GLOBAL HEALTH LAW LL.M.

Georgetown’s Global Health Law LL.M. programs train lawyers to apply their specialized legal skills to improve population health in the United States and globally. The innovative full- or part-time courses of study explore the intersections of health and law, including global health governance, health and human rights, food and drug law, access to health care, and legal interventions to prevent and control infectious and non-communicable diseases.

Created in 2007 as part of the O’Neill Institute for National and Global Health Law (http://oneill.law.georgetown.edu), our programs include the LL.M. in Global Health Law (see below), the LL.M. in Global Health Law and International Institutions (https://curriculum.law.georgetown.edu/llm/llm-joint-degree-programs/joint-degrees/jd-llm-global-health-law-international-institutions) with the Graduate Institute of International and Development Studies in Geneva, Switzerland, and certificate programs in Food and Drug Law (https://curriculum.law.georgetown.edu/llm/llm-certificate-programs/llm-food-drug-law-certificate) and U.S. Health Law (https://curriculum.law.georgetown.edu/llm/llm-certificate-programs/llm-us-health-law-certificate). We have more than 180 graduates from 30 countries working to improve the public’s health in the private sector, academia, non-government organizations, international organizations, and all levels of government.

LL.M. in Global Health Law

All Global Health Law LL.M. candidates are enrolled in the 4-credit Global Health Law course, which explores the roles that the law, lawyers, and legal institutions play in public health across the globe. Working with their academic advisor, candidates select additional courses from Georgetown Law’s unparalleled health law curriculum, with more than 35 courses addressing domestic and global health law.

As part of their program, Global Health Law students have the unique opportunity to engage with the O’Neill Institute, a premier research institution working at the intersection of health and the law. Candidates may work on O’Neill Institute projects as research assistants, enroll in its practicum courses, attend symposia and events, and participate in career and professional development programs including a mentorship program that partners students with O’Neill Institute staff and program alumni.

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<thead>
<tr>
<th>Requirement</th>
<th>U.S.-Trained Students</th>
<th>Foreign-Trained Students</th>
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<tbody>
<tr>
<td>Total Credits Required</td>
<td>24</td>
<td>20</td>
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<tr>
<td>Specialization Credits Required</td>
<td>16</td>
<td>14</td>
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<tr>
<td>Program Course Requirements</td>
<td>4-credit Global Health Law Course</td>
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<td>GPA Requirements</td>
<td>Students must earn a minimum grade point average of &quot;B-&quot; (2.67/4.00) in the courses that are counted toward the LL.M. in Global Health Law specialization requirements</td>
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Contact Information
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Please address any questions about admissions the Office of Graduate Admissions (https://www.law.georgetown.edu/admissions-aid/graduate-admissions).

Search LL.M Global Health Law Courses (http://curriculum.law.georgetown.edu/course-search/?program=program_83)

LAW 534 v01 Access to Health Care and Coverage: Law and Policy
(http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20534%20v01)
J.D. Course (cross-listed) 3 credit hours
The law governing access to health care has been in flux and in legal dispute in recent years. This course will examine America’s programs for health care access and finance, including employment-based private insurance, Medicare, Medicaid, CHIP, and VA. The course will also examine the enactment, implementation, and litigation of the Affordable Care Act. In addition to these payment systems, the course will also at laws affecting non-comprehensive systems such as emergency rooms, public hospitals, and community health centers. No previous knowledge of health law is required.

Note: The course will not focus on biomedical ethics, medical malpractice, or pharmaceutical regulation.
This is a required course for the U.S. Health Law Certificate.

LAW 1602 v00 Advanced Topics in Torts: Products Liability, Guns, and Drugs
(http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%201602%20v00)
J.D. Course (cross-listed) 3 credit hours
This upper level course will cover the law of products liability generally and take a close look at the state of products liability litigation and liability in relation to guns and to opioids. The goal is to combine a survey of the complicated field of products liability law with a sophisticated deep dive into two areas of cutting edge products liability litigation. The first part of the course will familiarize students with major topics applicable to all products manufacturers including: a product distributor’s liability for defect-caused harm, allocating responsibility inside and outside the commercial chain of distribution, causation, affirmative defenses, approaches to design defect litigation, and federal preemption of products liability claims. Later in the course, we will examine gun manufacturer liability, currently and prior to the passage of the Protection of Lawful Protection in Arms Act, which reshaped the landscape of gun litigation. Finally, we will end with a consideration of the growing litigation related to the opioids, litigation inflected by doctrines peculiar to prescription drug manufacturer liability. The final examination will be a self-scheduled 48 hour take home exam. Attendance and participation are crucial to the course, and significant credit will be given to those students who contribute thoughtfully and constructively to class discussion of cases and issues.
LAW 277 v02 Aging and Law Seminar

J.D. Seminar | 3 credit hours
This seminar explores, through lecture, discussion, role playing, and problem solving, the demographics, public perceptions, special legal problems, and public policy issues affecting older persons. Subject areas include income maintenance programs (Social Security, SSI); health and long-term care benefits (Medicare, Medicaid, long-term care insurance, state and federal financing issues); retirement housing and long-term care options and regulation (continuing care retirement communities, nursing homes, home and community-based care, home equity conversion); estate and personal planning issues related to incapacity (powers of attorney, trusts, guardianship and its alternatives, elder abuse, the right to refuse life-sustaining medical treatment, bioethical dilemmas, surrogate decision making, and health care advance directives); and ethical issues in representing the elderly. The seminar is both practice- and policy-oriented and integrative with respect to other coursework and related disciplines.

**Recommended:** Prior or concurrent enrollment in one or more of the following courses: Administrative Law; Family Law I: Marriage and Divorce; Constitutional Law II: Individual Rights and Liberties; Professional Responsibility.

**Note:** This seminar requires a paper. Students must register for the 3 credit section of the seminar if they wish to write a paper fulfilling the Upperclass Legal Writing Requirement. The paper requirements of the 2 credit section will not fulfill the Upperclass Legal Writing Requirement.

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LAW 369 v01 AIDS Law and Ethics Seminar

J.D. Seminar (cross-listed) | 2-3 credit hours
This course examines the social, legal, political, and ethical controversies surrounding the HIV/AIDS pandemic in contemporary society. It covers both domestic and international law and policy. The course is divided into several parts. Part I covers the role of social movements and mobilization in the response to HIV/AIDS. Part II, AIDS in the Courtroom, covers the major court cases related to HIV/AIDS in the United States and in key countries around the world like South Africa, India and Brazil that provide important comparative perspectives to understand the power of law. These cases demonstrate the social impact of AIDS—the effect of litigation on social institutions, constitutional law, and interpersonal relationships. Part III, Rights and Dignity, examines the role of international human rights, privacy, and discrimination. Part IV, Policy, Politics, and Ethics, covers a wide range of the most contentious debates of the HIV/AIDS pandemic, including testing, named reporting, civil and criminal confinement, sex work, drug law and policy, LGBT rights, and gender. The final Part, Governance and Financing, examines the absence of political leadership, the international trade system which militates against access to affordable treatment in low- and middle-income countries, the systems of financing for HIV in the U.S. and around the world, and the ethics of international collaborative research. The AIDS pandemic has reached deeply into all major spheres of modern life—e.g., law, medicine, economics, and politics. The pandemic has transformed society and restructured ethical values. This course provides an account of the major themes of the pandemic during the last three decades and offers an analysis of contemporary and future policy.

**Mutually Excluded Courses:** Students may not receive credit for this seminar and the course, AIDS Law and Ethics.

**Note:** This seminar requires a paper. J.D. students must register for the 3-credit section of the seminar if they wish to write a paper fulfilling the J.D. Upperclass Legal Writing Requirement. The paper requirements of the 2-credit section will not fulfill the J.D. Upperclass Legal Writing Requirement.
Alternative, Complementary, and Integrative Medicine ("CAM") is the fastest-growing sector of American Health Care and is one of the fastest growing fields in the United States. Presently, at least 50 percent of Americans are using some form of alternative and complementary therapy such as acupuncture, nutritional supplementation, herbs, massage, yoga, chiropractic and homeopathy. According to the Journal of the American Medical Association in 1997, visits to alternative health care practitioners exceeded total visits to all conventional primary care physicians. The number of clinics and hospitals that integrate some modalities of CAM alongside conventional medicine is growing rapidly. The Institute of Medicine, a part of the National Academy of Sciences, has held recent conferences on the values of both CAM and Integrative Medicine. The NIH is using significant resources to fund research in this area.

This development, of course, is raising legal issues. There is a growing but still unsettled body of law on this subject. Some but not all CAM modalities are now licensed and regulated by at least some states. Federal regulatory bodies, such as the FDA and FTC are trying, within the limits of their statutory authority, to protect what they perceive to be the interests of the public. Yet, they come at the problem through conventional, rather than alternative, eyes. Conventional law is based upon protecting the public from purveyors of the proverbial "snake oil" frauds. And to an extent this law is being used to keep out alternatives to the established health-care modalities. This seminar studies the tensions, legal, economic, and social, of this struggle as it unfolds. This seminar covers several areas of law including administrative law, medical malpractice, informed consent, FDA/FTC law, among others. A paper meeting the upperclass legal writing requirement is required.

Globalization has inevitably forced lawyers, public health professionals, health care professionals, and anthropologists alike to rethink the traditional approaches and methods relied upon within their disciplines. As international borders continue to disappear and countries and people throughout the world become increasingly interconnected and interdependent, public health threats can easily become global in scale and can only be properly addressed through multidisciplinary efforts at global, national, and local levels. Time and time again, the implementation of sound public health measures has proven difficult in communities when local culture and ideology are not considered or properly understood. As governments and international organizations increasingly rely on the law as a fundamental tool for solving critical health problems, it is of the utmost importance that the laws and regulations that they adopt with the aim of protecting and advancing population health, as well as their implementation, properly reflect the social and cultural context of those affected.

