CURRICULUM VITAE

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Name: Ellen Wright Clayton

Addresses:
Center for Biomedical Ethics and Society
2525 West End Ave., Suite 400
Nashville, TN 37203
(615) 322-1186

Vanderbilt University
School of Law
Nashville, TN 37240
(615) 322-1186

E-mail: ELLEN. CLAYTON@VANDERBILT.EDU

Education:

1970-74: B.S., Duke University
1974-76: M.S., Stanford University
1976-79: J.D., Yale Law School
1981-85: M.D., Harvard Medical School

Postgraduate Training:

1985-88: Resident, Department of Pediatrics, University of Wisconsin Hospital and Clinics, and Lecturer, Program in Medical Ethics

Licensure and Certification:

1988- : License to Practice Medicine, Tennessee
1989- : Certification, American Board of Pediatrics
1991- : Fellow, American Academy of Pediatrics

Academic Appointments:

1982-83: Research Assistant, Professor Jay Katz, Yale University, worked on his book, The Silent World of Doctor and Patient, Free Press, 1985

1985: Visiting Assistant Professor (Spring Semester), Law School and Program in Medical Ethics, University of Wisconsin

1988-96:	Assistant Professor of Pediatrics, Vanderbilt University Medical School Assistant Professor of Law, Vanderbilt University School of Law Fellow, Vanderbilt Institute for Public Policy Studies, Center for Health Policy
1993:	Visiting Professor of Law, University of Houston Law Center, Health Law and Policy Institute
1996-99:	Associate Professor of Pediatrics with tenure, Vanderbilt University Medical School Associate Professor of Law, Vanderbilt University School of Law Faculty Member, Vanderbilt Program in Human Genetics Senior Fellow, Vanderbilt Institute for Public Policy Studies, Center for Health Policy Associate, Center for Clinical and Research Ethics
1999-05:	Director, Center for Genetics and Health Policy
1999-2010:	Rosalind E. Franklin Professor of Genetics and Health Policy
2005-2011:	Director, Center for Biomedical Ethics and Society
Currently:	Craig-Weaver Professor of Pediatrics with tenure, Vanderbilt University Medical School Professor of Law, Vanderbilt University School of Law Professor of Health Policy, Vanderbilt University Medical School Co-Founder, Center for Biomedical Ethics and Society, Vanderbilt University School of Medicine Faculty Member, Vanderbilt Program in Human Genetics Senior Fellow, Vanderbilt Institute for Public Policy Studies, Center for Health Policy Senior Fellow, John F. Kennedy Center for Research on Human Development, Vanderbilt University Visiting Professor, Tokai University School of Medicine, Tokyo, JAPAN

Other Employment:

1979-80:	Law Clerk, The Honorable John C. Godbold, Judge U.S. Court of Appeals, Fifth Circuit
1980-81:	Attorney, Vinson and Elkins, Houston, TX, and Visiting Fellow, Institute for the Interprofessional Study of Health Law

Honors:

1970-73:	Class Honors, Freshman, Sophomore, and Junior years; Duke University National Merit Scholar
1974:	Phi Beta Kappa, Duke University Summa cum Laude, Duke University
1974-76:	National Institute of Health Traineeship, Stanford University
1978-79:	Editor, Yale Law Journal, Yale Law School
1985-86:	Henry M. Castello Memorial Award for the Outstanding Resident, University of Wisconsin Hospitals and Clinics
1997:	Sigma Xi, Vanderbilt University
1999:	Jay Healey Award for Outstanding Health Law Teacher
2004:	Excellence in Resident Education, Department of Pediatrics, Vanderbilt University School of Medicine
2006:	Member, Institute of Medicine
2007:	Founding Member, Vanderbilt Academy for Excellence in Teaching
2008:	Member, Alpha Omega Alpha
2010:	William G. Bartholome Award for Ethical Excellence, American Academy of Pediatrics
2012:	Fellow, American Association for the Advancement of Science
2013:	Member, American Pediatric Society
2013:	David P. Rall Medal, Institute of Medicine
2015:	Frank H. Morriss, Jr. Leadership Award, University of Iowa Department of Pediatrics and University of Iowa Foundation

Professional Memberships:

1980- :	State Bar of Texas and U.S. Court of the Southern District of Texas
1984- :	Member, American Society of Law, Medicine, and Ethics
1988- :	Member, American Bar Association
1989- :	Member, Lawyers Association for Women
1991- :	Member, Bioethics Section, American Academy of Pediatrics
1993- :	Member, Ambulatory Pediatric Association
1998- :	Member, American Society for Bioethics and Humanities
1998- :	Member, Health Law Section, American Bar Association
2010- :	Member, Public Population Project in Genomics

Vanderbilt Medical School/Medical Center Service:

1988-2008:	Member, Child Abuse Committee, Vanderbilt University
1991:	Member, Medicine in Society Committee
1991-93:	Consultant, Vanderbilt University Medical Ethics Committee
1993- :	Member, Vanderbilt University Medical Ethics Committee
1994-96:	Chair, Child Abuse Committee, Vanderbilt University
1996-2000:	Member, Credentials Committee, Vanderbilt University Medical Center
1997-2000:	Member, Joint Practice Subcommittee, Medical Staff Credentials Committee
1997-2000:	Member, Internal Review Committee, Graduate Education Review

Committee

1998: Member, Task Force on Compensation and Incentivization of Faculty (Education)

1998-2000: Member, Subcommittee on Protection of and Access to Patient Electronic Records

1998- 2001: Member, Software Oversight Committee

1998-2002: Member, Promotion Committee for the Class of 2002

1999- : Member, Ad Hoc Committee on Center for Fetal Intervention

1999-2001: Member, Academic Programs Committee

2000: Coordinator, Ecology of Medicine Course, sections on pregnancy and early childhood

2000- Member, Working Group on DNA Capture

> As part of this group, I oversaw the conduct of five focus groups to assess opinions of community members about collection of DNA in conjunction with medical information for use in epidemiologic research and continue to develop a program to develop the optimal approach to collecting DNA and medical records for research

2000- : Chair, Education on Research Integrity

2000-01: Chair, Conflict of Interest Committee

2001: Member, Emphasis Program and Synthesis Course Working Groups for Curriculum Reform

2001: Member, Search Committee for the Stahlman Chair in Medical Ethics

2001- : Member, Vanderbilt Child Advocacy Council

2001- : Faculty, Meharry-Vanderbilt Genetics Training Program

2002- : Member, Curriculum Advisory Committee, Meharry Clinical Research Education and Career Development (CRECD)

2002- : Member, Emphasis Program Planning Committee

2003- : Member, Steering Committee, PhD Program in Human Genetics

2003-5: Director, Genetics, second year medical school

2003-08: Director, Law and Policy Emphasis Program

2005- : Member, DNA Databank Oversight Committee

2008- : Member, Appointments and Promotions Committee, Department of Pediatrics

2009- : Member, Academic Advisory Council, Meharry-Vanderbilt Community Engaged Research Program, Vanderbilt Institute for Clinical and Translational Research

2009- : Member, Research Enterprise Communications Advisory Council

2009-11: Member, Appointments and Promotions Committee, VUMC

2011-12: Member, Executive Committee of the Executive Faculty, VUMC

2011- : Mentor, Hazinski Society, Department of Pediatrics

Vanderbilt Law School Service:

1988-2005: Faculty Advisor, Vanderbilt Bisexual, Gay, and Lesbian Law Students Association

1988-2005: Faculty Advisor, Vanderbilt Law Women's Association
1989-91: Member, Curriculum Committee, Vanderbilt Law School
1991-95: Member, Student-Faculty Relations Committee, Vanderbilt Law School
1995-97: Member, Diversity Lectures Committee
1997-05: Member, Admissions Committee

Vanderbilt University Service:

