

Curriculum Vitae

DEREK J JONES

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PERSONAL INFORMATION

Place of Birth: Swindon, England
Citizenship: U.S & Canada
Languages: Fluent in English & French

EDUCATION

1982-1985 **HARVARD UNIVERSITY LAW SCHOOL** CAMBRIDGE, MA
Received Juris Doctor, December 1985.

1981-1982 **INSTITUT D'ÉTUDES POLITIQUES** PARIS, FRANCE
As French Government Research Fellow, awarded under the Fulbright
Hayes International Scholarship Program, studied development of French
& European Union cosmetic and chemical safety legislation and policy.

1976-1980 **YALE UNIVERSITY** NEW HAVEN, CT
Received Bachelor of Arts in Political Science & Economics, June 1980.
Concentrated on public policy decisions involving health, safety & scientific
regulation. Played J.V. Hockey & wrote for the *Yale Daily News*.

WORK EXPERIENCE

2008-22 **SENIOR CONSULTANT IN LAW & PUBLIC POLICY**
1996-2001

Senior Consultant, advisor and scholar for government, universities, health care institutions and professionals in the analysis of pressing public policy and law issues. Expertise includes health law; risk-benefit and ethical-legal analyses of new technologies, health law matters, biotechnology; comparative & international law; public policy and human rights issues, often through interdisciplinary research teams. Sample clients have included Health Canada, Justice Canada, University of Toronto, Industry Canada, the Tri Council of Canada (Medical Research Council, Social Science & Humanities Research Council, Natural Science & Engineering Research Council); Government of Canada Interdepartmental Committees on Ethics & Biotechnology; Canadian Institutes for Health Research; Bayer International Advisory Council on Bioethics, UNESCO, European Parliament. Collaborations undertaken on a project, part-time or term basis.

1991-present

**LECTURER, ADJUNCT PROFESSOR, RESEARCH SCHOLAR
McGILL UNIVERSITY, MONTRÉAL**

TEACHING RESPONSIBILITIES: *FACULTY OF LAW: Psychiatry & the Law*: (1998-02; biennially 2012-22); *Law & Health Care*: (1993-2004: founder of this graduate level law, Bioethics, and Health care course for philosophy, theology, nursing, medical, health administration and other non-law students in McGill's Graduate Bioethics curriculum); *AIDS Law*: (1997-2002: one of the first AIDS courses in Canada, it was recognized by the Association of American Colleges and Universities for inclusion in National Leadership Resource Database as part of Learning for Our Common Health Program); *Comparative Medical Law*: (1991-1997, co-taught with Margaret Somerville). Courses emphasize (i) international comparative scholarship, (ii) interdisciplinary analysis for problem-solving, and (iii) role(s) of law in regulating science, technology, patient rights, professional relations, individual-societal risk-benefit analyses. *FACULTY OF NURSING*: Lectures in *Health Law & Ethics* (1998-99); Adjunct Professor to establish, design and teach *Nursing Law & Ethics* (2000-06). *FACULTIES OF MEDICINE, LAW & SOCIAL WORK* (2008-20). Guest lectures in selected coursework: medical confidentiality, neuro-ethics, comparative health law, social work and law, society, and medicine.

RESEARCH & SCHOLARSHIP: As former Senior Research Fellow for Centre for Comparative Law, former associate member of McGill's Centre for Medicine Ethics & Law, member of McGill's teaching hospitals clinical and research ethics committees, a founding and current member of the Research Group on Health Law, member of Faculty Editorial Advisory Committee of the *McGill Journal of Law and Health*, and member of the McGill Centre for Human Rights & Legal Pluralism, focus interdisciplinary research projects on a range of classic to emergent health law, ethics and human rights issues: e.g., mental health, disability and human rights; comparative privacy and confidentiality norms; ethics frameworks for conflicts of interest; human research, scientific conduct and biotechnology; global health law and bioethics; citizen engagement in ethics of life sciences; evolving ethico-legal standards for health professionals, therapeutic products, organ transplantation, medically assisted procreation, end-of-life care, etc.