Through the analysis of case studies from various areas of global health (including non-communicable diseases, infectious diseases, climate change and health, and gender and health), this course aims to underscore the importance both of incorporating anthropological methods into the practice of global health law and of utilizing a multidisciplinary approach when addressing global health challenges.

Recommended: Prior enrollment in Global Health Law and any coursework in public health, public health law, and cultural anthropology.

Global Health Law LL.M.

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LAW 284 v01 Bioethics and the Law Seminar (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20284%20v01)
J.D. Seminar (cross-listed) | 3 credit hours
This seminar investigates legal, ethical, and social problems raised by developments in health, medicine and the biological sciences through the study of selected subjects that vary from year to year. Issues covered might include death and dying, genomics, reproductive technologies, fetal treatment and research, experimentation with human subjects, and societal controls on scientific advances.

LAW 3038 v00 Biosecurity and the Law (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203038%20v00)
LL.M Seminar (cross-listed) | 2 credit hours
This course is designed to provide students an understanding of current and emerging issues concerning biosecurity and relevant law. Recent global acts of terrorism coupled with sophisticated advances in biotechnology present a host of complex issues driving biosecurity analysis and discussion. We will examine both the history and current state of bioterrorism threats with an emphasis on the legal and ethical challenges as we "sort out" best methods for moving forward. Our analysis will explore the recent Zika and Ebola threats as well as the threats posed by prior H1N1, SARS, anthrax, smallpox, SARIN, Ricin, and even the possibilities stemming from genetically engineered once thought "dead" bio-threats. Our method of instruction will focus on significant legal challenges each week. We will however, use past examples of quarantine and cordon sanitaire applications to instruct as how best to legally address future possible pandemics. Within our course exercises, students will be assigned healthcare and legal roles in order to gain insight as to management of a true healthcare crisis. Guest speakers with knowledge of crisis management will be invited to share personal experiences and advice concerning future pandemics. We will also examine recent legislation, both domestic and international, with a view toward a better understanding of the complex challenges within biosecurity.

LAW 2030 v01 Comparative Reproductive Technologies and "Reproductive Tourism" (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%202030%20v01)
LL.M Seminar (cross-listed) | 2-3 credit hours
The use of reproductive technologies—and crossing national borders to obtain them—has become a burgeoning multi-billion dollar, international industry. While the desire to have children may be universal, legal protections and restrictions on access to reproductive technologies vary immensely from country to country, and often reflect conflicting cultural and religious values.

This seminar will examine the fundamental elements of ART law and practice so that students have a foundation to explore and compare a diverse number of legal systems' approaches to selected reproductive technologies with a particular emphasis on the legal implications for "cross-border reproductive care" ("reproductive tourism"). Other topics will include: comparative access to and affordability of IVF, egg and sperm donation, and surrogacy; reprogenetics; treatment for same-sex couples; professional liability; and embryonic stem cell research (as it intersects with egg donation and the use of IVF embryos). Guest lectures will provide a medical and an ethical perspective to broaden an understanding of the legal and policy challenges in this unique field.

**Note:** This seminar requires a paper. J.D. students must register for the 3 credit section of the seminar if they wish to write a paper fulfilling the Upperclass Legal Writing Requirement for JD students. The paper requirements of the 2 credit section will not fulfill the Upperclass Legal Writing Requirement for JD students.

LAW 056 v00 Biotechnology and Patent Law Seminar (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20056%20v00)
J.D. Seminar (cross-listed) | 2-3 credit hours
This course examines a variety of legal and policy issues associated with the intellectual property protection available to biotechnological innovations, with an emphasis on patents.

Specific issues that will be examined within this framework include legal utility; conception and reduction-to-practice of biological molecules; anticipation and obviousness of nucleic acids; written description and enablement requirements for biological processes and molecules; experimental use; export and import issues; infringement; and potentially developments outside the US.

Course readings rely primarily on cases, statutes, and regulations. A biotechnology background is not necessary.

**Prerequisite:** Patent Law or patent law experience.

**Note:** J.D. students must register for the three-credit section of the course if they wish to write a paper fulfilling the J.D. Upperclass Legal Writing Requirement. A take home exam is required for the two-credit section.
LAW 1101 v00 Consumer Advocacy: Public Health Regulation of Tobacco and Personal-Care Products (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%201101%20v00) (Fieldwork Practicum) J.D. Practicum | 4 credit hours

In fieldwork practicum courses, students participate in weekly seminars and conduct related fieldwork at outside organizations. This fieldwork practicum course will focus on federal regulation of tobacco and personal-care products by the Food and Drug Administration (FDA) under the Food, Drug, and Cosmetic Act and how public interest groups advocate greater protections for consumers. Students will participate in a two-hour/week seminar and carry out one or two 10 or 15 hours/week of fieldwork as interns with one of two national consumer and environmental health organizations: the Campaign for Tobacco-Free Kids or the Environmental Working Group. (Two pass/fail credits will be awarded for 10 hours/week of fieldwork, and three pass/fail credits will be awarded for 15 hours/week of fieldwork.)

SEMINAR: In the two-credit, graded, seminar portion of the practicum, students will utilize legislative and administrative materials as well as case law to become familiar with the processes by which the federal government regulates tobacco and personal-care products, and to critique both the statutory framework and FDA’s performance in protecting consumers. The 2009 Tobacco Act created a new regulatory regime very different from the FDA’s existing authority to regulate other products within its jurisdiction. Examination of the agency’s initial steps to carry out this responsibility provides a chance for students to understand how a federal agency responds to a legislative mandate requiring innovative action to address a major public health problem. On the other hand, FDA has minimal authority to regulate personal-care products, raising real concerns for consumer protection. The course will also touch on related topics such as the role of the Federal Trade Commission in the regulation of trade practices, the Freedom of Information Act, the legislative process and the interaction of federal and state regulation.

FIELDWORK: In the two- or three-credit, mandatory pass-fail, fieldwork portion of the practicum, students will work as interns with the Campaign for Tobacco-Free Kids or the Environmental Working Group on projects aimed at strengthening the legislative or administrative processes, or on matters in litigation, under the supervision of attorneys connected with these organizations. (Students who have completed this course will have priority consideration if they opt to apply for the year-long Toni Stabile Graduate Fellowship at the Environmental Working Group after graduation.)

Prerequisite: J.D. students must complete the required first-year program prior to enrolling in this course (part-time and interdivisional transfer students may enroll prior to completing Criminal Justice, Property, or their first-year elective).

Recommended: Administrative Law; Food and Drug Law-related courses.

Mutually Excluded Courses: Students may not concurrently enroll in an externship or a clinic or another practicum course.

Note: LL.M. students must seek professor permission to apply.

Evening students who work during the day are encouraged to reach out to the professors to determine whether this practicum course would be compatible with their schedules.

This is either a four or a five credit course, depending on the number of fieldwork hours/week. Two credits will be awarded for the two-hour weekly seminar and either two credits (for 10 hours/week) or three credits (for 15 hours/week) for the fieldwork. The fieldwork will be conducted over a minimum of 11 weeks, to be arranged with the faculty members.

Students will have the ability to choose the credit option that best fits their schedules. However, the fieldwork must be completed during normal business hours. All students will initially be registered for a total of four credit hours.

LAW 1102 v00 Drug Law and Policy Seminar: A Critical Perspective on the War on Drugs in the Americas (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%201102%20v00)

J.D. Seminar (cross-listed) | 2-3 credit hours

This seminar will explore the policy of prohibition on drug production, distribution and consumption that embodies the long-standing hemispheric “war on drugs”. It will examine the legal regime that underpins current policies on drug trafficking (international, transnational and domestic regulations) related to criminalization but also to arms’ sale and money laundering. We will reflect on the economic and social consequences that drug-trafficking prohibition has had in countries and communities situated on different sides of the global drug trade. Our focus will be primarily on the United States, Mexico and Colombia. The seminar will explore the dominant “law and order” criminalization paradigm and contrast it with other potential approaches that focus on public health and economic development. We will conclude by considering a range of potential policy alternatives to the current model.

Note: This seminar requires a paper. J.D. students must register for the 3-credit section of the seminar if they wish to write a paper fulfilling the J.D. Upperclass Legal Writing Requirement. The paper requirements of the 2-credit section will not fulfill the J.D. Upperclass Legal Writing Requirement.

LAW 3003 v00 Employee Benefits: Health & Welfare Plans (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203003%20v00)

LL.M Seminar (cross-listed) | 2 credit hours

This course focuses on the tax and ERISA aspects of employer-sponsored health and welfare benefit plans. The tax discussion will concentrate on the conditions for favorable treatment of health and welfare benefits (and other statutory fringe benefits), the cafeteria plan rules, the applicable nondiscrimination requirements, and the special rules applicable to funded welfare benefits. The ERISA discussion will focus on plan design, reporting and disclosure, claims procedures, and fiduciary duty rules. The course will integrate the tax and labor aspects of the Affordable Care Act.

Prerequisite: Federal Income Taxation (formerly Taxation I).

Mutually Excluded Courses: Students may not receive credit for this course and Health and Welfare Benefit Plans: Tax & ERISA Aspects.

Note: This course is required for the Employee Benefits Certificate.