1989-93: Member, Advisory Board, Margaret Cuninggim Women's Center, Vanderbilt University
1998-2001: Member, Advisory Committee, Cal Turner Program in Moral Leadership
2000-01: Organizer, Seminar on Genetics and Mental Health, Center for Genetics and Health Policy and Robert Penn Warren Center for the Humanities
2001-09: Member, Steering Committee, Medicine, Health, and Society -- an interdisciplinary program that draws from many parts of the university
2001- 07: Member, Executive Board, Cal Turner Program in Moral Leadership
2002- : Member, Conflict of Interest Committee
2005-09 : Member, Technology Review Committee
2006-09: Member, Advisory Board, Center for Ethics
2006-09: Member, Faculty Advisory Board, Vanderbilt Center for Nashville Studies
2009-10: Member, Search Committee, Directors of Center for Medicine, Health, and Society and Program in Health, Environment, and Policy

Vanderbilt Health Plan Service:

1996-1997 Member, Pharmacy and Therapeutics Committee
1997-99: Member, Medical Resource Management Committee

Professional Service on National and International Level:

Institute of Medicine

1992: Consultant to Institute of Medicine Committee on Assessing Genetic Risks: Issues and Implications for Health, June 28-29, 1992, Irvine, CA
1993: Consultant, Institute of Medicine Committee on the Legal and Ethical Issues Relating to the Inclusion of Women in Clinical Studies
2002: Invited Speaker, Genomics and Its Implications for Human Health: A New Academies Initiative
2002-07: Member, Board on Health Sciences Policy
2003-04: Member, Committee on the Use of Third Party Toxicity Research with Human Research Participants, Science, Technology, and Law Program
2004-05: Member, Committee on Genomics and the Public's Health in the 21st Century

2005: Chair, Roundtable on Translating Genomic Information into Improved Health
2005-06: Member, Committee on the Disposition of the Air Force Health Study
2005-06: Member, Committee on Assessing Interactions Among, Social, Behavioral, and Genetic Factors of Health
2007: Member, Panel to Review the National Children's Study (NCS) Research Plan
2007-09: Chair, Committee to Evaluate Title X Family Planning
2009-11: Chair, Committee to Evaluate Vaccine Safety
2010- : Member, Advisory Council
2011-12: Chair, Standing Committee on Family Planning
2012- 13: Co-Chair, Committee on Commercial Sexual Exploitation and Sex Trafficking of Minors in the United States
2012- : Chair, Board on Population Health and Public Health Practice
2012: Member, Committee to Revamp "The Role of the Chair" of Consensus Committees
2012: Monitor, Reforming Juvenile Justice: A Developmental Approach
2012: Reviewer, The Appropriate Future Use of the Department of Defense Joint Pathology Center Tissue Repository
2012: Review Coordinator, Review of the California Institute for Regenerative Medicine (CIRM)
2013- : Executive Committee, Council, IOM
2013: Member, Report Review Committee, IOM
2013-14: Monitor, Integrating Workforce Health Protection at DHS
2013-15: Chair, Diagnostic Criteria for Myalgic Encephalomyelitis/Chronic Fatigue Syndrome
2014: Monitor, Treatment for Posttraumatic Stress Disorder in Military and Veteran Populations: Final Assessment
2014: Reviewer, Investing in the Health, Safety, and Well-Being of Young Adults
2014-15: Chair, Governance Committee
2014: Council Presentations (with Bill Stead), Grand Challenges in Health and Health Care

National Institutes of Health, National Human Genome Research Institute:

1994-1995: Chair, NIH-CDC Workshop on the Role of Informed Consent for Genetic Research on Stored Tissue Samples
1995-98: Member, National Advisory Council, National Human Genome Research Institute, National Institutes of Health
1997-2000: Member, ELSI Research Program and Evaluation Group, National Human Genome Research Institute
2000-01: ELSI Research Advisor
2001: Chair, Session on ELSI Issues at Developing a Haplotype Map of the Human Genome for Finding Genes Related to Health and Disease
2001-03: Co-Chair, Population/ELSI Group, International Haplotype Mapping Project

2003-05: Co-Chair, ELSI Group, International Haplotype Mapping Project
2007-11: Co-Chair, Consent and Community Consultation Working Group, eMERGE Network

National Institutes of Health -- other programs:

1990: Consultant, National Institutes of Health, Review Committee on Legal and Ethical Issues of Human Genome Initiative, Washington, D.C.
1991: Working Group on Reproductive Genetic Testing: Impact on Women, National Institutes of Health, Bethesda, MD
1996: Member, Workshop on the Inclusion of Children in Clinical Research, National Institute of Child Health and Human Development, American Academy of Pediatrics
2000- : Member, Workgroup on Ethical Issues in Pharmacogenomics, Clinical Ethics Section, Bethesda, MD
2002-05: Member, Ethics Working Group, National Children's Study, National Institute of Child Health and Human Development
2007: Group Leader, Custodianship and Ownership Issues in Biospecimen Research Symposium-Workshop, National Cancer Institute
2009-10: Member, National Children's Study Federal Advisory Committee, Eunice Kennedy Shriver National Institute of Child Health and Human Development
2010: Chair, National Cancer Institute, Workshop on Release of Research Results to Participate in Biospecimen Studies
2011- : Member, Observational Safety Monitoring Board, NHLBI OSMB for Next Generation Genetic Association Studies
2015- : Member, Expert Advisory Panel, NHLBI Trans-Omics for Precision Medicine Whole Genome Sequencing

Food and Drug Administration:

1999- : Consultant, Pediatrics Advisory Committee
2000: Consultant, Clinical pharmacology during pregnancy: Addressing clinical needs through science

Human Genome Organisation:

2010-13: Member, Working Group on Whole Genome Sequencing
2010-12: Member, Ethics Committee
2012- : Member, Committee on Ethics, Law, and Society

Public Population Project in Genomics and Society

2013- : Chair, Paediatric Platform

American Society of Law, Medicine, and Ethics:

1993-98: Member, Board of Directors, American Society of Law, Medicine, and Ethics
1994-95: Treasurer, American Society of Law, Medicine, and Ethics
1995-96: President-elect, American Society of Law, Medicine, and Ethics
1996-7: President, American Society of Law, Medicine, and Ethics
1997-2001: Editor-in-Chief, Journal of Law, Medicine, and Ethics
2003- : Member, Board of Editors, Journal of Law, Medicine, and Ethics

American Bar Association:

1998-2001: Chair, Clinical Ethics Issues Interest Group, Health Law Section

Executive Leadership in Academic Medicine:

1999-2000: Participant

National and International Committees and Workshops Not Listed Above:

1987-88: Workgroup on Reproductive Issues, MacArthur Foundation Research Group on Health Behavior, Key Biscayne, FL, and Washington, D.C.
1987: Workgroup on Genetic Screening, Hastings Center, Briarcliff Manor, NY
1989-95: Member, American Academy of Pediatrics Committee on Bioethics
1989: Consultant, American Academy of Pediatrics, Provisional Committee on Substance Abuse, Special Meeting on Drug Affected Infants
1990: Workgroup on Feminism and Bioethics, Hastings Center, Briarcliff Manor, NY
1992- : Member, Technical Advisory Board, National Emergency Medical Services for Children Resource Alliance, Torrance, CA
1993-95: Member, Research Group, "Setting Priorities in the Application of Human Genetic Research," Hastings Center, Briarcliff Manor, NY
1995: Member, Storage of Genetics Materials Committee, American College of Medical Genetics
1995-98: Member, Biological Resources Bank Working Group, National Action Plan on Breast Cancer
1996-99: Member, Social Issues Committee, American Society of Human Genetics
1997- : Member, National Advisory Committee, Pediatric Palliative Care Committee
1997-2000: Member, Advisory Committee, Center for Patients' Rights
1998: Consultant, New York State Task Force on Life and the Law, Taskforce on Genetics Research
1999-2000: Member, American Academy of Pediatrics Newborn Screening Task Force
1998-2001: Member, Expert Advisory Panel on Privacy and Confidentiality, American Medical Association Institute for Ethics
1999: Member, Classification-Nomenclature Group, EASL International Consensus Conference on Hemochromatosis, Sorrento, ITALY