2002-2007

**EXECUTIVE DIRECTOR, GOVERNMENT OF CANADA INTERAGENCY
ADVISORY PANEL & SECRETARIAT ON RESEARCH ETHICS OTTAWA**

As founding Executive Director of novel national advisory committee and supporting Secretariat, advised government on the revision, use, interpretation and governance of Canada's national policy for the ethical conduct of research involving humans. Results yielded second edition of Canada's national research ethics standards. Substantive issues included ethico-legal and international harmonization; emerging questions at the interface of scientific research, technology, and ethics; removing anachronisms and reforming consent/privacy/confidentiality standards,

working definitions, principles and exceptions; building national consortium for ethical guidelines with peoples from Aboriginal communities; improving ethical norms for research in the social sciences and humanities. Issues processed through a policy development cycle — from identification to research and analysis, policy options, textual proposals, and amendments – for deliberative democratic definition of a new generation of human research ethics norms. Oversaw, researched, and wrote advisory opinions on interpretation of national ethics questions. Special responsibilities for international standards and liaison (e.g., Australia, US, France, UK, EU, UN), national ethics governance, intergovernmental representation and strategic partnerships, design of public engagement framework, and operational guidance of Secretariat team to support a dozen national interdisciplinary working sub-committees. *Ex officio* member of Panel, and substantive lead on conflicts of interest, clinical trials information, stem cell research, legal and regulatory issues, and principles for ethical research in public emergencies.

1994-1996

**DIRECTOR, NATIONAL COUNCIL ON BIOETHICS IN HUMAN RESEARCH
OTTAWA**

As Director of this national advisory council on the ethics of human research, contributed to analysis of issues at the confluence of scientific policy, ethics, and law, as illustrated by the novel questions presented by embryo research, experimentation on incompetent persons, new drug development and access thereto and justice in the allocation of the benefits and burdens of research. Operationally responsible for administration of Council secretariat, assisting in the development, coordination and execution of work agendas and projects of substantive NCBHR working committees; drafting budget; organizing meetings and national ethics workshops; editing publications and reports. Special responsibilities for outlining NCBHR research ethics education initiatives; serving as draft-person for written responses to formal research ethics queries; consolidating guidelines, codes and laws on research ethics norms; and editor-in-chief of the research ethics periodical, *NCBHR Communiqué CNBRH*.

1992-1994

**VISITING FELLOW, CENTRE DE RECHERCHE EN DROIT PUBLIQUE,
UNIVERSITÉ DE MONTRÉAL
MONTRÉAL**

As a Visiting Fellow in the Centre for Public Law Research of the Faculty of Law, University of Montreal, served as consultant to the Department of Justice Canada on legal and ethical issues presented by medical testing and biomedical monitoring in the federal workplace (e.g., mandatory substance abuse testing, genetic testing, pre-employment medical examinations). Focussed on reconciling public safety and human rights arguments, evolving medico-legal standards, and comparative international law insights. Convened and coordinated multidisciplinary team drawn from law, philosophy, and medicine in initial research on this project while at the Law Reform Commission of Canada.

- 1988-1992 **SENIOR LEGAL ADVISOR, LAW REFORM COMMISSION OF CANADA**
 MONTRÉAL
- As senior legal adviser in health law and bioethics section of the Commission, examined legal facets of national biomedical, health & ethics issues with view towards recommending to Parliament advisable changes in federal law. Special responsibilities for monitoring and advising on major U.S. health law and bioethical developments and trends. Supervised or participated in analysis of medical device & drug law to ensure safe human therapeutics, assisted procreation law, patenting human life forms, proprietary rights in human cellular materials, genetic engineering, occupational health law, medical waste, human research, fetal tissue transplants, evolving legal regimes for biotechnology, organ procurement & tissue banking law, euthanasia, international bioethics -- many of which challenge societal norms, conventional thought & fundamental values.
- 1986-1988 **ATTORNEY, PRIVATE PRACTICE** PORTLAND, MAINE
- As an associate with Pierce, Atwood, Scribner et al, assisted in litigating wrongful discharge, employment discrimination, hazardous waste siting, personal injury & medical liability cases. Subsequent solo practice concentrating in health law involved advising on AIDS screening legislation, hospice liability issues, US-Canadian occupational health & worker compensation systems.
- Autumn 1987 **DALHOUSIE UNIVERSITY LAW SCHOOL** HALIFAX, NOVA SCOTIA
- Visiting Scholar in law & medicine. Discharging special advisory responsibilities to Deans of Law & Medicine, helped develop and establish the Dalhousie University Health Law Institute.
- 1985-1986 **RESEARCH ASSOCIATE, BOSTON UNIVERSITY** BOSTON, MA
- At Health Law Center of School of Medicine & Public Health conducted research and analysis of comparative organ transplantation law & policy, functions & limits of informed consent, human experimentation law, termination of life support, artificial reproduction law.
- Summer 1985 **SUMMER ASSOCIATE, PIERCE, ATWOOD, SCRIBNER** PORTLAND, ME
- Summer Associate in 60-attorney law firm. In representation of plaintiff & defense actions, researched & wrote legal motions & memoranda on legislative history of Maine Dram Shop Act, bankruptcy, labour arbitration, health and constitutional law.
- Summer 1984 **RESEARCH ASSOCIATE, UNIVERSITY OF CALIFORNIA**
 SAN FRANCISCO, CA
- As special consultant to collaborative research effort between Institute for Health Policy Studies & the Public Hospitals of Paris, assessed state-of-