This course is open to J.D. students by professor permission. Interested students should contact Ellis Duncan via email at ged5@law.georgetown.edu no later than August 1, 2018 for permission to take this class. Students may not withdraw from this class after the add/drop period ends without the permission of the professor.
LAW 754 v01 Epidemiology for Lawyers
This course will introduce students to the foundational laws and policies governing the production and distribution of foods, drugs, devices, cosmetics and dietary supplements in the United States, focusing on the Federal Food, Drug, and Cosmetic Act (the "Act") and the role of the Food and Drug Administration in enforcing the Act. The course will cover key concepts and definitions -- e.g., "food," "drug," "labeling" - and federal statutory provisions designed to assure that such products are not adulterated or misbranded. Students will also receive an overview of the different agencies that have jurisdiction over foods, devices and drugs on the state and federal levels, as well as an introduction to the ways in which FDA exercises its authority under the Act through rulemaking and guidance. Time permitting, there will be a portion of the course devoted to the extent to which third parties -- outside of the regulatory agency and the regulated business -- seek to influence policy and decision making through communications and lobbying.

Note: This is a required course for the Food and Drug Law Certificate.

LAW 1208 v00 Food Law Seminar
This course will introduce students to the foundational laws and policies governing agriculture, including laws and regulations related to farm subsidies, farm stewardship, biotech regulation, food safety, food labeling, food assistance, farm labor, animal welfare, agricultural trade, and antitrust issues related to crop and livestock production.

The Farm Law and Policy Seminar complements other courses offered by the Law Center, including courses on Food Law and Environmental Law.

Recommended: A course in food law or environmental law.
LAW 1272 v00 Gender and Sexuality (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%201272%20v00)
J.D. Course (cross-listed) | 3 credit hours
This course will provide an introduction to the legal contexts and theoretical debates surrounding sex, gender, sexuality, and their intersections. We will explore the ways that gender and sexuality have been debated, defined, and redefined in the contexts of gender identity and performance, sexual pleasure, reproductive rights, sexual violence, marriage, family organization, and work. In each context, we will canvas the evolution of the law as we consider how feminist and queer theorists have conceptualized gender and sexuality in order to reimagine and critique prevailing legal rules and cultural norms. In short, the class will probe the ways that law is gendered, sexualized, and raced, and with what overall effects on social institutions and practices.

Key topics will include:

- The influence of identitarian politics on law and vice versa
- Regulation of Sexual Conduct
- Regulation of Reproduction
- The Meanings of Same Sex Marriage
- Sex, Law & Consent
- Gender & Sexuality at Work
- Equality, Stereotypes, and Pregnancy
- Sexual Harassment

Students will be graded primarily on the basis of a take-home exam at the end of the semester with some consideration of class participation. There may also be short response/essay papers or small group projects required.

Strongly Recommended: Constitutional Law II.

Mutually Excluded Courses: Students may not receive credit for this course and Sexual Orientation and the Law: Selected Topics in Civil Rights.

LAW 3028 v00 Global Drug Law and Regulation (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203028%20v00)
LL.M Seminar (cross-listed) | 2 credit hours
Globalization and the international trade of drugs and medical products have progressed beyond any single regulatory authority's ability to effectively ensure the quality, safety, and effectiveness of these products. In the U.S., the importation of foreign sourced products has increased tremendously, accounting for over 80% of the active pharmaceutical ingredients. However, varying drug regulations have resulted in gaps in oversight causing differing views on the acceptable level of risk in public health leading to drug quality-related deaths and other serious harms. One clear reason for this compromised system is the differences in how these products are regulated from country to country. Nevertheless, the pharmaceutical and related industries are thriving in the global marketplace. This course is intended to be the first comparative survey into the regulatory frameworks of certain key countries, both developed and developing markets, along with international institutions, such as the World Health Organization, involved in promoting the access and development of safe, effective, and quality medical products. This course will also identify the major international non-governmental stakeholders, and the multi-lateral schemes and treaties in which they operate that are intended to assist in the convergence of pharmaceutical laws and regulations.

Recommended: Prior enrollment in Food and Drug Law

LAW 493 v01 Global Health Law (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20493%20v01)
LL.M Course | 3-4 credit hours
Global Health Law is the flagship course for Georgetown University Law Center’s O’Neill Institute for National and Global Health Law. This course is open to both Georgetown JD and LLM students and is a compulsory unit in the Global Health LLM.

No longer an emerging field, global health law encompasses international law and policy that directly or indirectly affects global health, including treaties, regulations, global strategies, and non-binding standards, national and international jurisprudence etc. The field of study includes both legal instruments designed to protect public health as well as the interaction between legal instruments from other international legal regimes and public health considerations and concerns. This course provides a strong foundation in these laws and policies, including governance of the World Health Organization, the International Health Regulations, the WHO Framework Convention on Tobacco Control, international trade and investment law, and human rights treaties such as the International Covenant on Economic, Social, and Cultural Rights.

In examining the application and effectiveness of global health law, this course provides a normative foundation for global health issues including infectious diseases (such as Ebola, HIV/AIDS, tuberculosis, malaria, and influenza) and noncommunicable diseases (such as diabetes, cancer and cardiovascular disease and their causes, including obesity, tobacco, and alcohol).

In this course, students will hear from leading voices in global health and the law and benefit from the expertise of Georgetown Law’s O’Neill Institute.

Recommended: Prior enrollment in International Law I.

Note: Required for the Global Health Law LL.M.
LAW 594 v00 Global Health Law: An Intensive, Problem-Based Exploration

In this intensive course, students will work with faculty and fellows at the O'Neill Institute for National and Global Health Law to gain an in-depth understanding of global health law through intensive examination and role play of one or more major problems in global health. Potential problems could include a naturally occurring infectious disease epidemic such as extensively drug-resistant tuberculosis; a future epidemic such as pandemic influenza (A) H5N; an intentional introduction of a lethal pathogen such as anthrax; and/or major chronic diseases caused by obesity or tobacco use. Students should come to this course with a basic level of understanding of global health law, including the major international health treaties and governing structures. When studying and role playing these kinds of problems, students will be asked to construct innovative methods of global health governance, drawing upon existing international health law and institutions, along with a vision for more ideal models. The course will also capitalize on materials developed in collaboration with faculty from the School of Nursing and Health Studies for use in the “health care situation room”.

Full attendance and participation is required at all sessions. Class sessions will consist of a combination of lecture, case simulations, and discussion. Grades are based on student participation, a daily journal to be kept by students, and a final paper.

Note: The first weekend of the course will be held at the Law School and the second weekend will be on the Main Campus at St Mary's at the School of Nursing and Health Studies.

A student will be permitted to drop a course that meets for the first time after the add/drop period, without a transcript notation, if a student submits a written request to the Office of the Registrar prior to the start of the second class meeting. Withdrawals are permitted up until the last class for this specific course. ATTENDANCE IS MANDATORY AT ALL CLASS SESSIONS. All enrolled and waitlisted students must be in attendance at the start of the first class session in order to be eligible for a seat in the class and must attend each class session in its entirety. This course requires full participation at all class sessions.

LAW 1028 v00 Health Care Fraud and Abuse Seminar

One-fifth of the U.S. economy centers around health care industry sectors. This seminar examines criminal, civil and administrative tools used by federal and state enforcement authorities to police the U.S. healthcare system. We will focus on cases brought under federal and state False Claims Acts (FCA), the Anti-Kickback Statue (AKS), Stark laws, Federal Food Drug and Cosmetic Act (FDCA), and Federal Corrupt Practices Act (FCPA). The seminar provides a survey of the enforcement activities of the U.S. Department of Justice (DOJ), the Office of Inspector General at Department of Health and Human Services (OIG), and state Medicaid Fraud Control Units (MFCUs) in matters against pharmaceutical and medical device manufacturing companies, physicians, hospitals, clinical practices, nursing homes, laboratories, and others. The seminar materials thoroughly cover the statutes, safe-harbors, and regulations that govern the health care industry. We will also discuss risk mitigation strategies and compliance program best practices across industry sectors to provide insight into the impact enforcement has on (1) clinical decision-making, (2) costs to providers, payers, and patients, (3) patient safety, and (4) quality of care. In an effort to maintain a broad perspective with the diverse and frequently changing legal landscape in the area, in addition to the case book, materials discussed and presented in this course draw from news reports, trade publications, and U.S. government agency materials.

The class requires a paper of approximately 20-25 pages in length.

Recommended: Criminal Justice (or Democracy and Coercion) or Criminal Procedure.

LAW 2037 v00 Health Information Technology and the Law

Health care decision-making and innovation are increasingly driven and made possibly by vast stores of data. The importance of data has created an inevitable push-pull dynamic between concerns for confidentiality and demands for medical progress and cost containment. Data is both a privacy risk and a tremendous asset. This course will explore the legal and ethical issues at the intersection of health information, including where data comes from, how it is and should be protected, how it can be used, and risks to its integrity and security. In doing so, this course will cover a range of topics including health information privacy, future use of data assets, and conflicts of interest.

LAW 206 v03 Health Law and Policy

Health law is a vast and expanding field. No single course can survey it all. This course focuses on the organization, financing, and provision of medical care, with an eye toward issues not yet resolved by courts, legislators, regulators, and American society. It also considers some related ethical questions. Topics and themes include the economics of health insurance and managed care, regulatory responses to the market's perceived failures, medical tort law, access to care, consumer choice and patient autonomy, defining and assessing quality, health care providers' conflicts of interest, privacy and confidentiality, and socio-economic and racial disparities in health and medical care.
Beyond health insurance and the delivery of health care, goods and services related to individual and public health are highly regulated in the United States, and often serve as a basis for international regulations. These goods and services are a large and growing part of the U.S. and world economy, with some estimates being that more than one-quarter of U.S. food and medical products are regulated by the FDA alone. This regulation is carried out directly by a variety of State and Federal agencies (such as the FDA, the CDC, and the NIH) as well as indirectly through the purchasing power of federally financed programs, such as Medicare.