1999- : Member, External Advisory Board, Public Health Genetics Program, University of Washington
2000-02: Member, Ethics and Security Advisory Board, Marshfield Clinic Personalized Medicine Program
2000-02: Consultant, Council of International Organizations of Medical Sciences, Geneva, SWITZERLAND
2000- : Member, Human Biological Materials Focus Group, Adult AIDS Clinical Trials Group
2001-03: Member, Episcopal Task Force on Issues in Human Genetics
2002-05: Member, Evaluating Genetic Tests: A Models Process, University of Washington
2003: Consultant, ELSI Genetic Variation Consortium, Colliding Categories: Haplotypes, Race, and Ethnicity, University of Minnesota
2003: Participant, Newborn Screening for Cystic Fibrosis Meeting, National Center on Birth Defects and Developmental Disabilities (NCBDDD) at the Centers for Disease Control and Prevention (CDC) and the Cystic Fibrosis Foundation (CFF)
2003-05: Member, Ethical Decision Making for Newborn Genetic Screening, Hastings Center, New York
2004-06: Chair, American Society of Human Genetics Working Group on HIPAA and Genetics Research
2004-09: Member, Ethics Review Panel, Shanghai Health Study
2005-08: Member, Fellows Council, Hastings Center
2005: Chair, Nominating Committee, AAP Section on Bioethics
2006-07: Chair, AALS and ASRM Planning Committee for Workshops on Reproductive Medicine and the Law
2009-12: Member, External Advisory Board, ESRC Centre for Genomics in Society, University of Exeter, UK
2014- : Member, External Advisory Board, CITI Japan

Professional Service on State Level:

1991- : Member, Genetics Advisory Committee, Tennessee Department of Health and Environment
1992- : Chair, Committee on Bioethics, Tennessee Chapter American Academy of Pediatrics
1993- : Member, Tennessee Lawyers Association for Women Judicial Appointments and Elections Committee
1996-2000: Member, Family Law Committee -- Code Revision Committee, Tennessee Bar Association

1997-2000:	Member, Board of Directors, American Civil Liberties Union of Tennessee
2001- :	Chair, Ethics/Confidentiality Committee, Statewide Genetics Planning Committee
2004- :	Chair, Government Relations Committee, Genetics Advisory Committee

Service at Other Universities:

1978-79:	Educational Planning Committee, Yale Law School; Treasurer, Yale Law Women's Association, Yale Law School
1982-84:	Member, Genetics Discussion Group, Harvard Medical School
1983-84:	Director, Medicine and Society, Harvard Medical School
1986-88:	Moderator, Ethics Forums, University of Wisconsin Hospital and Clinics
1986-88:	Member, Ethics Discussion Group, University of Wisconsin
1986-88:	Human Subjects Committee, University of Wisconsin
1987-88:	Ethics Committee, VA Hospital, Madison, WI

Community Service:

1989- :	Member, Davidson County Child Abuse Team
1995- :	Member, Board of Directors, Tennessee Health Decision Ethics Guild
1996-2002:	Member, Nashville Opera Guild
2012- :	Member, Board of Directors, Planned Parenthood of Middle and East Tennessee
2013- :	Member, Board of Directors, Tennessee Advocates for Planned Parenthood

Editorial Board:

Genomics, Society and Policy
Journal of Law, Medicine, and Ethics

Teaching Interests:

General Pediatrics; Biomedical Ethics and Law; Medical Ethics; Legal and Ethical Issues Raised by Genetics; Genetics and Health Policy; Genetics; Research Ethics; Public Health Law; Legal and Ethical Issues in Children's and Women's Health; Family Law; Gender and the Law

Teaching Experience:

1973-74:	Teaching Assistant, graduate course in cellular physiology, Duke University
1974-76:	Teaching Assistant, Introductory Biochemistry and Human Genetics, Stanford University
1982-84:	Lecturer in Introductory Genetics (Spring), Harvard Medical School
1984-85:	Seminar in Law and Medicine (Spring), University of Wisconsin Law School

1988- : Family Law, Gender and the Law, Bioethics and Law, Torts, Health Law and Policy, Genetics and the Law, Seminar on the Health Care Safety Net, Genetics, Medicine, and Law, Vanderbilt University School of Law

1988- : Within the medical school, I have taught in various capacities. I have assisted a number of medical students with emphasis projects over the years. I am currently developing part of the ethics curriculum for Curriculum 2.0. The multidisciplinary courses that I teach in the law school are open to medical students. I teach in the MSCI and other graduate programs. I directed a Hazinski mentoring group in pediatrics and give grand rounds in pediatrics approximately once a year.

1990- : In the last several years, I have guest lectured in classes in the divinity, business, nursing, and engineering schools, the Kennedy Center, and in the college of arts and sciences on an array of issues in law, medicine, and ethics. I currently do several such presentations every semester.

Grant and Funding Support:

Co-Investigator, Birth Outcomes, Satisfaction with Care, and Malpractice, Agency for Health Care Policy and Research (Frank A. Sloan & Gerald B. Hickson, Principal Investigators) HS06499 ($660,538) (9/30/90-3/31/94)

Co-Investigator, Cystic Fibrosis Screening: An Alternative Paradigm, National Institutes of Health 1R01 HG 00638-02 (John A. Phillips, III, Principal Investigator) ($386,271) (9/30/91-6/30/95)

Principal Investigator, Charles E. Culpeper Foundation Scholarship in Medical Humanities ($90,000) (7/1/92-6/30/95)

Principal Investigator, Shannon Award, National Center for Human Genome Research ($100,000) (1995-1997)

Consultant, Psychosocial Effects of Neonatal Hearing Screening, National Institutes of Mental Health 1R03 MH53965-01 (Anne Marie Tharpe, Principal Investigator) ($32,100)(~1995-1997)

Contract, Informed Consent for the Use of Stored Tissue Samples for Genetic Research, National Center for Human Genome Research (Ellen Wright Clayton, Principal Investigator) ($5,000) (1995)

Contract, Informed Consent for Creation of DNA Libraries for Large Scale Sequencing, Department of Energy (Ellen Wright Clayton, Principal Investigator) ($2,500) (1996)

Contract, Inclusion of Children in Research, National Institute of Child Health and Human Development (Ellen Wright Clayton, Principal Investigator) ($2,500) (1996)

Contract, Services Grant for Children with Genetic Conditions, Tennessee Department of Health (~$2,000/year) (1999)

Principal Investigator, Public Health and Genetics, National Human Genome Research Institute, 1 R01 HG01974-01 ($103,168) + a supplement for the Public Dialogue on Human Gene Patenting (1999-2002)

Principal Investigator, Japan HapMap Community Engagement, National Human Genome Research Institute ($68,342) (2002-2003)

Co-Investigator, Primary Pulmonary Hypertension: Mechanisms and Family Registry, National Heart, Lung, and Blood Institute (1998-2008)(Jim Loyd, Principal Investigator) ($83,000) (08/03-3/09) – 19% effort

Co-Investigator Genetics in Literature, Film, and Popular Culture, National Human Genome Research Institute, R03 HG-03031-01 (Jay Clayton, Principal Investigator) ($100,000) (2003-2005)

Co-Investigator, Clinical Use of Research Genetic Tests for Arrhythmia, National Human Genome Research Institute, R03 HG-2545-01, (Carol Freund, Principal Investigator) ($100,000) (2003-2005) – 2% effort

Principal Investigator, Impact of Newborn Screening on Families, March of Dimes, 6-FY05-85 (6/1/05-5/31/08) ($98,194) – 0% effort

Co-Principal Investigator, Vanderbilt Engaging Nashville Survey (Daniel Cornfield, Principal Investigator) (2007) – 0% effort

Principal Investigator, Religion and Genomics: Navigating Pathways and Perspectives of Patient Care, National Human Genome Research Institute, 1 R13 HG004689-01 ($31,900) (03/08- 02/09) – 0% effort

Co- Investigator, Fogarty "Creating Collaborative Research Ethics Education with Costa Rica," Elizabeth Heitman, PI (09/06-06/11) – 4% effort

Co-Investigator, NHLBI "Treatment of Pulmonary Edema in Organ Donors" Lorraine Ware, PI (04/07-03/11) – 1% effort

Co-Investigator, NHGRI "Vanderbilt Genome-Electronic Records Project" Dan Roden, PI (09/07-07/11) – 15% effort