the-art, legal status & economics of some of the newest medical technologies emerging in Europe & the US: lasers, transplantation, magnetic imaging, *in vitro* fertilization, genetic prenatal diagnostic biotechnologies.

- Summer-
Autumn 1983 **PUBLIC ADVOCATES** SAN FRANCISCO, CA
During leave of absence from law school, served as principal draftsman on international petition to the World Bank. Drawing on US & European export law & relevant UN, European, OECD initiatives, petition proposed legal remedies to ensure proper information exchange on -- and informed, judicious use of -- hazardous agrochemical substances/pesticides in sustainable development and international trade. The issue impacts farmer safety, public health, regional ecology, agricultural development, international pesticide residues on food, and global food safety. In 1985, the World Bank established innovative pesticide standards and pest management controls in Bank-financed projects. The initiative became a model for environmental standards for public international development banks.
- January
August 1981 **RESEARCH ASSOCIATE, UNIVERSITY OF PITTSBURGH** PITTSBURGH, PA
Headed research project that evaluated preventive health status of Pennsylvania relative to 100 national prevention objectives;
Literary editor of *Vital Signs*, monthly publication of the American Nurses' Association, Pittsburgh chapter.
- Autumn 1980 **O'REILLY FOR CONGRESS** ANN ARBOR, MI
As aide for congressional candidate, formulated & implemented media strategies, candidate background briefings, & issued papers on subjects ranging from national nurse shortage to regional mass transit needs.
- Summer 1980 **UNITED MINeworkERS' HEALTH & PENSION FUND** WASHINGTON, D.C.
Assisted policy analysis division in investigating duplicate payments problems in 2 major fraud & abuse cases, & in identifying & analyzing duplicate payments made to health care providers who had rendered services to miners.
- Summer 1979 **CONSUMER FEDERATION OF AMERICA** WASHINGTON, D.C.
Under research grant from Yale University, assisted NGO staff in analysis & articulation of consumer interests in legislative & regulatory food & drug safety & health policy.
- Autumn 1978 **HEALTH SYSTEMS AGENCY OF S.W. PENNSYLVANIA** PITTSBURGH, PA
Provided research assistance in finance, long-term care & regional, state, national health care policy. Involved analyzing impact of additional health

care resources on existing regional health care systems. Fall leave of absence from Yale.

Summer 1978 **CONSUMER PROTECTION BUREAU** PITTSBURGH, PA
 Mediated consumer complaints against allegedly unscrupulous businessmen. Exposed to difficulties of enforcing laws & acts governing fair trade, debt collection, credit & contracts.

SELECTED MEMBERSHIPS

- Admitted to practice before Courts of Maine & Massachusetts; Member of American Bar Association (1988-).
- American Society of Law, Medicine & Ethics (1987-).
- Canadian Bioethics Society (1988-2007).
- Science & Technology, Health Law, International Law, and Human Rights Sections of the American Bar Association (1988-).
- National Health Lawyer's Association (US) (1988-2002).
- Maine & Massachusetts Bar Associations Health Law Sections (1988-2002).
- Advisory Council of the Canadian Bioethics Society (1997-99).