This course will include an introduction to the basic legal and regulatory frameworks within public health and an overview of the Constitutional limits and policy choices that have led to current law. The course will then move to a review of several major fields of regulation. From a high-level, this includes the regulation of health professionals, health systems, and medical or food products impacting human health. The course will then conclude with an examination of several contemporary problems, such as the safe and effective use of human drug products, infectious-disease prevention and control, ethical research practices, and rationing and allocation of limited resources.

The primary objective of the course is to teach students about the regulation of public health at the intersection of state and federal levels, recognizing that such regulatory frameworks often become the template for international policies. Students will be called on to learn the basics of two fundamental statutes—the Public Health Service Act (PHSA) and the Food, Drug and Cosmetic Act (FDCA). Students will also be called upon to follow examples of administrative change under these statutes, each year analyzing a new set of proposed regulations and sub-regulatory guidance documents. By the end of the course, students will be able to describe the major means by which goods and services used in both personal and public health are controlled, as well as areas in which future changes are likely.

Currently, there is no text or case book on this subject. The primary readings will be assigned by the professor.

LAW 3054 v00 Health Rights Litigation Intensive (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203054%20v00)
LL.M Course (cross-listed) | 2 credit hours
This one-week intensive course offers students an opportunity to develop specialist-level knowledge in litigating health-related rights at the national, regional, and international levels.

During the course, globally renowned experts will lecture on a range of topics, including: regulation of private actors; sexual and reproductive health and rights; rights issues arising in health-care settings; approaches to health-care rationing and factors to consider in assessing the equity impacts of judgments; access to medicines and intellectual property; judicial legitimacy in deciding issues with budgetary and policy implications; and judicial effectiveness and impact of judgments.

The course is highly participatory, and uses case-based teaching and group exercises extensively. Students will be evaluated based on their participation in lectures and group exercises throughout the week, as well as their participation in either the moot court competition or the fundraising pitch on the final day of the course.

Recommended: Prior enrollment in International Human Rights; International and Comparative Law on Women’s Human Rights; Gender, Sexual and Reproductive Health and International Human Rights Law; O’Neill Institute Practicum: Health and Human Rights

Note: This class will meet on the following Summer 2017 dates: 6/26, 6/27, 6/28, 6/29, and 6/30.
LAW 3058 v00 Health, Human Rights and Social Justice (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203058%20v00)

LL.M Seminar (cross-listed) | 1 credit hour

Human rights is now the dominant language for claims of human emancipation around the world; human rights theory and practice have permeated many domains beyond the law, including health. Yet the landscape of global health is marked by vast inequities and brutal deprivation, and it is not yet clear how bringing human rights concepts and strategies to bear will change the lives of the millions of people around the globe who are suffering. In this course, we will explore these questions and see how human rights provides not the only, but one, critical framework and set of tools through which to advance social justice in health. Nonetheless, the use of human rights to advance social justice faces vexing challenges, including being reduced to rhetoric by powerful actors and becoming overly legalistic.

The class will explore the conceptual and practical implications of adopting human rights frameworks relating to health policymaking and programming, including emphases on accountability, participation and non-discrimination. We will examine how human rights discourses are shaped and contested, and how this determines the relevance of ‘human rights-based approaches’ to addressing the health needs of different populations. Throughout the course, as we discuss specific issues, we will examine potential limitations as well as strengths of using human rights to improve global health.

The course seeks to answer the following questions:

- What are international human rights standards that relate to health?
- What does it mean in practice to set out a “right to health,” and how might such a right be implemented?
- What is (and should be) the role of courts in enforcing health rights?
- What are the key elements of ‘rights-based approaches’ in programs and policies, with reference to specific health issues and affected populations?
- How might adopting a rights-based approach to global health issues challenge traditional human rights assumptions and practices?
- How can human rights be used to create meaningful social change in health, and what are the limitations to using human rights frameworks?

LAW 1408 v00 Human Genetic Engineering: Law and Policy (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%201408%20v00)

J.D. Seminar (cross-listed) | 2 credit hours

This two-credit seminar will open a window into the fast-developing world of human genetic engineering. It begins with a review of international and regional efforts to ban or restrict human germline modification (HGM), along with a brief world overview of relevant laws and trends. It then focuses on cutting-edge techniques like CRISPR/Cas9 and organized research efforts, particularly in China, that may nonetheless spark a race to create designer babies within a decade or less, as regulation lags behind technology and human affairs. Next, we examine two more well-established reproductive technologies, mitochondrial replacement and pre-implantation genetic diagnosis (PGD), to identify forces that are likely to guide HGM regulation as relevant technologies become safer and more efficient. We'll study the U.K.'s recent adoption of mitochondrial transfer to reduce birth defects or enhance fertility and then learn about evolving U.S. policy. We'll follow the spread of PGD, initially used to identify embryos bearing genes causing incurable childhood diseases, first to other less serious conditions, then to the creation of “savior siblings” and finally to non-medical uses like sex selection. Then we'll return to HGM and view a sample of public policy proposals and religious views likely to influence the coming debate. Finally, we'll close with an introduction to futuristic impulses to implement theoretically limitless improvements to human capabilities, sometimes balanced by the desire to use HGM to improve human moral character and tempered by doubts regarding the moral status to be accorded new HGM creations.

LAW 233 v01 Intellectual Property and Medicines (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20233%20v01)

J.D. Seminar (cross-listed) | 3 credit hours

This course examines the special legal and policy issues arising from the use of intellectual property rights in the pharmaceutical and biotechnology fields. The course will cover: (1) U.S. case law impacting intellectual property, patents, trademarks and copyrights in the pharmaceutical and biotechnological arts; (2) the interplay of the regulatory approval process for therapeutic and diagnostic products with intellectual property rights; (3) the Hatch-Waxman Act and its impact on how patent rights for pharmaceuticals are procured and enforced; and (4) major legislative developments affecting the use of intellectual property rights in the drug, biotechnology and medical device fields, such as the Biologics Price Competition and Innovation Act of 2009 and the America Invents Act of 2011. Other topics may be included depending on current judicial or legislative developments. A background in biologics or pharmaceuticals is not required, although completion of a basic patent law or a food and drug law course is recommended.

Students will have the option of taking this course for either two or three credits. The three credit option will require a paper that satisfies the upperclass legal writing requirement in compliance with Law Center regulations. The two credit option will require completion of several shorter legal writing samples on student-selected or assigned topics.

**Strongly Recommended:** Prior or concurrent enrollment in a basic patent law course or food and drug law course is highly recommended.

**Note:** This seminar requires a paper. J.D. students must register for the 3 credit section of the seminar if they wish to write a paper fulfilling the Upperclass Legal Writing Requirement. The paper requirements of the 2 credit section will not fulfill the J.D. Upperclass Legal Writing Requirement.
LAW 802 v01 International Assistance for Global Health (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20802%20v01)
LL.M Course (cross-listed) | 1 credit hour
The overall objective of this course is to explore the present and potential role of international human rights law – and the right to health in particular – in international assistance for global health.

During the first class, we will discuss the recent history and present practice of international assistance for global health. We will discuss tensions between the objectives of development and relief, and between the objectives of promoting 'health security' and equity in global health. During the second class, we will explore the concept of equity in global health: how it is central to several definitions of global health, what the practice of international assistance for global health should look like, if the predominant objective of that assistance were equity. During the third class, we will examine the meaning of the right to health, the freedoms and entitlements it generates and the corresponding national and international responsibilities. During the fourth class, we will discuss if and how the right to health can be used to modify/improve the practice of international assistance for global health, taking into account that such assistance serves other objectives as well.

Strongly Recommended: Completion of coursework in the area of international human rights law.

Note: A student will be permitted to drop a course that meets for the first time after the add/drop period, without a transcript notation, if a student submits a written request to the Office of the Registrar prior to the start of the second class meeting. Withdrawals are permitted up until the last class for this specific course. The take-home exam in this course may be administered mid-semester and the specific exam date will be provided by the professor after the add/drop period.

LAW 691 v00 International Trade and Health (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20691%20v00)
LL.M Seminar (cross-listed) | 2 credit hours
The objective of this 2 credit seminar is to introduce students to the relationship between international laws governing trade and efforts to protect and promote human health. The course will focus on the impact of the law of the World Trade Organization (WTO) on domestic health measures as well as on international efforts carried out under the auspices of the World Health Organization. Students will learn, and be asked to think critically about, how the international trade regime affects national regulation in the interests of human health.

The course will address the following subject matter:

1. Introduction to 'trade and health': issues and underlying theories.
2. The prohibitions and exceptions of the General Agreement on Tariffs and Trade in a health context.
3. Risk regulation, the precautionary principle and sanitary measures (such as import safety measures).
4. International standards and technical barriers to trade, such as labeling measures.
5. Trade in health services, health worker migration and the General Agreement on Trade in Services.
6. Access to essential medicines under the Agreement on Trade-Related Aspects of Intellectual Property Rights (TRIPs) and TRIPs plus agreements.
7. The fragmentation of international law: cross-fertilization between international trade law and global health law.
8. Policy coherence, legitimacy and participatory governance at the WTO.

Note: A student will be permitted to drop a course that meets for the first time after the add/drop period, without a transcript notation, if a student submits a written request to the Office of the Registrar prior to the start of the second class meeting. Withdrawals are permitted up until the last class for this specific course.