Co- Investigator, NCR "The Vanderbilt Institute for Clinical and Translational Research (VICTR)" Gordon Bernard, PI (09/07-05/12) – 5% effort

Principal Investigator, NHGRI, "Returning Results of Pediatric Genomic Research to Participants," (9/30/2011-9/29/2013) -- 8% effort, 20% effort year 2

Co-Investigator, NHGRI "Vanderbilt Genome-Electronic Records Project" Dan Roden, PI (8/12-7/16) – 5% effort

Co-Investigator, NIGMS, “A Risk Management Framework for Identifiability in Genomics Research,” Brad Malin PI (09/30/2012 – 09/29/2016) – 10% effort

Publications:

Articles:

1. Ellen E. Wright, Father and Mother Know Best: Defining the Liability of Physicians for Inadequate Genetic Counseling, Yale Law Journal 1978; 87:1488-515

2. Ellen E. Wright, The Legal Implications of Refusing to Provide Prenatal Diagnosis in Low-Risk Pregnancies or Solely for Sex Selection, American Journal of Medical Genetics 1980; 5:391-97

3. Ellen E. Wright & Margery W. Shaw, Legal Liability in Genetic Screening, Genetic Counseling, and Prenatal Diagnosis, Clinical Obstetrics and Gynecology 1981; 24:1133-49; reprinted in Genetic Disease Diagnosis and Treatment: Proceedings of the Fifth Arnold O. Beckman Conference in Clinical Chemistry 17-40 (A.A. Dietz, ed., 1983)

4. Ellen Wright Clayton, Legal Aspects of Medical Genetics in Wisconsin, Wisconsin Medical Journal 1985; 84:28-33

5. Ellen Wright Clayton, From Rogers to Rivers: The Rights of the Mentally Ill to Refuse Medication, American Journal of Law and Medicine 1987; 13:7-52

6. Ellen Wright Clayton & Jay Clayton, Voices and Violence, Vanderbilt Law Review 1990; 43:1807-1818

7. Ellen Wright Clayton & Gerald B. Hickson. Compensation Under the National Childhood Vaccine Injury Act, Journal of Pediatrics 1990; 116:508-513

8. Gerald B. Hickson, William A. Altemeier & Ellen Wright Clayton, Should Promethazine Be Available Without Prescription? Pediatrics 1990; 86:221-25

9. Ellen Wright Clayton, Screening and Treatment of Newborns, Houston Law Review 1992; 29:85-148

10. Gerald B. Hickson, Ellen Wright Clayton, Penny B. Githens, & Frank A. Sloan, Factors That Prompted Families to File Medical Malpractice Claims Following Perinatal Injuries, Journal of the American Medical Association 1992; 267:1359-1363

11. Ellen Wright Clayton, Issues in State Newborn Screening Programs, Pediatrics 1992; 90:641-645

12. Ellen Wright Clayton, A Ray of Light About Frozen Embryos, Kennedy Institute Ethics Journal 1992; 2:347-359

13. Ellen Wright Clayton, Reproductive Genetic Testing: Regulatory and Liability Issues, Fetal Diagnosis and Therapy 1993; 8 (Suppl 1):39-59

14. Gerald B. Hickson, Ellen Wright Clayton, Are You and Your Waiting Room's Televised "Expert" Saying the Same Thing? Clinical Pediatrics 1993; 32:172-174

15. Vickie L. Hannig, Ellen Wright Clayton, & Kathryn M. Edwards, Whose DNA Is It Anyway?: Relationships between Families and Researchers. American Journal of Medical Genetics 1993; 47:257-260

16. Ellen Wright Clayton, Newborn Screening for the Pediatrician, Contemporary Pediatrics 1993; 10:34-46

17. Ellen Wright Clayton, Gerald B. Hickson & Cynthia S. Miller, Parents' Responses to Vaccine Information Pamphlets. Pediatrics 1994; 93:369-372

18. Gerald B. Hickson, Ellen Wright Clayton, Stephen S. Entman, Cynthia S. Miller, Penny B. Githens, Kathryn Whetten-Goldstein, Frank A. Sloan, Obstetricians' prior malpractice experience and patients' satisfaction with care, Journal of the American Medical Association 1994; 272: 1583-1587

19. Ellen Wright Clayton, What are the law's priorities about the dispersion of genetic technologies? Special Supplement, Hastings Center Report 1995; 25(3):S13-S15

20. Ellen Wright Clayton, Removing the shadow of the law from the debate about genetic testing of children, American Journal of Medical Genetics 1995; 57: 630-634

21. Ellen Wright Clayton, What is really at stake in Baby K: A Response to Ellen Flannery, Journal of Law, Medicine, and Ethics 1995; 23:13-14

22. Ellen Wright Clayton, Vickie L. Hannig, Jean P. Pfotenhauer, Robert A. Parker, Preston W. Campbell, III, John A. Phillips, III. Teaching about cystic fibrosis carrier screening by using written and video information, American Journal of Human Genetics 1995; 57: 171-181

23. Ellen Wright Clayton, Karen K.Steinberg, Muin J. Khoury, Elizabeth Thomson, Lori Andrews, Mary Jo Ellis Kahn, Loretta M. Kopelman, Informed consent for genetic research on stored tissue samples, Journal of the American Medical Association 1995; 274: 1786-1792

24. Ellen Wright Clayton, Panel Comment: Why the use of anonymous samples for research matters, Journal of Law, Medicine, and Ethics 1995; 23: 375-377.

25. Ellen Wright Clayton, Vickie L. Hannig, Jean P. Pfotenhauer, Robert A. Parker, Preston W.

Campbell, III, John A. Phillips, III, Lack of interest by nonpregnant couples in population based cystic fibrosis carrier screening, American Journal of Human Genetics 1996; 58: 617-627

26. Ellen Wright Clayton, Pediatrics ethics: A view from the United States, Acta Paediatrica Japonica 1996; 38: 299-303

27. Ellen Wright Clayton, Problems posed by genetics for law and ethics: American policies, Annual Review of Law and Ethics 1996; 4: 3-19

28. Ellen Wright Clayton, Potential Liability in Cases of Child Abuse and Neglect, Pediatric Annals 1997; 26: 173-178

29. Ellen Wright Clayton, Genetic Testing in Children, Journal of Medicine and Philosophy 1997; 33: 233-251

30. Ellen Wright Clayton, The Dangers of Reading *Duty* Too Broadly, Journal of Law, Medicine, and Ethics 1997; 25: 19-21

31. Anne Marie Tharpe, Ellen Wright Clayton, Newborn Hearing Screening: Issues in Legal Liability and Quality Assurance, American Journal of Audiology 1997; 6: 5-12

32. Gerald B. Hickson, James Pichert, Charles Federspiel, Ellen Wright Clayton, Development of an Early Identification and Response Model of Malpractice Prevention, Law and Contemporary Problems 1997; 60:7-29

33. Gerald B. Hickson, Ellen Wright Clayton, Cynthia S. Miller, James W. Pichert, Stephen S. Entman, Satisfaction with Obstetrical Care: Relation to Neonatal Intensive Care, Obstetrics and Gynecology 1998; 91:288-92

34. Ellen Wright Clayton, What Should the Law Say About Disclosure of Genetic Information to Relatives, Journal of Health Care Law & Policy 1998; 1: 373-390

35. Ellen Wright Clayton, Genetic Testing in Children: Law's Role in Addressing the Challenges Posed by New Technology, Children's Legal Rights Journal 1998; 18:39-42

36. Wylie Burke, Elizabeth Thomson, Muin J. Khoury, Sharon M. McDonnell, Nancy Press, Paul C. Adams, James C. Barton, Ernest Beutler, Gary Brittenham, Allen Buchanan, Ellen Wright Clayton, Mary E. Cogswell, Eric M. Meslin, Arno G. Motulsky, Lawrie W. Powell, Elliott Sigal, Benjamin S. Wilfond, Francis S. Collins, Hereditary Hemochromatosis: Gene Discovery and Its Implications for Population-Based Screening, Journal of the American Medical Association 1998; 280: 172-178

37. Ellen Wright Clayton, What Should Be the Role of Public Health in Newborn Screening and Prenatal Diagnosis? American Journal of Preventive Medicine 1999; 16: 111-5