PROFESSIONAL & COMMUNITY ENGAGEMENT

- Associated Senior Research Fellow, Paris School of Economics, Université de Paris 1/Hospinnomics Research Centre, (Nov 2016-)
- McGill University Faculty of Law, Centre for Human Rights & Pluralism (2011-).
- McGill University Faculty of Law, Research Group on Health & Law (2005-).
- *McGill Journal of Law and Health*, Faculty Editorial Advisory Committee (2005-).
- Canadian National Council on Bioethics in Human Research, Task Force on Accreditation of Participant Protection Program, (observer 2004-06).
- Government of Canada Interdepartmental Committee on UNESCO *Draft Declaration on Bioethics and Human Rights* (2004-2005).
- Government of Canada Interdepartmental Committee on UNESCO *International Declaration on Human Genetic Data* (2003).
- Government of Canada Interagency Advisory Panel on Research Ethics, *ex officio* member (2002-2007).
- Government of Canada Interdepartmental Committee on Human Research Ethics (2002-2007).
- McGill University Health Centre Bioethics Subcommittee on Resuscitation Policy (2000-02).

- Clinical Ethics Committee, Montreal Children's Hospital (1997-2002).
- Research Ethics Committee, Sir Mortimer Davis Jewish General Hospital (Montreal) (1997-2000).
- Reproduction & Genetics Technology Task Force of the American Bar Association Section on Family Law (1993-98).
- Associate Member, McGill University Centre for Medicine, Ethics & Law (1991-97).
- Clinical Ethics Committee, Sir Mortimer Davis Jewish General Hospital (Montreal) (1992-96).
- Clinical Ethics Committee, Ottawa General Hospital (1995-97).
- Standing Committee on Ethics, Medical Research Council of Canada (Observer, 1994-96).
- Government of Canada Interdepartmental Committee on New Reproductive Technologies (1990-92).
- Government of Canada Interdepartmental Committee on Biotechnology, Subgroup on Safety & Regulations (1991-92).
- Board Member, Hospice of Maine (1986-88).

SELECTED PRESENTATIONS

- "Mental Health Restraints & Seclusion: Human Rights Insights," Mental Health Commission of Canada Forum: Ottawa, Canada, March 2018.
- "Enabling Justice: Mental Health & Human Rights at Work" Hospinomics Institut, Paris School of Economics Hôtel-Dieu: Paris, May 2017.
- "Mental Health in the Workplace: A Global Human Rights Challenge?" Centre for Disability & Law, University of Ireland: Galway Ireland, March 2017.
- "Global Health Law & Human Rights," University of Aix-Marseille Law School: Aix en Provence, France, April 2017.
- "Enabling Justice: Mental Health Information Privacy & Equality at Work," International Labour Organization Expert Meeting, Geneva: October 2016.
- "Mental Health Privacy and Discrimination on the Job," Annual Conference of the Canadian Association of Statutory Human Rights Agencies (CASHRA), Montreal: May 2016.
- "Mental Health & Human Rights in the Workplace," World Congress of Law & Mental Health, Vienna: July 2015.