LAW 3006 v00 International Right to Health (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203006%20v00)
LL.M Course (cross-listed) | 2 credit hours
The course provides an introduction to the human right to health and its implementation in a few hot topic issues. Issues covered will include criminalization of drug use and sexual and reproductive health, vulnerable groups, HIV/AIDS, the intellectual property regime and access to medicines, and a critique of the current framework and the challenges that impede the realization of the human right.

Recommended: Introductory course in public international law or human rights, introduction to public international law and/or introduction to human rights

Note: A student will be permitted to drop a course that meets for the first time after the add/drop period, without a transcript notation, if a student submits a written request to the Office of the Registrar prior to the start of the second class meeting. Withdrawals are permitted up until the last class for this specific course.
LAW 708 v00 International Trade, Intellectual Property Rights, & Public Health

This course will cover the interface between the intellectual property rights, international trade and public health, focusing in particular on the WTO TRIPS Agreement and subsequent decisions. It will provide an introduction to the provisions of WTO agreements relevant to public health (other than TRIPS), and to the law and economics relating to IPRs and public health; it will cover the provisions of the TRIPS Agreement relevant to public health, and discuss the relevant disputes settled in the WTO. It will examine the background, content and implications of the Doha Declaration on the TRIPS Agreement and Public Health and of the subsequent TRIPS amendment implementing compulsory licensing for exports. It will also discuss the relevance of bilateral or regional free trade area agreements to the subject.

The course would study relevant national/regional implementing legislation, for example on compulsory licenses, and discuss use of the WTO export compulsory license provisions, namely the Rwanda-Canada case. In addition to the final paper, students will be graded on class participation, individual presentations and group exercises, an example of the last being a hypothetical case study of exports, with students taking up the role of legal advisors based in either the importing country or the exporting country.

Finally, the course will also cover recent work on trade, intellectual property and public health in other intergovernmental organizations, in particular in the World Health Organization.

Recommended: Coursework in International Trade, Intellectual Property Rights, or Public Health.

Note: ATTENDANCE IS MANDATORY AT ALL CLASS SESSIONS. All enrolled and waitlisted students must be in attendance at the start of the first class session in order to be eligible for a seat in the class and must attend each class session in its entirety.

A student will be permitted to drop a course that meets for the first time after the add/drop period, without a transcript notation, if a student submits a written request to the Office of the Registrar prior to the start of the second class meeting. Students may not withdraw from this class after the start of the second class session without the permission of the professor.

LAW 2071 v00 Law and Functioning of International Organizations

Multilateral diplomacy and the international organizations that support much of it have come to play a pervasive role in international relations particularly since the end of World War II. These international organizations serve as instruments of consultation, co-operation and standard setting in almost every sphere of public policy and governmental activity. This may involve broad concerns such as the maintenance or restoration of international peace and security, protection of economic development and stability, advancement of human rights, protection of health, protection of the environment, and the facilitation of trade and investment. Or they may address more specific tasks such as settlement of disputes, codification and progressive development of international law, civil aviation and maritime safety and security, protection of intellectual property, understanding world weather, internet governance, management of the geo-stationary orbit, police cooperation, education and biotechnology. They vary greatly in nature, mission, powers, structure and size. There are inevitable conflicts and frictions in their functioning.

It will examine a number of aspects common to the universe of public, i.e., intergovernmental, international organizations: their legal nature; their need, as the collective instruments of their member states, to remain independent of the individual control of these states, including control through the exercise of some normal state jurisdiction; their immunities and counter-balancing obligations to co-operate; their special body of labor law, including the duties and ethics of the international civil service and the use of international administrative tribunals to adjudicate their employment disputes. The course will examine a sampling of international organizations: how they are governed and financed, how they go about fulfilling their functions, the powers and instruments they use to advance agreed policies, some examples of their output including the development of soft law and hard law instruments and the increasing involvement of civil society in monitoring and influencing their actions. It will also consider the crucial advisory and operational roles lawyers play in a typical international organization, whether as members of the secretariat or of member state delegations.

Students will have the option of preparing a paper that they may elect, after grading, to include in the calculation of their final course grade.

Recommended: Prior enrollment in Public International Law or International Law I.

LAW 199 v03 Law and Regulation of Drugs, Biologics and Devices

This course explores the legal, regulatory and policy issues that shape the research, development, and commercialization of drugs and biologics in the United States and Europe, with a particular emphasis on public policy issues. We will consider: the role of Federal, State and international regulation; regulatory and ethical issues in the development and testing of new therapies; managing incentives for innovation, including patent, regulatory and data exclusivity; tort liability and its function in the regulation of the pharmaceutical industry; pricing and payment systems and controls; and evolving medical technologies.

Recommended: Prior or concurrent enrollment in Administrative Law.

Note: This is a required course for the Food and Drug Law Certificate.
Technological advances in diagnostics and therapeutics have the potential to revolutionize health care and improve the lives of millions of people. However, many of these technologies remain out of reach to those who need them, particularly the poor in low- and middle-income countries.

With increased investment and global attention over the last decade, there has been tremendous progress in building a pipeline of candidate medical technologies to meet the health needs of the world’s poorest people. This course will explore the legal, regulatory and policy issues that are shaping the research, development, and delivery of those drugs, vaccines, and diagnostics. Part I of the course will provide an overview of the burden of neglected diseases in low- and middle-income countries and the new institutions and initiatives that have arisen to address that burden. Part II will examine the incentives for global health innovation, including intellectual property management, regulatory and tax incentives, and prizes and advance market commitments. Part III will consider the role of national and international regulation, international clinical trials and the globalization of research, and World Health Organization’s policy processes for ensuring drug and vaccine safety and recommending their use. Part IV of the course will explore the legal, regulatory and policy issues that arise in the delivery and use of global health technologies, particularly supply chain contracting, drug resistance, and post-market surveillance in low- and middle-income countries.

Law’s relationship with tobacco, alcohol and food is complex and contested. Nevertheless, governments around the world are experimenting with a wide range of legal strategies to encourage healthier lifestyles. This course places U.S. developments in a global and comparative context, offering comparisons with legal strategies for encouraging healthier lifestyles in Australia and other countries.

During the course, we will confront some important over-arching questions. What are the global determinants of NCDs, and how are these diseases being managed, globally? What do the global solutions look like? To what extent should law intervene to influence the behavior of populations – as distinct from treating lifestyle-related risk factors as the personal responsibility of each individual? Does a regulatory approach to the prevention of NCDs imply coercion? Does it signal the emergence of the “nanny state”? Does progress necessarily depend on motivating people to consciously improve their habits and lifestyles? Is it possible to regulate business without micro-managing or dictating commercial decisions and “legislating the recipe for tomato ketchup”?

Most people want to live longer and healthier lives. Yet no country can achieve this without addressing the preventable risk factors that drive non-communicable diseases. This course gives students the conceptual tools to think powerfully about law’s role in the prevention of NCDs, and to participate effectively in debates about appropriate, workable, legal interventions.

Note: WEEK ONE COURSE. This seminar will meet for one week only on the following days: Monday, January 6, 2020 through Friday, January 10, 2020, 9:00 a.m. - 11:35 a.m.

This course is mandatory pass/fail and will not count toward the 7 credit pass/fail limit for J.D. students. Attendance at all class sessions is mandatory and all enrolled students must attend the first class in order to remain enrolled. Students on the wait list must attend the first class in order to be admitted off the wait list. Enrolled students will have until the beginning of the second class session to request a drop by contacting the Office of the Registrar. Once the second class session begins, students may only seek a withdrawal by contacting their academic advisor in the Office of Graduate Programs. Withdrawals are permitted up until the last class for this specific course.
Despite significant progress in legal protections for — and the visibility of — LGBT people over the past decade, LGBT communities continue to face systemic obstacles to quality health care such as refusals of care, substandard care, and inequitable policies and practices in health care settings. These experiences of discrimination correlate with significant health disparities, including greater exposure to violence, higher rates of tobacco and other substance use, mental health concerns, HIV, and cancer. These disparities are even more pronounced for LGBT people who are also members of other groups that face discrimination because of race, ethnicity, or other aspects of identity — such as people of color and immigrants, among others.

In this seminar, students will learn about LGBT health law and policy issues from a variety of perspectives — including medicine, public health, women's studies, and U.S. foreign policy — and gain a better understanding of the social mistreatment and ostracism of LGBT people at both the individual and community level. Topics covered will include international human rights law, LGBT-inclusive data collection, clinical and cultural competency, and health issues facing LGBT young people and elders. This course will also examine the ways in which LGBT individuals and families are treated under federal, state, and international (U.S. foreign policy) law and how these policies impact access to health care and contribute to health disparities.

“Into each house I go,” the Hippocratic Oath promises, “I shall go only for the good of my patients.” We think of doctors as devoted to their patients’ well-being, but they increasingly serve social and legal purposes. To control medical costs, physicians ration care, often unbeknownst to their patients. To protect us from foreign enemies, doctors wage war, designing and overseeing the interrogation of terror suspects. When threats to public health loom, physicians make clinical decisions that protect society at their patients’ expense. In our criminal and civil justice systems, medical judgment answers moral and legal questions about the scope of personal responsibility, the reach of civil rights law, and more. And in our politics, medical opinion both masks and imposes moral and cultural beliefs. This Seminar will explore medicine’s myriad social and legal roles, with an eye toward conflict between these roles and physicians’ traditional commitment to their patients’ interests.

As part of this class you will be involved in projects designed to support the establishment of an MLP at Georgetown. The latter part of the semester will also include joint classes with first-year Georgetown medical students enrolled in a medical school “selective” on MLP. Travel to main campus for those classes will be provided, timing and details will be arranged at the start of the semester. Students will be evaluated based on the quality of their class participation, several short written assignments, and their MLP project. There is no end-of-semester exam.