38. Ellen Wright Clayton, Lessons (Re)Learned from Cystic Fibrosis Carrier Screening, Journal of Gender-Specific Medicine 1999; 2: 21-24

39. Ellen Wright Clayton, Using Newborn Blood Samples for Genetics Research: Thinking About the Context, Annual Review of Law and Medicine 2001; 9: 21-31

40. Ellen Wright Clayton, Through the Lens of the Sequence, Genome Research 2001; 11: 659-64.

41. Ellen Wright Clayton, You Have to Protect Physician Reporters if You Want to Protect Children from Abuse and Neglect, Houston Journal of Health Law and Policy 2001; 1: 133-146

42. Ellen Wright Clayton, Regulating Germ-Line Gene Transfer in the United States, Annual Review of Law and Ethics 2002; 10: 147-60

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"Regulaciones de marcadores genéticos en la investigación," II Simposio Internacional de Bioética, Hospital Nacional de Niños, San José, COSTA RICA, March, 2008

"Ten Fingers, Ten Toes," Newborn Screening Conference, Case Western Reserve University Law-Medicine Center, Cleveland, OHIO, April, 2008

"Where Should We Go with Newborn Screening?" Harvey and Bernice Jones Lecture, Arkansas Children's Hospital, Little Rock, AK, December, 2008

"Pros and Cons of Screening for Untreatable Disorders," American College of Medical Genetics annual meeting, Tampa, FL, March 2009

"Why Parents Should Not Have the Authority to Consent on Behalf of Their Children, but Adolescents May Consent for Themselves," American College of Medical Genetics annual meeting, Tampa, FL, March 2009

“Return of Results in Genet(om)ics Research,” David Green Memorial Lecture, University of Utah, April, 2009

“Regulating Clinical Genomics,” Yale Law School Reunion, New Haven, CT, October, 2009

“Ethical Issues in Moving Clinical Genomics from the Bench to the Bedside,” Medical Genome Science in the Personal Genome Era, International Symposium on Applied Genomics 2009, TOKYO, JAPAN, December, 2009

“Clinical Genetics and Identity,” Genomics, Medicine and the Humanities Workshop, University of Exeter, EXETER, UK, March, 2010

“Governance Issues in Whole Genome Diagnostics,” Changing Landscape of Genetic Testing and Its Impact on Clinical and Laboratory Services and Research in Europe, GOTEBORG, SWEDEN, June, 2010

“Ethical Issues in Expanded Newborn Screening,” European Society of Human Genetics, GOTEBORG, SWEDEN, June, 2010

“Newborn Screening: Informed Consent Through the Back Door,” Is Medical Ethics Really in the Best Interest of the Patient?” UPPSALA, SWEDEN, June, 2010

“Newborn Screening: The Canary in the Genomics Mine” and “Dealing with Genomics in the Clinic,” Hagop S. Mekjian Lectures in Medical Ethics and Professionalism, Ohio State University, Columbus, OH, September, 2010

“What are Advances in Genomics Really Going to Mean for Patients? Issues in Translation, Hartnell College, Salinas, CA, October, 2010

“Surfing the Tsunami of Whole Genome Sequencing,” Pacific Symposium on Biocomputing, Kona, HI, January, 2011

“Surfing the Tsunami of Whole Genome Sequencing,” Advances in Genome Biology and Technology, Marco Island, FL, February, 2011

“Challenges Posed by New Technologies for the Standard of Care,” Personalized Medicine Symposium, Birmingham, AL, February, 2011

“Challenges Posed by Translating Whole Genomes to the Clinic,” Presidential Commission for the Study of Bioethical Issues, Washington, DC, February, 2011 and HUGO 15th Human Genome Meeting: Genomics of Human Diversity and Heritable Disorders, DUBAI, UAE, March, 2011

"Imagined Futures: Capturing the Benefits of Genome Sequencing for Society," HUGO 15th Human Genome Meeting: Genomics of Human Diversity and Heritable Disorders, DUBAI, UAE, March, 2011

"What should we do with whole genomic sequences in the clinic and in research?" Personal Genomes, Cold Spring Harbor, NY, October, 2011

"Returning Results of Genome Wide Tests in Research and the Clinic – Addressing the Issues" Sanger Center and HeLEX, Oxford, UNITED KINGDOM, November, 2011

"Consent," Human Microbiome Project, Houston, TX, March, 2012

"Whole Genome Sequencing in the Clinic," Law Gene Centre, University of Tasmania Law School, Hobart, AUSTRALIA, March, 2012

"Imagined Futures: Capturing the Benefits of Genome Sequencing for Society," HUGO Annual Meeting, Sydney, AUSTRALIA, March, 2012

"How Not to Drown in the Whole Genome in the Clinic," Pediatric Academic Societies Meeting, Boston, MA, April, 2012

"Ethical Issues in Non-Invasive Prenatal Diagnosis," IOM Annual Meeting, Interest Group, Washington, DC, October, 2012

"Pediatricians Beware! Direct to Consumer Marketing of Genetic Testing for Children," AAP Annual Meeting, New Orleans, LA, October, 2012

"The Role of Genomics in Primary Prevention," University of Michigan, Ann Arbor, MI, March, 2013

"Genomics in Primary Prevention," Greenwall Fellows Program, Chicago, IL, May, 2013

"Transdisciplinarity – Why it matter for the Future, Exeter University, UNITED KINGDOM, June, 2013

"Who Should Decide About Genetic Testing in Kids? A Response to the ACMG," Bioethics Seminar, Seattle Children's Hospital, October, 2013

"Major Policy Questions in Genomic Medicine," University of Washington Center for Genomics and Health Care Equality, 2013 CGHE All-Investigators' Meeting, Seattle, WA, October, 2013

"Return of results in research and the clinic – Getting back on the right track," University of Pittsburgh Law School, Pittsburgh, PA, October 2013

"Challenging the Duty to Hunt," ASHG Annual Meeting, Boston, MA, October, 2013

"Newborn Screening: From Guthrie to the Uniform Panel and Beyond," AAP Annual Meeting, Orlando, FL, October, 2013

"From Peapods to Whole Genome Sequencing: Old and New Ethical Challenges," AAP Annual Meeting, Orlando, FL, October, 2013

"Challenges Posed by the ACMG Recommendations on Incidental Findings in Clinical Genomic Sequencing," Eighth Annual Cynthia Jean Stolman Lecture in Medical Ethics, Rutgers New Jersey Medical School, Newark, NJ, November, 2013

"What findings, if any, from archival data and samples should be offered to research participants? Some reflections and practical issues in the US," Returning Genetic Results in Biobanks: Opening an International Dialogue," Brocher Foundation, Hermance, Switzerland, November, 2013

"Return of Results? No, Unless," HandsOn Biobank Annual Meeting, The Hague, NETHERLANDS, November, 2013

"What to Ask Your Doctor About Genetic (and Other Complex) Tests," Policy Forum: Policy implications of genomic sequencing for individual and population health, Brown School, Washington University, St. Louis, MO, February, 2014

"Whole Genome Sequencing in Newborn Screening?," Human Genome Meeting 2014, Geneva, SWITZERLAND, April, 2014

"What Should You Know and Be Doing About Genome Privacy?" Workshop on Genome Privacy held in conjunction with the 14th Privacy Enhancing Technologies Symposium (PETS 2014), Amsterdam, NETHERLANDS, July, 2014

"On the Cusp of Adulthood: Ethical and Legal Issues Raised by Genomics Research and Care Involving Adolescents," Grand Rounds, St. Jude Research Hospital, Memphis, TN, January, 2015

"Current Controversies in Whole Genome Sequencing," Challenging the Norm: Fostering Ethics, University of Wisconsin, March, 2015

"When Participants in Genomic Research Grow Up: Contact and Consent at the Age of Majority," Annual Meeting, Paediatric Platform, P3G, June, 2015, Montreal, CANADA

Ethical and Legal Issues in Research and Patient Care (often includes genetics):

"Informed Consent for Genetic Research Using Stored Tissue Samples," The Responsibility of Oversight in Genetic Research and Technology Assessment, Public Responsibility in