- “Developments in Mental Health Human Rights Law,” Quebec Human Rights Tribunal Annual Retreat. Estérel, Quebec: March 2014.
- “Silencing Expertise: Conflicts of Interest in International Clinical Practices Guidelines,” to the French National Expert Committee on Economic Evaluation and Public Health of the Haute Autorité de la Santé, French Ministry of Health. Paris: June 2013.
- “Transformative Justice: The Roles of Law in Shattering Stigma in the Workplace,” 5th International Stigma Conference: Ottawa: June 2012
- “Managing Conflicts of Interest in Expert Advisory Committees' Practice Guidelines: Regulatory Decisions & International Insights,” American Society of Law Medicine & Ethics National Symposium: Conflicts of Interest in the Practice of Medicine. Pittsburgh: October 2011.
- “Conflict of Interest in the Development of Medical Practice Guidelines,” Canadian Gastroenterology Society Annual Meeting. Banff Alberta: March 2009.
- “Ethics Frameworks to Manage Conflicts of Interest in Evidence-Based Clinical Practice Guidelines.” International Consensus Conference of the Canadian, European & Asian Gastroenterology Medical Societies. Vienna: October 2008.
- Pluralistic Perspectives on Public Health Ethics”, session moderator for the First Canadian Roundtable on Public Health Ethics: Exploring the Foundations. Montreal: November 2007.
- “Public Emergencies Research: Ethical Norms and Principles.” Canadian Association of Research Ethics Board Annual Meeting. Montreal: May 2007.
- “Imagine: A National Research Ethics Education Strategy,” 5th Annual Workshop of the Government of Canada Interagency Advisory Panel on Research Ethics. (co-presenter Glen Greiner). Ottawa: February 2007.
- “Towards a National Research Ethics Education Strategy.” National Council on Ethics in Human Research, Education Subcommittee. Alymer, Quebec: December 2006.
- “Harmonization and National Research Ethics Identity: Canada’s Path”, presented at the International Town Hall Meeting of the Annual Meeting of Public Responsibility in Medicine (PRIM&R). Washington, DC: November 2006.
- “The Duty to Share New Information in Clinical Trials: Working Recommendations for Canada’s National Research Ethics Policy,” 16th World Congress on Medical Law. Toulouse, France: August 2006.
- “Public Process Values, Deliberative Democracy, and New Models of the Ethics of Research Involving Humans,” International Conference of the Association canadienne-française pour l’avancement des sciences (ACFAS). McGill University. Montréal: May 2006.

- “Drawing on Research Ethics Boards’ Culture as a Voice of Change for Research Ethics Norms” presented to the Annual Conference of the Canadian Research Ethics Boards, (co-presenter: Bruce Clayman), Toronto: May 2006.
- “Negotiating Space for Indigenous Knowledge: Dialogue on TCPS Research Ethics” presented to the Government of Canada Aboriginal Policy Research Conference. Ottawa: March 2006.
- “Informed Consent Issues in Canadian Research Ethics Norms: A Sampling,” International Conference on Informed Consent in a Changing Environment of Research Involving Humans. Oxford University, United Kingdom: June 2005.
- “Towards New TCPS Ethics Guidelines for Research involving Aboriginal Peoples: The Emerging Process” (co-presenters: Marlene Brant Castellano & Willie Ermine), Trent University. Ontario: June 2005.
- “The Evolving Interface between Research Ethics and Human Rights”, presented to the Government of Canada Biotechnology Secretariat Conference - A Brave New World: Where Biotechnology and Human Rights Intersect. Ottawa: April 2005.
- “The TCPS Work-in Progress: Partnered Responses to Privacy Developments”, presented to the Canadian Institutes of Health Privacy Workshop. Ottawa: May 2004.
- “Update on Activities of the Government of Canada Interagency Advisory Panel on Research Ethics”, Canadian Bioethics Society Annual Meeting (co-presenter) Calgary. October: 2004.
- “Longitudinal Research Ethics: A TCPS Perspective on Canadian Lifelong Health Initiatives”, Canadian Institutes for Health Research Planning Workshop, Legal and Ethical Issues facing the Canadian Lifelong Health Initiative. Montreal: November 2003.
- “From Recruitment to Participation in Research Ethics”, Presented to an International Workshop at the Joint Annual Meeting of the American Society for Bioethics and Humanities and the Canadian Bioethics Society. Montreal: October 2003.
- “Listening to the Community: A Case Study on Procedural Reform of the TCPS”, presented to the Annual Conference of the Learned Societies of the Humanities and Social Sciences of Canada. Halifax, Nova Scotia: June 2002.
- “Selected International Legal Norms on the Protection of Personal Information in Health Research,” presented to Annual Meeting of the Canadian Bioethics Society. Winnipeg: October 2001.
- “Human Rights, Personal Data & International Norms,” presented to the Genetic Data Working Group of the International Bioethics Committee of UNESCO. Paris: June 2001.
- “Repairing the Past: Ethico-Legal Standards for the Duplessis Orphans,” presented to the International Academy of Mental Health Law Congress. Montreal: June 2001.