**Required Text**

TBA
LAW 593 v00 National and Global Health Law: O'Neill Colloquium
(http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%20593%20v00)
J.D. Seminar (cross-listed) | 2-3 credit hours
In this interdisciplinary colloquium, leading national and international scholars in a range of domains will explore fundamental normative and policy problems of contemporary concern in health law. Topics will include health care, public health, global health, science, regulation, politics, ethics, and policy. The colloquium will have participants from across the Georgetown University campus among faculty, senior administration, and students, as well as participants in the Washington health policy and legislative community.

Each seminar session will focus on a presentation by, or structured dialogue with, distinguished guest speakers. Students from the Law Center and other schools within Georgetown University (including Nursing and Health Studies, Medicine, Arts and Sciences, Foreign Service, Business, and other graduate programs) will be expected to prepare for intensive discussions in which experts, faculty and students explore, analyze and deepen their understanding of issues selected for consideration each month. The colloquia will be open to other students and faculty members across Georgetown University as well as interested members of the public, particularly professionals working in health law and policy in Washington.

Note: This seminar requires a paper. J.D. students must register for the 3 credit section of the seminar if they wish to write a paper fulfilling the Upperclass Legal Writing Requirement. The paper requirements of the 2 credit section will not fulfill the J.D. Upperclass Legal Writing Requirement.

LAW 2099 v00 Nutrition Law and Policy
(http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%202099%20v00)
LL.M Course (cross-listed) | 2 credit hours
Proper nutrition is one of the many contributor’s to overall health and could be one of the most cost-effective approaches to address many of societal, environmental and economic challenges facing the world today. Increasingly, law and policy has been recognized as a high-impact and robust approach for accelerating progress toward reducing and managing nutrition-related chronic diseases such as obesity, cardiovascular disease, type 2 diabetes mellitus and certain types of cancer. In various jurisdictions, policymakers enact courses of action, regulatory measures, laws and policies, and set funding priorities designed to address food insecurity, hunger, obesity prevention, chronic diseases, among other health and well-being concerns. This course focuses on policies, programs and practices across the globe—at the national, tribal, state and local levels—that improve or hinder healthy eating. Students will examine the evidence informing these courses of action, along with the historical and contemporary legislative, regulatory and judicial aspects. Topics and themes include dietary guidance, food and nutrition assistance programs, food and nutrition labeling, and other environmental and policy strategies to improve access to healthier foods and beverages.

Student Learning Goals:
• Identify key law and policy approaches used across the globe that have been used or have the potential to improve or hinder healthy eating;
• Discuss and debate the historical and contemporary legislative, regulatory and judicial aspects of the key nutrition law and policy approaches during collaborative in-class exercises; and
• Execute analytical and strategic planning for developing, implementing, evaluating, and sustaining a nutrition law and policy approach during in-class exercises and as part of the mid-term and final projects.

Recommended: Constitutional Law, Property, Contracts, Torts, as well as Food and Drug Law, Introduction to Health Law, and Land Use and Planning.
LAW 1209 v01 O’Neill Institute Practicum: Health and Human Rights (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW %201209%20v01) (Project-Based Practicum)
J.D. Practicum | 4 credit hours
In a project-based practicum course, students participate in a weekly seminar and work on a project under the supervision of their professors. This project-based practicum course will give students the opportunity to work with Georgetown Law’s O’Neill Institute (http://www.law.georgetown.edu/oneillinstitute/index.cfm) and its civil society partners to use international human rights law to advocate for positive health outcomes. Students will participate in a two-hour/week seminar and carry out 10 hours/week of project work under the direction of the course professors.

SEMINAR: In the seminar, students will explore the connections between global health and human rights. We begin by examining the emergence of health and human rights as a distinct field. Following this, we will carefully consider the meaning of the international right to health, stressing the material differences between civil and political rights, on the one hand, and economic, social, and cultural rights on the other. After exploring a series of foundational themes and issues through the first half of the semester, the remainder of the class will focus on in-depth case studies (e.g., HIV/AIDS, mental health, access to essential medicines, reproductive rights). While we will use the International Covenants on Economic, Social and Cultural Rights, and on Civil and Political Rights as the dominant legal instruments in our analyses, our discussions will also reference other international instruments, as well as a variety of national cases, constitutions and legislation. Class time will be devoted to developing practical advocacy and drafting skills to support students in their project work. Students will also learn how to use epidemiological data to support and craft compelling arguments for advancing the right to health.

PROJECT WORK: Students will work with external partners of Georgetown Law’s O’Neill Institute on legal and policy projects related to health and human rights. For example, students may draft alternative reports to UN bodies analyzing compliance with human rights obligations related to tobacco control (e.g., “shadow reports”). A report such as this could analyze the prevailing legal frameworks for ensuring universal access to maternal health services in a particular country and highlight any weaknesses in the statutory and regulatory language. A report such as this would also suggest recommendations for the UN body to consider. By working with the O’Neill Institute, Inter-Governmental Organizations (e.g., PAHO), and civil society organizations, the course will give students the opportunity to use international human rights law to advocate for legal mechanisms to address critical health challenges.

Prerequisite: J.D. students must complete the required first-year program prior to enrolling in this course (part-time and interdivisional transfer students may enroll prior to completing Criminal Justice, Property, or their first-year elective).

Mutually Excluded Courses: Students may not concurrently enroll in this practicum course and a clinic or another practicum course. Students may concurrently enroll in this practicum course and an externship.

Note: This practicum course is open to LL.M. students, space permitting. Interested LL.M. students should email Louis Fine (fine@law.georgetown.edu) to request admission.

Evening students who work during the day are encouraged to reach out to the professor to discuss whether this practicum course would be compatible with their schedules.

This is a four-credit course. Two credits will be awarded for the two-hour weekly seminar and two credits will be awarded for approximately 10 hours of supervised project work per week, for a minimum of 11 weeks. Both the seminar and the project work will be graded.

LAW 1209 v02 O’Neill Institute Practicum: Regulating Alcohol, Tobacco & Food in International and Comparative Law (http://curriculum.law.georgetown.edu/course-search/?keyword=LAW %201209%20v02) (Project-Based Practicum)
J.D. Practicum | 4 credit hours
In a project-based practicum course, students participate in a weekly seminar and work on a project under the supervision of their professors. This course will give students the opportunity to work with Georgetown Law’s O’Neill Institute (http://www.law.georgetown.edu/oneillinstitute/index.cfm) and its external partners in government and civil society to gain experience in using law to prevent non-communicable diseases (NCDs), including cancer, cardiovascular diseases, chronic respiratory diseases and diabetes. Law is a key tool to reduce the prevalence of key NCD risk factors: tobacco and alcohol use, physical inactivity, and unhealthy diets. Students will participate in a two-hour/week seminar and carry out 10 hours/week of project work under the direction of the course professors.

SEMINAR: In the seminar, students will explore the challenges and opportunities of using law to address risk factors that contribute to the rising prevalence of NCDs. The course will take a global approach grounded in international law, including international human rights law - the right to health, and World Health Organization (WHO) law and policy instruments, such as the Framework Convention on Tobacco Control and the Global Action Plan for the Prevention and Control of NCDs 2013-2020. Further, case studies will explore a variety of best practice examples from jurisdictions spanning the United Kingdom, South Africa, and Latin American countries, including taxes to discourage consumption of unhealthy products, laws restricting advertising and promotion, and laws and policies to promote physically active lifestyles.

Students will be equipped with an understanding of specific issues, such as the role of law compared with policy, the strengths and weaknesses of different regulatory strategies and the role and responsibilities of the relevant industries in promoting the right to health. After exploring a series of foundational themes and issues through the first half of the semester, the remainder of the class will focus on in-depth case studies and experiences in regulating the risk factors (e.g., industry litigation challenging NCD-related laws, challenges in monitoring and evaluating the health impacts of NCD-related laws, and civil society’s role in NCD law-making). Students will also learn how to use epidemiological data to craft compelling arguments in support of adoption of NCD-related laws and policies and to defend these laws when challenged by industry. Class time will be devoted to developing practical advocacy and drafting skills to support students in their project work.

PROJECT WORK: Students will work with external partners of the O’Neill Institute for National and Global Health on legal and policy projects related to NCDs, law and human rights. For example, students may draft alternative reports to UN bodies analyzing compliance with human rights obligations related to tobacco control and unhealthy diets (e.g., “shadow reports”). A report such as this could analyze the prevailing legal frameworks in a particular country and highlight any weaknesses in the statutory and regulatory language. By working with the O’Neill Institute, Inter-Governmental Organizations (e.g., the Pan American Health Organization, World Health Organization), and civil society organizations (e.g., Campaign for Tobacco Free Kids, Inter-American Heart Foundation), the course will give students the opportunity to use law to advocate for legal mechanisms to address critical health challenges.

Prerequisite: J.D. students must complete the required first-year program prior to enrolling in this course (part-time and interdivisional transfer students may enroll prior to completing Criminal Justice, Property, or their first-year elective).

Mutually Excluded Courses: Students may not concurrently enroll in this practicum course and a clinic or another practicum course. Students may concurrently enroll in this practicum course and an externship.

Note: This practicum course is open to LL.M. students, space permitting. Interested LL.M. students should email Louis Fine (fine@law.georgetown.edu) to request admission.

Evening students who work during the day are encouraged to reach out to the professor to discuss whether this practicum course would be compatible with their schedules.