Medicine and Research Annual Convention, Boston, MA, May, 1996

"Genetic Screening and Research," First International Conference on DNA Sampling -- Human Genetic Research: Ethical, Legal and Policy Aspects, Montreal, Quebec, CANADA, September, 1996

“Underlying Ethical Principles in Research Using Repository Material,” Science to Practice: The Role of Research in Public Health, Emory University, Atlanta, GA, April, 1997

“Issues Related to the Re-Analysis of Stored Samples,” Biomarkers: The Genome and the Individual: Workplace and Medical Implications of a Rapidly Evolving Technology, Medical University of South Carolina, Environmental Hazards Assessment Program, Charleston, SC, May, 1997

“Protection of Privacy vs. Promotion of Research,” Privacy and Confidentiality: Is It a Privilege of the Past? Massachusetts Medical Society, 1997 Annual Education Program, Boston, MA, May, 1997

“Human Research: A Step Forward for Newly Included Populations?” 18th Annual Health Law Teachers Conference, Seton Hall Law School, Seton Hall, NJ, June, 1997

“Whose Tissue is It? The Patient as Product” Military Research Symposium: Protection of Human Subjects, Uniformed Services University of the Health Sciences, Bethesda, MD, June, 1997

“Using Stored Tissues for Genetics Research,” Interagency Human Subjects Conference, Genetics Research and Human Subjects: The Changing Landscape, Department of Energy, Bethesda, MD, June, 1997

“Impact of Regulation on Medical Research,” Regulation of Health Care: Latest Developments, ABA Society of Administrative Law and Regulatory Practice, ABA Midyear Meeting, Nashville, TN, January, 1998

“Research Ethics,” Health Law Teachers Meeting, Houston, TX, June, 1998
“Who Needs Informed Consent in Human Subjects Research,” Tennessee Technological University, Cookeville, TN, October, 1998

“DNA Banks and Human Research on Cancer Genetics,” Ethical Boundaries in Cancer Genetics, St. Jude’s Children’s Research Hospital, Memphis, TN, May, 1999

“Ethical Issues in Research in Pediatrics,” Pediatric Subcommittee, Anti-Infective Drug Committee, Food and Drug Administration, Chevy Chase, MD, November, 1999

“Genetics Research,” Ethical and Regulatory Aspects of Human Subjects Research Protocol, National Institutes of Health, Bethesda, MD, December, 1999

"Genetics," Privacy and Confidentiality in Clinical and Social Science Research: Myth or Reality?, University of Texas Health Science Center at Houston, February, 2000

"Children as Research Subjects," Research Ethics: Institutional Review Board Responsibilities and Related Competencies, Georgetown University Center for Clinical Bioethics, March, 2000

"Ethical Issues in Human Subjects Research: Staying Oriented in a Shifting Landscape," Ethical Issues in Health Care: The Nancy Tatum O'Neal Series, University of Mississippi Medical Center, Jackson, MS, April, 2000

"Legal Aspects of Studying Pregnant Women," Clinical Pharmacology During Pregnancy: Addressing Clinical Needs Through Science, FDA/NICHD Conference, Washington, DC, December, 2000

"Creating a Process to Collect Human Biological Materials and Medical Records for Research from Patients in Teaching Hospitals," A Decade of ELSI Research, Bethesda, MD, January, 2001

"Genetics Research in the International Context – The CIOMS Experience," A Decade of ELSI Research, Bethesda, MD, January, 2001

"The Academic Medical Center Perspective," Conflicts of Interest in Clinical Research: Legal and Ethical Issues, Widener University School of Law, Wilmington, DE, March, 2001

"So What Should We Do About Genetics Research?", Bringing Ethics to Life in Human Subjects Research: A Case Approach, Meharry Medical College, Program in Clinical and Research Ethics, Nashville, TN, September, 2001

"Fusing/Confusing Ethics and Regulation," "President's Advisory Committee on Stem Cell Research," "Third Party Human Subjects, Including Genetic Privacy," IRBs and New Approaches: Assessing, Evaluating, Discarding, and Moving Forward, PRIM&R's 2001 IRB Conference, Boston, MA, December, 2001

"Protecting the Privacy of Research Specimens," Inaugural Symposium, Genetic Privacy and Discrimination, University of Rochester, Rochester, NY, June, 2002

"Ethics, Politics, and the Haplotype Map," Genetics Symposium 2002, Nashville, TN, September, 2002

"Race and Genetics," "Race and Law," Research Ethics, Meharry CRECD Program, Nashville, TN, October, 2002

"Ethical Issues in Research Design," Research Ethics, Meharry CRECD Program, Nashville, TN, December, 2002

"Ethical Issues in Genetic Epidemiology – International Perspectives," US-Japan Symposium: Advances in Genetic Research in Cardiovascular Disease, Osaka, JAPAN, and Ethical Concerns in Molecular Epidemiology, Tokyo, JAPAN, February, 2003

"Collecting Human Biological Materials – The CPCRA Protocol," Terry Beirn Community Programs for Clinical Research on AIDS, Reston, VA, May, 2003

"Genetic Research in the Global Community," Globalization and Human Rights: Bioethics, Mexico City, MEXICO, June, 2003

"Ethics and Policy," 44th Annual Short Course in Medical and Experimental Mammalian Genetics, Jackson Laboratories, Bar Harbor, ME, July, 2003

"Legal Issues of HAPMAP: Individual and Community Rights," Working at the Frontiers of Law and Science: Applications of the Human Genome, The American Bar Association, American Medical Association and the American Association for the Advancement of Science, Chapel Hill, NC, October, 2003

"Ethical, Legal, and Social Issues in the International Haplotype Map Project," Colliding Categories: Haplotypes, Race, and Ethnicity, Minneapolis, MD, October, 2003

"Ethical Issues in Genomic Research," International Ethical Guidelines for Research Involving Human Beings, SANTIAGO, CHILE, October, 2003

"Legal and Policy Issues in the HapMap," The Genomics Revolution: Science, Law, and Policy, Louisiana State University, Baton Rouge, LA, February, 2004

"'Privacy' in Research," American Society of Experimental NeuroTherapeutics, Bethesda, MD, March, 2004

"The Web of Relations: Thinking About Physicians and Patients," A World Less Silent: Celebrating Jay Katz's Contributions to Law, Medicine, and Ethics, Yale Law School, New Haven, CT, October, 2004

"Ethical Issues in International Genetics Research," Distinguished Lecture Series, University of Texas at Houston Health Science Center – IRB, February, 2005

"Investigaciones genéticas con los niños," I Simposio Internacional de Bioética, Hospital Nacional de Niños, Dr. Carlos Sáenz Herrera, San Jose, Costa Rica, May, 2005

"Sheathing the Sword of Conscience," Legal Issues Affecting Academic Medical Centers and Other Teaching Institutions, American Health Lawyers Association, Washington, DC, January, 2006

"The Future of Ethics in Genetic Research: Informed Consent, Oversight, and Engagement, Tufts New England Medical Center, Boston, MA, January, 2006
"The HapMap Experience: Lessons for the Future," Wellcome Trust Advanced Course, Sanger Centre, Hinxton, UNITED KINGDOM, August, 2006

"Incidental Findings in Genetics Research Using Archived DNA," Managing Incidental Findings in Human Subjects Research, Minneapolis, MN, May, 2007

"Data Sharing in the Genomic Era," Egenis, Exeter, UNITED KINGDOM, May, 2007

"Ethical Challenges in Exploring Genomic Diversity," HGM2007, Montreal, CANADA, May, 2007

"Data Sharing and Its Implications," Second Circuit Judicial Conference, Bolton Landing, NY, June, 2007

"Genome-Wide Association Studies: What are the Issues?," Translating ELSI, Cleveland, OH, May, 2008

"Biobanking in the 21st Century: Lessons Learned About Navigating in Turbulent Waters from the HapMap and Other Projects, Instituto Superiore de Sanità, Rome, ITALY, June, 2008

"Returning Individual Results," SACHRP, Arlington, VA, July, 2008

"BioVU: Vanderbilt's Approach to Biobanking," Biobanking, Bioethics, and the Law, Stanford Law School, Stanford, CA, September, 2008