- “Genetic Testing & Human Rights,” presented to the Health Canada Advisory Committee on Genetic Testing for Late-Onset Diseases. Ottawa: June 2000.
- “Role & Structural Models for Ethics within National Institutes of Health Research: A Comparative Perspective,” presented to the Canadian Institutes of Health Research Interim Governing Board Subcommittee on Ethics. Montreal: July 1999.
- “Proportionate Review of Ethics Assessment,” in Round Table on Implementing the Tri-Council Policy Statement on Ethical Conduct for Research Involving Humans: Ottawa, Canada: October 1998.
- “Managing Conflict in Bioethics: Alternative Dispute Resolution & Mediation,” Workshop developed for the University of Toronto Centre for Bioethics. Toronto: Spring 1998.
- “Legal & Ethical Duties of the Disabled Doctor.” Radio Interview, Canadian Broadcast Corporation, Montreal: Winter 1998.
- “Resolving Bioethics Disputes through ADR,” presented to the Canadian Bioethics Society Annual Meeting: Halifax, Nova Scotia: October 1997.
- “Positive & Negative Rights in Psychiatric Research Ethics,” presented to the 22nd International Congress on Law & Mental Health, Montreal: June 1997.
- “Genetic Confidentiality & the Family: Whose Rights Prevail?” presented to the Canadian Bioethics Society Annual Meeting (co-presenter, L. Arbour) Montreal, Canada: October 1996.
- “Canada’s Draft Code of Conduct for Research Involving Humans,” panellist on Round table at the University of Ottawa, Faculty of Medicine: June 1996.
- “Psychiatry & Research Ethics,” presented to a National Conference on Research Methods in Psychiatry at the Royal Ottawa Hospital Institute of Mental Health Research: Ottawa, Canada: February 1996.
- “Ethics & Compassionate Access to Experimental Therapies,” presented to the HIV/AIDS Subcommittee of the Standing Committee on Health of the House of Commons of Canada (co-presenter, Benjamin Freedman, televised) Ottawa, Canada: December 1995 & May 1996.
- “Judicialization of Genetic Medicine,” presented as part of a Professional Norms in the Practice of Human Genetics research project to the Canadian Bioethics Society Annual Meeting: (co-presenters, BM Knoppers, C. Laberge, D. Wertz et al) Ottawa, Canada: November 1994.
- “Ethico-Legal Regimes to Regulate the Trade in Human Bodily Parts & Genetic Materials,” Workshop chairperson for the 8th International Biotechnology Meeting: Toronto: May 1994.
- “Human Rights & Bioethical Issues in AIDS Testing,” presented to the Department of Justice Canada 5th Annual Conference on the Canadian Charter of Rights & Freedoms: Ottawa, Canada: November 1993.

- “Ethical & Legal Aspects of Genetic Diagnostics,” presented to the Annual Meeting of the German Society of Clinical Chemistry & German Society of Laboratory Medicine: Dresden, Germany: October 1993.
- “Does the Law Care Whether Doctors Lie to Patients?” presented to the Jewish General Hospital, Grand Medical Rounds. Montreal: April 1993.
- “Transplantation & the Law: Recent Proposals from the Law Reform Commission of Canada,” presented to the Third International Conference on Health Law & Ethics. Toronto, Canada: July 1992.
- “Evolving Medico-Legal Standards of Practice in the Maternal-Fetal “Rights” Discourse: The Case of Electronic Fetal Monitoring,” Presented to the Third International Conference on Health Law & Ethics, (co-presenter, L. Arbour, M.D.). Toronto, Canada: July 1992.
- “Transplantation, Biotechnology & the Law: An Inquiry into Values;” radio interview with Canadian Broadcasting Corporation, Radio Noon. Montreal: June 1992.
- “Evolving Legal Regimes for Tissue Replacement Technology in a Transplant & Biotechnological Era”, Presented to an International Symposium on the Definition of Death & Organ Transplantation at the European Centre or Tokai University (Japan), Vedbaek, Denmark: August 1991.