This is a four-credit course. Two credits will be awarded for the two-hour weekly seminar and two credits will be awarded for approximately 10 hours of supervised project work per week, for a minimum of 11 weeks. Both the seminar and the project work will be graded.
This course will use the legal challenges to the Affordable Care Act as an initial case study of how political disputes play out in litigation, from the trial courts to the Supreme Court. We will follow the pattern of political litigation into the Trump Administration, and assess the lessons from the partisan shift in the litigation. We will consider how and why partisan litigation has proliferated. Increasingly, the losers in legislative battles have leapt immediately into the judicial arena, challenging the constitutionality of the laws enacted over their objection. Increasingly, politicians and the entities they control have deployed litigation as another tool in the partisan arsenal. And increasingly, federal courts have become the arbiters of political disputes between the other two branches of the Federal Government, and between the Federal Government and the States.

This course will address:

1. the theoretical perspectives underlying the international law of investment and the concept of police powers (particularly the power to protect health);
2. the different types of legal tests used to determine the regulatory legitimacy of a measure as well as the impact of varying standards of proof on analysis of this type;
3. rules governing expropriation of property rights and the circumstances in which health and environmental health measures might be considered equivalent to expropriation;
4. rules governing fair and equitable treatment as used in determining the legitimacy of domestic regulatory measures;
5. rules governing non-discrimination and their potential impact on domestic health measures that are non-discriminatory in form;
6. rules governing arbitrary or discriminatory measures; and
7. procedural issues relating to the participation of civil society in the negotiation of investment treaties and the settlement of investment disputes.

The course will address:

**Prerequisite:** Constitutional Law I: The Federal System (or Democracy and Coercion).

**Recommended:** Prior or concurrent enrollment in Federal Courts and the Federal System.
Increasing globalization compounds the complexity in preparing for and responding to public health emergencies. Identifying the numerous stakeholders, legal and policy considerations, and authorities in the midst of responding to a potential or declared public health emergency is a difficult endeavor. Fragmented or siloed preparedness activities and efforts before the public health emergency or after the response to the emergency or event inevitably result in duplicative or even confounding efforts, initiatives, authorities, or mandates. In spite of this, governments, international organizations, and non-government organizations continue to combat waning prioritization and urgency to initiate, maintain, and enable preparedness and response activities, capabilities, and functionality. Now more than ever, coordinated integration and implementation of national and international law and policy is critical to ensuring and enabling effective operationalization of stakeholders and resources globally during a response as lives hang in the balance.

Through the analysis of case studies from various global events (including infectious diseases, chemical and radiological events, and natural disasters), this course aims to establish a better understanding of the various global and national legal and policy fora, considerations, and influences and how they have (or have not) been applied in preparedness and response various efforts.

This course will focus on four core issues: quarantine and criminal penalties, access to affordable medicines and intellectual property, the international and constitutional right to health, and how political institutions and race intersect and condition the effect of law on population health. By looking at these issues in both the United States and abroad, particularly in low- and middle-income countries, students will gain a comparative perspective about how law works in practice.

Utilizing HIV and tuberculosis as core case studies, students will explore how law and policy tools can have a significant impact on population health. Today, state actors continue to use law to address public health issues—with examples of both great successes and significant failures.

As public health law embodies both thorny ethical issues and empirical questions about the power of the state to affect health, examining the intersection of law and social science will reveal substantive issues in key health policy issues as well as the conditions under which law and legal advocacy affects wellbeing. HIV and TB, the world’s biggest infectious killers, provide a lens through which to better understand central issues in public health law and policy, which will then be linked to other health concerns in infectious, non-communicable, and environmental health.

This one-week class will be a blend of lectures, discussion, and small group work. Readings will include case law and legal theory as well as texts from political science, economics, and public health. At the end, students will have a better understanding of how and when the law matters for health—which will be of particular interest to students interested in litigation, lawmaking, and public health regulation.

Note: WEEK ONE COURSE. This seminar will meet for one week only on the following days: Monday, January 7, 2019 through Friday, January 11, 2019, 9:00 a.m. - 11:35 a.m.

This course is mandatory pass/fail and will not count toward the 7 credit pass/fail limit for J.D. students. Attendance at all class sessions is mandatory and all enrolled students must attend the first class in order to remain enrolled. Students on the wait list must attend the first class in order to be admitted off the wait list. Enrolled students will have until the beginning of the second class session to request a drop by contacting the Office of the Registrar. Once the second class session begins, students may only seek a withdrawal by contacting their academic advisor in the Office of Graduate Programs. Withdrawals are permitted up until the last class for this specific course.
This course is intended to provoke thought and legal and ethical debate over pressing public policy issues surrounding the major health problems facing America and globally—e.g., infectious diseases, smoking, obesity, violence, injuries, and the environment. First, the course will examine the Constitutional and historical foundations of public health law in the United States. This section will discuss the Constitutional and statutory powers and duties of government to assure the conditions for a healthy and safe population, including the implementation of the Affordable Care Act. Second, the course will examine the conflicts between public health and civil liberties. For example, the course will probe conflicts between: (1) injury and disease surveillance and privacy; (2) labeling and advertising restrictions and freedom of expression; (3) personal control measures (e.g., screening, forced medical treatment and quarantine) and liberty; (4) commercial public health regulation and property rights. Finally, the course will examine the future of public health law. This “Future” includes a careful analysis of biosecurity—both naturally occurring (e.g., Ebola, pandemic influenza, Zika Virus) and intentional (e.g., bioterrorism such as smallpox and the anthrax attacks after September 11, 2001). This course should be important for all students considering careers in health law as well as those simply interested in exploring and debating the state of public health in America. It is a particularly unique opportunity for students given the expertise of Georgetown Law’s own O’Neill Institute for National and Global Health Law.

This course will proceed in three movements. First, it will consider the conceptual foundations of public health law in the United States through the lens of governance, ethics, and human rights, affording particular attention to the statutory and regulatory powers and duties of federal and state governmental entities to protect the health and safety of the population. Second, it will examine the sources of tension between the objectives of public health and civil liberties including, among others: labeling and advertising restrictions and free speech; disease surveillance and privacy; and considerations in personal versus population-based conceptions of health. Lastly, this course will conclude by considering emerging public health issues at the intersection of law and ethics, such as biosecurity, vaccine policy, and the role for public health in broader discussions around health care reform in the United States.

In a fieldwork practicum course, students participate in a weekly seminar and conduct related fieldwork at an outside organization. This practicum course will focus on regulation of food and personal-care products by the Food and Drug Administration (FDA) under the Food, Drug, and Cosmetic Act. Students will participate in a two-hour/week seminar and also undertake 10 hours/week of fieldwork at the Environmental Working Group (EWG), a public-interest nonprofit that advocates on behalf of consumers of both food and personal-care products, and other non-governmental organizations.

SEMINAR: The FDA’s authority to regulate the safety and nutritional aspects of food products dates back to 1906, and has evolved as food production has become increasingly industrialized and affected by an array of new technologies. On the other hand, FDA’s authority over the safety of personal-care products has remained rudimentary, and presents unique challenges in the face of both long- and short-term risks linked to their use. This seminar will utilize legislative and administrative materials as well as case law to enable students to become acquainted with the processes by which the federal government regulates food and personal-care products, will compare FDA’s regulatory authorities with regulatory schemes for consumer products, and will critique both the statutory framework and the performance of FDA in carrying out its duties. The course will also touch on related topics such as the role of the Federal Trade Commission in the regulation of trade practices related to food and cosmetics, the roles of the Environmental Protection Agency and the Consumer Product Safety Commission in the regulation of chemicals in consumer products, and the interaction of federal and state regulation.

FIELDWORK: In the fieldwork component of this course, students will be assigned to projects at the Environmental Working Group or a similar non-governmental organization. They will have an opportunity to learn how such institutions play a role in representing consumer interests in product safety issues being debated in both the administrative and legislative processes, and in matters subject to litigation.

Prerequisite: J.D. students must complete the required first-year program prior to enrolling in this course (part-time and interdivisional transfer students may enroll prior to completing Criminal Justice, Property, or their first-year elective).

Recommended: Administrative Law, as well as food and drug law-related courses, are recommended but not required.

Mutually Excluded Courses: Students may not concurrently enroll in this practicum and an externship, a clinic, or another practicum.

Note: This practicum course is open to LL.M. students, space permitting. Interested LL.M. students should email Louis Fine (fine@law.georgetown.edu) to request admission.

This course is suitable for evening students who can commit to attending class and working 10 hours/week (during business hours) on site at their field placements. This is a four credit course. Two credits will be awarded for the two-hour weekly seminar and two credits for approximately 10 hours of fieldwork per week, for a minimum of 11 weeks, to be scheduled with the faculty. The fieldwork must be completed during normal business hours. The two credit seminar portion of this practicum will be graded. The two credits of fieldwork are mandatory pass/fail. Students will be allowed to take another course pass/fail in the same semester as the field work. Students who enroll in this course will be automatically enrolled in both the seminar and fieldwork components and may not take either component separately. After Add/Drop, a student who wishes to withdraw from a practicum course must obtain permission from the faculty member and the Assistant Dean for Experiential Education.
LAW 3067 v00 Public-Private Partnerships: Law and Governance
(http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%203067%20v00)

LL.M Seminar (cross-listed) | 2 credit hours
Public-Private Partnerships (PPPs) are a modern means for pursuing social and policy outcomes. When they work, they mobilize the comparative advantages of the public and private spheres to address key challenges. When they fail, critics cite their worst aspects and argue for a retreat into more traditional roles. Their prevalence means that everyone is likely to come across, and be affected by, such a partnership at some point in their careers.

Bridging theory and practice, students will have the opportunity to understand why PPPs have emerged and the theoretical basis under which they operate. They will also receive practical tools and knowledge to allow them to work with or within a PPP and discuss and debate sophisticatedly how they should be managed and governed.

