- **Challenges in the use of direct-to-consumer personal genome testing in children.** *The American journal of bioethics : AJOB*
  Tabor, H. K., Kelley, M.
  2009; 9 (6-7): 32-4

- **Ethical implications of array comparative genomic hybridization in complex phenotypes: points to consider in research** *GENETICS IN MEDICINE*
  Tabor, H. K., Cho, M. K.
  2007; 9 (9): 626-631

- **Candidate-gene approaches for studying complex genetic traits: practical considerations** *NATURE REVIEWS GENETICS*
  Tabor, H. K., Risch, N. J., Myers, R. M.
  2002; 3 (5): 391-A396