"Data Access v. Confidentiality: Balancing Risks and Benefits," American Society of Bioethics and Humanities, Cleveland, OH, October, 2008

"Data Access versus Confidentiality," Committee on Population and Committee on Collecting, Storing, Protecting and Accessing Biological Data Collected in Social Surveys, National Academies, Washington, DC, November, 2008

"Looking at the Ends of the Spectrum: What That Can Tell Us about the Big Question," Genetics Results Reporting Workshop, National Heart, Lung, and Blood Institute, Bethesda, MD, January, 2009

"What's Next in Genomic Research," What's Next in Law, Health and the Life Sciences? Debating Openness, Access, and Accountability, University of Minnesota, March, 2009

"Research on Residual Bloodspots: When Should We Report Back Results?," American College of Medical Genetics annual meeting, Tampa, FL, March 2009

"Informed Consent and What Else in Genomics?" Northwestern University, Chicago, IL, September, 2009

"What Data Do We Owe Back to Participants in Genetics & Genomics Research? Everything? Nothing?" American Society of Bioethics and Humanities, Washington, DC, October, 2009

"Biobanking at Vanderbilt University: Lessons Learned and Future Opportunities and Challenges," Medical Genome Science in the Personal Genome Era, ISAG2009, Tokai University, TOKYO, JAPAN, and Kyoto University, KYOTO, JAPAN, December, 2009

"What is the policy landscape for addressing IFs and IRRs in research using biobanks & archives?," University of Minnesota, January, 2010

"Biobanking of Residual Samples: Opt-Out?" Public Population Project in Genomics, MONTREAL, CANADA, April, 2010

"What Data Do We Owe Back to Participants?" OHRP Research Community Forum, Regulatory Responsibility and Innovative Research: An Opportunity for Partnership, Chicago, IL, May, 2010

"Legal Issues Related to the Usage of Newborn Screening Samples," Challenges and Opportunities in Using Newborn Screening Samples for Translational Research: A Workshop, Roundtable on Translating Genomic-Based Research for Health, Institute of Medicine, Washington, DC, May, 2010

"Legal Issues Related to the Use of Newborn Blood Spots for Research," Health Law Professors, Austin, TX, June, 2010

"What Role Should Research Participants' Preferences About Data Sharing Play?," International Data Sharing Conference, Oxford, UNITED KINGDOM, September, 2010

"What Data Do We Owe Back to Participants in Genetics & Genomics Research? Everything? Nothing?," Human Subjects Protection: A Time to Remember, Covington, KY, October, 2010

"Retos éticos para las aplicaciones clínicas de la medicina genómica" 10th Anniversary, INMEGEN, Mexico City, MEXICO, October, 2010

"National policy statements on the retention and use of residual samples for research," Methods for Promoting Public Dialogue on the Use of Residual Newborn Screening Samples for Research, Salt Lake City, UT, January, 2011

"Challenges in Genetic Research" and "Panel on Genetic Research," 6th Annual Columbia

University IRB Conference, New York NY, March, 2011

“Hot Topics in Genomics Research,” 6th annual Columbia University IRB conference, New York, NY, March, 2011

“Workshop on Return of Results,” Exploring the ELSI Universe, University of North Carolina, Chapel Hill, NC, April, 2011

“Reservations about Return of Results,” ISBER Conference, Arlington, VA, May, 2011

“The Misguided Debate about Return of Research Results,” Should We Return Individual Research Results and Incidental Findings from Genomic Biobanks and Archives? University of Minnesota, Consortium on Law and Values in Health, Environment, and the Life Sciences, Bethesda, MD, May, 2011

“Panel on Joint Consideration of Special Complexities in Genotype-Driven Recruitment,” Ethical Approaches to Genotype-Driven Research Recruitment,” Durham, NC, May, 2011

“Using Newborn Blood Spots for Research and the Common Rule,” American College of Medical Genetics, Salt Lake City, UT, September, 2011

“Biospecimen Exceptionalism,” The Future of Human Research Regulation, Harvard Law School, Boston, MA, May, 2012

“Should Results of Genomics Research Be Returned to Study Participants?” Personalized Medicine for the Care of the Child with Heart Disease: Discovery to Application, Toronto, CANADA, October 2012

“How Should We Use Genomic Information for Primary Prevention?” 2013 Research Education Symposium: Life at the Interface of Genomics & Clinical Care, University of Michigan, Ann Arbor, MI, March, 2013

“The Role of Bioethicists in Policy Formation,” Greenwall Fellows Meeting, Chicago, IL, May, 2013

“Ethical and Community Perspectives: The need for community consultation in biospecimen collection”, 2013 Biospecimen Science and Cancer Health Disparities Conference, Johns Hopkins Center to Reduce Cancer Disparities and Meharry Medical College, Nashville, TN, July, 2013

“Lessons Learned from Biobanking,” Treuman Katz Lecture, Seattle Children’s Hospital, Seattle, WA, October, 2013

“A Review of the SUPPORT Study and Ethical Challenges in Assessing and Disclosing Research Risk,” CTSA Clinical Research Ethics Key Function Working Group Workshop on

Medical Practice Research, Atlanta, GA, October, 2013

“Ethical Issues in Anticipate and Communicate,” eMERGE Consortium Steering Committee Meeting, Bethesda, MD, January, 2014

“Ethical Perspectives on Big Data,” Ethical Issues in Data Science and Digital Medicine, Icahn School of Medicine, New York, NY, February, 2014

“Ethical Challenges in Genome-Based Cancer Research: Liability and Other Challenges,” IOM Workshop on Contemporary Issues in Human Subjects Protections in Cancer Research, Washington, DC, February, 2014

“Return of Results – From the Clinic to Research,” Sophia University and 164th Committee on Genome Technology, TOKYO, JAPAN, March, 2014

“Ethical Issues in Anticipate and Communicate,” CSER Consortium meeting, Bolger Center, Potomac, MD, May, 2014

“Ethical Issues in Human Genetics Research,” Statistical Genetics and Genomics Short Course, University of Alabama at Birmingham, Birmingham, AL, July, 2014

“How Much Individual Choice?” Institute for Public Health 7th Annual Conference: Public Good and Individual Choice, Washington University, St. Louis, MO, October, 2014

“Finding Your Way on Shifting Sands: Reflecting on the Changing Ethical, Regulatory, and Policy Landscape of Genomics Research,” Novartis, Boston, MA, November, 2014

“How much control do adolescents have over parental access to and sharing of their genomic test results? How much should they?” Should We Offer Genomic Research Results to a Participant's Family, Including After Death? University of Minnesota, Minneapolis, MN, November, 2014

“Balancing autonomy with social benefit: From broad or tiered consent to notification?” International Biobanking Summit, Phoenix, AZ, May, 2015

“When Participants in Genomic Research Grow Up: Contact and Consent at the Age of Majority,” 3rd P3G International Paediatric Platform Conference, Montreal, CANADA, May, 2015

Pediatrics:

"Compensation for Vaccine-Related Injuries: Developments on the National and State Level," Annual Meeting, American Society of Hospital Pharmacists, Nashville, TN, June, 1989

"Infant Care Review Committees," Huntsville Hospital, Huntsville, AL, January, 1990

"Drug Testing in Teenagers," PRIDE International Convention, Orlando, FL, April, 1990

"Decision-Making for Children: Key Legal Cases, Statutes, Regulations, and Policy Issues in Pediatric Cases," Ethics Committees and the Young: Families, Hospitals and the Courts Trying to Do the Right Thing, St. Louis University Schools of Law, Medicine, and Nursing in conjunction with American Society of Law, Medicine and Ethics, St. Louis, MO, May, 1994

"Legal vs. Religious Exemptions: A View From the American Academy of Pediatrics," National Council of Juvenile and Family Court Judges, Nashville, TN, July, 1994

"Health Care Decision-Making and the Law," Medical Ethics for Physicians and Nurses: End of Life Issues, University of Alabama in Huntsville School of Primary Medical Care, Huntsville, AL, July, 1994