SELECTED PUBLICATIONS

- Sheppard, NC & Jones, DJ. *Bill C-7's Express Exclusion of Individuals Whose Sole Underlying Medical Condition Is Mental Illness from Canada's Evolving MAiD Regime: (Un) Justified Human Rights Discrimination?* Ottawa: Brief to the Senate of Canada Standing Committee on Legal & Constitutional Affairs, Feb. 2021.
- Sheppard, NC, Thermitus T, Jones DJ. Understanding How Discrimination Becomes Systemic, *Globe and Mail [Toronto]*, 24 July 2020.
- Lu, Y, Jones DJ et al. Transparency ethics in practice: Revisiting financial conflicts of interest disclosure forms in clinical practice guidelines. *PLoS ONE*, 2017.
- Jones DJ, Bush P, Macauley AC. Beyond Consent: Respect for Community in Genetics Research. *eLife Sciences (eLS)*, 2014.
- Jones, DJ, Barkun AN, Lu Y et al. Conflicts of Interest Ethics: Silencing Expertise in the Development of International Clinical Practice Guidelines. *Ann Internal Med* 2012; 156:809-816.
- Jones DJ, Sheppard NC. Advancing America's Right to Health. *Globe and Mail [Toronto]* June 2012.
- Barkun AN, Bardou M, Kuipers EJ, Sung J, et al. International Consensus Upper Gastrointestinal Bleeding Conference Group. International consensus recommendations on

the management of patients with nonvariceal upper gastrointestinal bleeding. *Ann Intern Med.* 2010;152:101-13. (contributing ethics author).

- *Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans* 2d ed., Government of Canada, Ottawa, 2010 (contributing author).
- *The Duty to Share Information in Clinical Trials*, Government of Canada Interagency Advisory Panel & Secretariat on Research Ethics, Working Committee on Clinical Trials Information, Ottawa: 2008 (principal author).
- “Interface of Law and Ethics in Canadian Research Ethics Standards: An Advisory Opinion on Confidentiality, its Limits and Duties to Others,” *McGill Journal of Law & Health* 2007;1:101-116.
- Interagency Advisory Panel & Secretariat on Research Ethics. *Advisory Opinion, Researchers & the Duty to Warn: Limits on the Spectrum of Confidentiality?* (2006), Ottawa: 2007, (principal author).
- Interagency Secretariat on Research Ethics, *Developments in the Literature: Social Sciences and Humanities Research Ethics: Selected Annotated Bibliography*, Ottawa: Dec. 2007 (executive editor & research director).
- Interagency Advisory Panel & Secretariat on Research Ethics, *Interpreting the TCPS, Volume II* (Ottawa: Government of Canada, Dec. 2006) (contributing author).
- Interagency Advisory Panel & Secretariat on Research Ethics. *Advisory Opinion, Occasional Videoconference Meetings for Research Ethics Boards*. Government of Canada, December 2006 (contributing author).
- Interagency Advisory Panel & Secretariat on Research Ethics. *Advisory Opinion, Use of Student Subject Pools in Research*, 2005. (contributing author).
- Interagency Advisory Panel/Secretariat on Research Ethics, *Interpreting the TCPS* (Ottawa: Government of Canada, 2004) (contributing author).
- *Selected Human Research Ethics Norms: Policies, Laws and Guidelines*, Interagency Secretariat on Research Ethics, 2003-2007, (principal author: semi-annual updates) (e-publication).
- Interagency Advisory Panel/Secretariat on Research Ethics, *Process Principles for Ethics Governance Systems*, 2002, with 2006 update (contributing author).
- Interagency Advisory Panel & Secretariat on Research Ethics. *Advisory Opinion, US Ethics Committee Review of Clinical Trials in Canada*, 2003, (principal author).