To do this, the course uses a mix of lecture, discussion, video, and experiential learning. It will start by examining and contrasting the key organizational, legal, and governance components of public-sector institutions and corporate entities. Students will then analyze several kinds of PPPs including global health partnerships, innovative finance institutions, and biomedical research consortiums. In particular, they will examine applicable international, corporation, and regulatory law and seek to understand the choices these partnerships make in applying the formation of a PPP using the tools they develop during the semester.

LAW 1071 v00 Reproductive Health and International Human Rights Law
(http://curriculum.law.georgetown.edu/course-search/?keyword=LAW%201071%20v00) (Project-Based Practicum)
J.D. Practicum | 4 credit hours
In a project-based practicum course, students participate in a weekly seminar and work on a project under the supervision of their professor. This project-based practicum course will focus on the interaction between international human rights law and reproductive health and rights. Students will participate in a two hour/week seminar and carry out 10 hours/week of project work under the direction of the course professor.

SEMINAR: The seminar will begin by providing an overview of international human rights law as it pertains to reproductive rights. The course will then focus on access to reproductive health from an international perspective, examining States’ obligations on a variety of issues, such as maternal mortality and coerced sterilization. Analyzing recent decisions emerging from regional and international human rights bodies, such as the European Court of Human Rights, the Inter-American Commission and Court on Human Rights and the CEDAW Committee (UN Committee on the Convention on the Elimination of All Forms of Discrimination against Women), the seminar component will provide a solid legal foundation for students to develop their experiential/field placement projects.

PROJECT WORK: Students will work with external partners on legal and policy projects related to reproductive health. Some of the projects may include drafting amicus briefs for cases currently pending before international bodies, and drafting briefs assessing a particular State’s compliance with human rights law regarding sexual and reproductive rights to be filed in front of UN bodies (shadow reports). Through these projects, students will learn how to conduct an analysis of existing legal and regulatory frameworks for sexual and reproductive health from a human rights perspective. Students will also learn how to use epidemiological data to support and craft compelling human rights law arguments for advancing public policy on, for example, maternal mortality and sexual violence prevention and eradication. By working with external civil society organizations, the course will give students the opportunity to develop practical projects using international human rights law to advocate for the advancement of reproductive health rights. In the past, external partners have included organizations working on women's rights issues, such as: the Center for Reproductive Rights, Women’s Link Worldwide, Human Rights Watch (Women’s Rights Division), IPAS, and Southern Africa Litigation Centre, among others.

Prerequisite: J.D. students must complete the required first-year program prior to enrolling in this course (part-time and interdivisional transfer students may enroll prior to completing Criminal Justice, Property, or their first-year elective).

Mutually Excluded Courses: Students may not concurrently enroll in this practicum course and a clinic or another practicum course. Students may concurrently enroll in this practicum course and an externship.

Note: This practicum course is open to LL.M. students, space permitting. Interested LL.M. students should email Louis Fine (fine@law.georgetown.edu) to request admission. Evening students who work during the day are encouraged to reach out to the professor to discuss whether this practicum course would be compatible with their schedules.

This is a four credit course. Two credits will be awarded for the two-hour weekly seminar and two credits will be awarded for approximately 10 hours of supervised project work per week, for a minimum of 11 weeks. Both the seminar and the project work will be graded.

Students who enroll in this course will be automatically enrolled in both the seminar and project components and may not take either component separately. After Add/Drop, a student who wishes to withdraw from a
LAW 837 v00 Research with Human Subjects: Law, Policy & Ethics
This course provides an opportunity to explore issues related to the use of humans as research subjects. The course will begin by examining the history of human subject research, including the abuses that led to the creation of our modern legal protections. Following a discussion of the ethical principles of research involving humans, we will carefully explore the legal treatment of human subjects, focusing on US regulations and international instruments. Core sources will include: the Common Rule (45 CFR 46), Belmont Report, Declaration of Helsinki, Nuremberg Code, and CIOMS. After exploring a series of foundational themes and issues through the first half of the semester, the remainder of the class will focus on in-depth case studies. Topics may include: international research, research involving vulnerable populations (children, prisoners, and pregnant women), informed consent, research on subjects with impaired decision-making abilities, genetic/genomic research, risk-benefit analysis, coercion/undue inducement, use of placebos, and IRB governance.

This seminar provides opportunities for participants to engage in a research and writing project related to humans as research subjects. Participants will conduct independent research and scholarly writing on important problems at the intersection of law, policy and ethics.

Note: This seminar requires a paper. J.D. students must register for the 3 credit section of the seminar if they wish to write a paper fulfilling the Upperclass Legal Writing Requirement for JD students. The paper requirements of the 2 credit section will not fulfill the Upperclass Legal Writing Requirement for JD students.

LAW 3017 v00 Survey of Employee Benefits Law
This course provides a general overview of the federal labor and tax law policies and principles that apply to employment-based retirement plans, health care plans, and compensation arrangements. Students will examine the principles of the Internal Revenue Code and ERISA that govern the form and operation of employer-sponsored plans, especially tax-qualified pension plans, executive compensation arrangements and welfare plans (with a focus on health care). This class will concentrate on the structure and basic rules that govern such arrangements. In the context of tax-qualified and nonqualified plans and health and welfare programs, this course will explore plan administration, fiduciary duties, claims appeals and litigation.

Prerequisite: Federal Income Taxation (formerly Taxation I).

Mutually Excluded Courses:
- Employee Benefits: Qualified Retirement Plans
- Employee Benefits: Executive Compensation
- Employee Benefits: Health & Welfare Plans

Note: Students enrolled in the Employee Benefits Certificate Program may not register for this course.

LAW 2080 v00 The Affordable Care Act: Law and Policy Governing Private Health Insurance
The Patient Protection and Affordable Care Act has significantly impacted the health care system. Through changes to the regulation of private health insurance, federal subsidies for low-income people, individual and employer mandates, and expansion of the Medicaid program, millions of Americans have gained health insurance. Yet, implementation of the law's most significant reforms has been challenging, and its provisions have been subject to extensive litigation and continued threats of repeal.

This course will examine the regulation of private health insurance with an emphasis on the issues that the Affordable Care Act was designed to address. The course will explore regulatory changes such as new market reforms and health insurance marketplaces; the impact of federal and state approaches to implementation; and legal and regulatory challenges. Guest lectures by speakers—including insurance industry representatives, legal experts, and regulators—will provide students with an in-depth understanding of how the Affordable Care Act has been implemented and what it means for millions of consumers across the country.

The primary objective of the course is to teach students about the regulation of private health insurance at the federal and state level, with the rare opportunity to witness real-time changes to this complex system. Through this course, students will gain practical experience in identifying and analyzing federal and state laws, regulations, and administrative materials. Students will also gain a broader understanding of administrative law and how it has affected Affordable Care Act implementation. By the end of the course, students will be able to describe how private health insurance is regulated at the federal and state level, the major reforms ushered in by the Affordable Care Act, federal and state implementation, and likely areas of changes to the law given real-time debates.

Note: This is a required course for the U.S. Health Law Certificate.

LAW 496 v01 The Mind and the Law
Developments in neuroscience and the psychological study of cognition and emotion are transforming our understanding of the mind. These developments have large implications for law and lawyering. They challenge some of civil and criminal law's central premises - about people's rationality, free choice, and consistency over time. This course will examine some of these challenges, and it will weigh the law's possible responses. Legal topics to be explored will include health and safety regulation, the idea of the reasonable person, intent and culpability, mental disability, and the roles of revenge, regret, and other motives in civil and criminal justice. Other themes, relevant to lawyering and to the management of conflict, will include the psychology of negotiation, the nature of intuition and judgment, and the roles of trust and social norms. The course will also consider the causes and control of violence and extremism, as well as the use of emerging neuroscience technologies for legal and national security purposes.
WHO is the sole global health organization and, as such, has been traditionally considered the lead actor in this domain at the center and the forefront of technical, policy and normative developments in global health governance. Its Constitution and institutional structure represent both the embodiment of a new way of thinking about global health and its governance at the time of creation of the UN system as well as the outcome of political compromises that have affected the life of the organization and its positioning in the global health landscape. The historical development of WHO’s functions and activities are also an emblematic reflection of the changing role of health in the global economic, political and development agendas and, as such, can shed light on broader issue of international law and relations. WHO’s role and significance as the central actor in global health governance has been the object of contestation and much critical reflection starting in the 1990s and culminating with the organization’s role in responding to the Ebola crisis. At the same time, global health governance has become more complex, fragmented and politicized and the future role of WHO has to be seen in the context of those developments and the search for a coherent global health architecture.

The course will focus on four broad areas covering the most important aspects of WHO as an international organization, an actor in global health governance, and a forum for policy and legal developments: 1) Historical, constitutional and institutional aspects; 2) normative functions and the role of WHO in the development of international law; 3) directing and coordinating functions, both in terms of how to address the most important health challenges as well as with regard to WHO’s interaction with a number of political and economic regimes; 4) the position of WHO in the current global health landscape, both with regard to the role of other actors as well as to how the organization should look to its own future. The course will allow students to gain not only an in-depth knowledge and appreciation of WHO from an institutional and structural perspective, but also most importantly to appreciate the main contemporary challenges in global health as well as the role of health in a number of critical policy and normative regimes. The approach of the course will be based on lecturing, class discussions and at least one class exercise. Prof. Burci’s former tenure as the Legal Counsel of WHO and a long-term senior lawyer in the organization will allow him to contribute real-life examples and experiences that will better root the course in the realities of the life of an international organization.

This course is mandatory pass/fail and will not count toward the 7 credit pass/fail limit for J.D. students.