"Protection and Justice in the Care of Maltreated Children," "Legal Ethics in Child Protection," Cross Training: Knowing Your Way Around the Court, Case Western Reserve Child Protection Resource Center, Cleveland, OH, April, 1995

"Pediatric Ethics -- A View from the United States," Japanese Pediatric Society Annual Meeting, Kumamoto, JAPAN, April, 1996

"Inclusion of Children in Clinical Research," National Institute of Child Health and Human Development, American Academy of Pediatrics, Bethesda, June, 1996

"Legal and Ethical Issues Associated with Newborn Hearing Screening," Fourth International Symposium on Childhood Deafness, Kiawah Island, SC, October, 1996

"Families and Health," Family Re-Union 7, Nashville, TN, June, 1998
"Who Speaks for the Child: Ethical Issues in Pediatric Autonomy," Section on Bioethics, Annual Meeting, American Academy of Pediatrics, San Francisco, CA, October, 2001

"Medical Ethics and Developmental Disabilities," The Young Child with Special Needs, 13th Annual Conference, Nashville, TN, March, 2004

"Legal and Ethical Issues in Child Abuse Reporting," Annual Review of Pediatrics, University of Wisconsin, Madison, WI, September, 2007

"How Lawyers Can Help Patients (and Their Pediatricians), Arkansas Children's Hospital, Little Rock, AK, December, 2008

"Legal and Ethical Issues in Reporting Child Abuse and Neglect," Arkansas Children's Hospital, Little Rock, AK, December, 2008

“Pediatric Ethics: What Has Law Got to Do with It?” William G. Bartholome Award for Ethical Excellence Lecture, Section on Bioethics, AAP Annual Meeting, San Francisco, CA, October, 2010

“Vaccines: From Science to Policy,” Western Conference of the Bench and Bar, United States Court of Federal Claims, Berkeley, CA, October, 2011

“Do Adolescents Have Rights to Make Decisions About Their Health Care,” Buhl Lecture, University School of Nashville, TN, April, 2012

“Confronting Commercial Sexual Exploitation and Sex Trafficking of Minors in the United States,” Eighth Annual Cynthia Jean Stolman Lecture in Medical Ethics, Rutgers New Jersey Medical School, Newark, NJ, November, 2013

“Lessons Learned on my Path to Child Advocacy,” Frank H. Morriss, Jr. Leadership Award Lecture, University of Iowa, Iowa City, IA, February, 2015

Women's Health:

"Women and Advances in Medical Technology: The Legal Issues," The Effects of Pharmacology, Technology, and Other Modern Interventions on Women's Health, MacArthur Foundation Network on Health Behavior, Key Biscayne, FL, June, 1987

"The Law and Advances in Technology: Who Decides How Women and Their Unborn Children Should Live?," Reproductive Technology and Ethics for the Year 2000, Spokane, WA, April, 1988

"The Legal Status of the Fetus: Reflections on the Duties of Pregnant Women and of Society," Birth Defects Symposium XIX, Albany, NY, September, 1988

"Maternal-Fetal Conflicts," Hillsborough and Pinellas County Associations of Women Lawyers and for Women Physicians, Tampa, FL, February, 1989

Grand Rounds, Department of Obstetrics & Gynecology, Staten Island Hospital, Staten Island, NY, June, 1989

"Reproductive Technology and Human Values: A Health Care Professional and Woman's Perspective," Reproductive Technology: Ethical Issues, Ethical Alternatives, Maryville College, Maryville, TN, October, 1990

"Legal Issues and Reproductive Care," Legal, Medical and Ethical Issues in Women's Reproductive Health and Neonatology, American Society of Law, Medicine and Ethics, Naples, FL, January, 1995

“Legal Aspects of Studying Pregnant Women,” FDA.NICHD Conference, Clinical Pharmacology during Pregnancy: Addressing Clinical Needs Through Science, Washington, DC, December, 2000

“Legal and Ethical Issues in Embryo Mix-Ups,” AALS/ASRM Workshop on Reproductive Medicine and Law, Vancouver, CANADA, June, 2007 and ASRM Annual Meeting, Washington, DC, October, 2007

“Predictive Genetic Testing of Fetuses and Newborns: Mapping the Ethical and Legal Landscape,” Obstetrics and Gynecology Grand Rounds, Pittsburgh, PA, October, 2013

“A Call to Action: Child exploitation and sex trafficking in the US,” AAP Annual Meeting, San Diego, CA, October, 2014

Medical Malpractice and Health Care Reform:

"Conversations with Patients: How Patients See Their Physicians and Why They Say They Sue," Patient Grievances in Health Care: A Research Seminar, Ohio State University Center for Sociolegal Studies and Office for Health Services Research, Columbus, Ohio, November, 1992

"Health Care Reform: Ethics and the Practicing Pediatrician," American Academy of Pediatrics Annual Meeting, Dallas, TX, October, 1994

Visiting Professor Lecture Series, Department of Pediatrics and Center for Medical Ethics, Pediatric Grand Rounds -- "Why People Sue Their Doctors," University of Pittsburgh Medical Center, Pittsburgh, PA, December, 1994

"The Manifestations of Women's Dissatisfaction: Why Women Sue Their Physicians and How the Health Care System Has Responded," Legal, Medical and Ethical Issues in Women's Reproductive Health and Neonatology, American Society of Law, Medicine and Ethics, Naples, FL, January, 1995

"Medicaid, Managed Care, and the New Health Services Markets," Managed Care, Integrated Delivery Systems, and Consolidation -- Law, Medicine, and Ethics, American Society of Law, Medicine, and Ethics, Boston, MA, September, 1995

"Restructuring Healthcare: Jeopardizing Patient Care," Annual Meeting, Oncology Nursing Society, Nashville, TN, November, 1995

“Lawsuits--Why Do They Happen and What Can We Do About It?” Annual Conference, Pediatric Nursing, Integrating Mind, Body, and Spirit to Enhance Pediatrics Outcomes: New Challenges, Nashville, TN, September, 2000

Other:

"Teaching Across Professional Boundaries," 17th Annual Health Law Teachers Conference, Wilmington, DL, June, 1996

"The Confidential Medical Record in the New Millennium - Who Has Access and Where Does It Go?" 26th Annual Course, Society for Gastroenterology Nurses and Associates, Nashville, TN, May, 1999

"The Ethics Quagmire – Lifelines for Health Care Practitioners and Lawyers," Tennessee Bar Association, Brentwood, TN, October, 2008

Conferences Organized:

Law, Literature, and Social Change, Vanderbilt Law School and Robert Penn Warren Center for the Humanities, Nashville, TN, March, 1990 (with Jay Clayton)

Informed Consent for Genetic Studies on Stored Tissue Samples, Ethical, Legal, and Social Implications Branch/NCHGR/NIH and U.S. Centers for Disease Control and Prevention, Bethesda, MD, July, 1994 (with Elizabeth Thomson)

Legal, Medical and Ethical Issues in Women's Reproductive Health and Neonatology, American Society of Law, Medicine and Ethics, Naples, FL, January, 1995 (with Cindy Hylton Rushton)

The Changing Health Care Environment: Issues for the Future, American Society of Law, Medicine and Ethics and Vanderbilt University School of Medicine, Nashville, TN, March, 1997 (with H. Lee Barfield)

Children, Parents, Professionals, and the State: Ethical, Clinical, and Legal Issues in Health Care for Children, American Society of Law, Medicine and Ethics, Cambridge, MA, September, 1997 (with John Lantos)

A Public Dialogue about Gene Patenting, Vanderbilt University Center for Genetics and Health Policy and the American College of Medical Genetics, Washington, DC, March, 2000 (with R. Rodney Howell)

First Annual Meharry/Vanderbilt Genetics Symposium, Nashville, TN, November, 2000 (with Al George, Jonathan Haines, John Phillips, Shirley Russell, Scott Williams)

Second Annual Meharry/Vanderbilt Genetics Symposium, Nashville, TN, September, 2001 (with Al George, Jonathan Haines, John Phillips, Shirley Russell, Scott Williams)

Third Annual Meharry/Vanderbilt Genetics Symposium, Nashville, TN, September, 2002

(with Jonathan Haines, John Phillips, Shirley Russell, Scott Williams)