- *Compendium of Selected International Legal Norms on the Protection of Personal Information in Health Research*. Ottawa, Canadian Institutes of Health Research, 2001 (principal author), 78 pp.
- *Selected Legal Issues in Genetic Testing: Guidance from Human Rights*. Ottawa: Health Canada, 2001, 85 pp.
- *Socio-Ethical Issues in Biotechnology: Emerging Legal Norms in Biotechnology in Selected Countries -- Laws, Conventions & Guidelines*. Ministry of Industry Gov. of Canada, 2000-01.
- *Towards a Coherent Ethics Framework for Biotechnology in Canada*, Canadian Biotechnology Advisory Committee, Ottawa: Government of Canada, 1999.
- *The Ethics Mandate of the Canadian Institutes of Health Research: Implementing a Transformative Vision (1999-2000)* (principal consultant and draftsman for the Canadian Institute for Health Research (CIHR) Interim Governing Council Sub-Committee on Ethics, with B.M. Knoppers et al).
- *Ethical & Legal Issues in the Supply of Blood Products: Should We Sell Blood?* Bayer Advisory Council on Bioethics. Toronto, 1999, 110 pp.
- Tri-Council of Canada (Medical Research Council, Natural Sciences & Engineering Research Council, Social Sciences & Humanities Research Council). *Policy Statement on Research Involving Humans*. Ottawa, 1998 (contributing draftsman, contributing editor).
- "Government & Biotechnology: Ethics Frameworks to Manage Moral Uncertainty & Policy Development," Ottawa, 1998; prepared for the Government of Canada Working Group on the Advisory Body, Ethics & Public Confidence of the Canadian Biotechnology Strategy Task Force.
- "Ethics & Biotechnology: The Role of the Government of Canada," in *Renewal of the Canadian Biotechnology Strategy Roundtable Consultation: Background Documents*. Ottawa, 1998 :1-69.
- "Health Law & Bioethics: Requiem or Renaissance for the Law Reform Commission of Canada," (1995) 46 *International Digest Health Legislation* 117; reprinted with modifications in *Annals of the Royal College of Physicians & Surgeons of Canada* 1996:29(3):167-170.
- National Council on Bioethics in Human Research. "Protecting & Promoting the Human Research subject: A Review of the Function of Research Ethics Boards in Canadian Faculties of Medicine," *NCBHR Communiqué CNBRH* 1995/6(1):3-32 (executive editor & contributing author).
- "Conflict of Interest in Research Ethics," *National Council on Bioethics in Human Research Communiqué CNBRH* 1995/6(2):5-10.
- "Divided Loyalties: An Anthology of Conflict of Interest Duties," *NCBHR Communiqué CNBRH* 1995/6(2):11-16 (principle author and research director).

- National Council on Bioethics in Human Research [of Canada]. "Selected Bibliography: Bioethics of Research with Human Subjects in the Health Sciences (1983-1993)," *NCBHR Communiqué CNBRH* 1995/6(2) supplement: 1-16 (executive editor).
- "Medical Screening & Monitoring in the Federal Workplace" (1994) internal technical document of Department of Justice Canada, 200 pp.
- "Retrospective on the Future: Brain Death & Evolving Legal Regimes for Tissue Replacement Technology" (1993) 38 *McGill L.J.* 396-415.
- *Procurement & Transfer of Human Tissues & Organs*, Law Reform Commission of Canada (1992), 230 pp. (principal author).
- A. Meagher et al., "Doctors & Hospitals: Legal Duties" (1992) 71 *Can. Bar Rev.* 634-639 (book review).
- "Reproductive Hazards in the Workplace" (1992) 147(10) *Can. Med. Assoc. J.* 1412-1414 (letter).
- "Law Reform Commission of Canada Brief to the Royal Commission on New Reproductive Technologies" (1992) 13(1) *Health Law in Canada* 119-124.
- *Medically Assisted Procreation*, Law Reform Commission of Canada (1992) (contributing author to ch. III, Role of the State, esp. pp. 113-119).
- "Brain Death & Evolving Legal Regimes for Tissue Replacement Technologies" (1992) 41 *J. of Behavioral & Social Sciences* 57-74 (Japan).
- *Toward a Canadian Advisory Council on Biomedical Ethics*, Law Reform Commission of Canada (1990), 56 pp. (contributing analyst).
- "AIDS & Disability Employment Discrimination in & Beyond the Classroom" (1989) 12(1) *Dalhousie L.J.* 103-130 (co-author, Colleen Sheppard).
- "Artificial Procreation, Societal Reconceptions: Legal Insight from France" (1988) 36(3) *American J. Comparative L.* 525-545.
- "Time to Treat MD's Ailing Training System" [*Toronto*] *Globe & Mail* (6 January 1988) A7.
- "Les procédures de remboursement des technologies nouvelles aux USA" (août 1984) 82 *Hôpital à Paris* 44.