

CAMILA LOSADA STRASSLE

Department of Bioethics, National Institutes of Health Clinical Center • 10 Center Drive, Suite 1C118 • Bethesda, MD 20892-1156 •
camila.strassle@nih.gov • camilas@alumni.stanford.edu • (626) 710-6173

EMPLOYMENT:

2018-20 Predoctoral Fellow, Department of Bioethics, National Institutes of Health (NIH)
Research topics: the legal issues around workplace wellness programs (with Benjamin Berkman), a framework for guiding patient involvement in conducting clinical research (with Steven Pearson), the ethical use of patient contracts in clinical care (with Benjamin Berkman), the moral (in)significance of genetic parenthood (with David Wasserman), a working group on direct-to-consumer genetic testing (with Benjamin Berkman and Leila Jamal)
Participation and training: Bioethics Consultation Service, Clinical Center Ethics Committee, Institutional Review Board meetings, clinical rounds, Course on Ethical and Regulatory Aspects of Clinical Research, Bioethics seminars

EDUCATION:

2018 Stanford University, 3.816 GPA
B.A., Human Biology, with interdisciplinary honors in Ethics in Society
Minors in Philosophy and Spanish
Study Abroad in Madrid and Buenos Aires

JOURNAL PUBLICATIONS:

1. **Camila Strassle** and Benjamin E. Berkman. (2019). “Workplace Wellness Programs: Empirical Doubt, Legal Ambiguity, and Conceptual Confusion.” *William & Mary Law Review*, Forthcoming.
2. **Camila Strassle**. (2019). “Fair Subject Selection in Cystic Fibrosis Trials.” *Journal of Cystic Fibrosis*, 18(5), e47.
3. Gregory Sawicki, Joshua Ostrenga, Kristofer Petren, Aliza Fink, Emma D’Agostino, **Camila Strassle**, Michael S. Schechter, and Margaret Rosenfeld. (2018). “Risk Factors for Gaps in Care during Transfer from Pediatric to Adult Cystic Fibrosis Programs in the United States.” *Annals of the American Thoracic Society*, 15(2), 234-240.

Under Review: “A Framework for Patient Engagement Throughout the Broader Research Enterprise” with Steven Pearson

REFEREED CONFERENCE PRESENTATIONS:

1. **Camila Strassle** and Steven Pearson. (2019). “A Framework for Patient Engagement Throughout the Research Life Cycle.” Paper presented at the American Society for Bioethics and Humanities 21st Annual Conference, Pittsburgh, PA, October 24-27.
2. **Camila Strassle** and Benjamin Berkman. (2019). “Workplace Wellness Programs: Empirical Doubt, Legal Ambiguity, and Conceptual Confusion.” Paper presented at the American Society for Bioethics and Humanities 21st Annual Conference, Pittsburgh, PA, October 24-27.
3. **Camila Strassle**, Breck Gamel, David Hansen, Meghann Barloewen, Maria Bellefeuille, Annette Campbell, Emma D’Agostino, Laurie Eddie, Jessica Franklin, Preston Hinkle, Kat Porco, Margaret Smith, Heather Walter, Victoria Danner, and Aliza Fink. (2017). “Advancing the Research that Matters Most to CF Patients and Families: Preliminary Findings from the Insight CF Registry Research Project.” Poster presented at the 31st Annual North American Cystic Fibrosis Conference, Indianapolis, IN, November 2-4 and abstract published in *Pediatric Pulmonology*, 52(47), 424.
4. **Camila Strassle**, Diana Naranjo, Rosa Girón, Colleen Dunn, Luis Máiz-Carro, and Carlos Milla. (2017). “Social Support and Its Implications for Health in Patients with Cystic Fibrosis: Results of an International Study Across Two Cultures.” Poster presented at the 31st Annual North American Cystic Fibrosis Conference, Indianapolis, IN, November 2-4 and abstract published in *Pediatric Pulmonology*, 52(47), 483.

CAMPUS TALKS AND MISCELLANEA:

1. **Camila Strassle**. (2019). “Bioethics and Genetics.” Presentation to high school students at the National Human Genome Research Institute, Bethesda, MD, May 3.
2. Ben Schwan, Dominic Mangino, and **Camila Strassle**. (2019). “Introduction to Medical Ethics.” Presentation to Chaplain Interns at the National Institutes of Health Clinical Center, Bethesda, MD, January 15 (repeated July 24).

3. **Camila Strassle.** (2018). “Contemporary Controversies in Workplace Wellness and Disability.” Paper presented at the annual Undergraduate Ethics Symposium at DePauw University, Greencastle, IN, April 19-21.
4. **Camila Strassle.** (2018). “How Workplace Wellness Programs Harm People with Disabilities.” [Justice Everywhere blog post].
5. **Camila Strassle.** (2017). “Wisconsin v. Yoder and the Libertarian View on Publicly Funded Education.” *A Priori: The Brown Journal of Philosophy*, 2:165-176.

AWARDS, GRANTS, AND HONORS:

2018	Lyle and Olive Cook Prize for Outstanding Honors Thesis (~500 USD), Stanford’s Center for Ethics in Society
2018	Cardinal Service Notation for Commitment to Public Service, Stanford’s Haas Center for Public Service
2018	Comunidad Recognition for Academic Excellence, Stanford’s El Centro Chicano y Latino
2017	Public Interest Law Fellowship (~5000 USD), Stanford’s Haas Center for Public Service
2016	Prize for Best Paper in Justice Course, Stanford Department of Political Science
2016	Major Grant for Original Research (~7500 USD), Stanford Undergraduate Advising and Research
2013	Boothe Prize for Excellence in First Year Writing Nominee, Stanford Program in Structured Liberal Education

EXPERIENCE:

2017-18	Research Assistant , Stanford Law School Co-wrote a proposal with Professor Rabia Belt to promote disability scholarship and access as part of Stanford’s Long-Range Planning initiative. Assisted research on the litigation history of post-traumatic stress disorder and psychological trauma.
2017	Public Interest Law Fellow , Esperanza Immigrant Rights Project Led legal orientations in Spanish at the Los Angeles Immigration Court and Adelanto Detention Center. Performed “intakes” of minors and adults in immigration removal proceedings to answer questions in a confidential environment, assess eligibility for legal relief, and make referrals to ancillary services. Researched country conditions for asylum applicants detained at Adelanto Detention Center. Assisted direct representation program for adults deemed mentally incompetent and unable to represent themselves.
2015-17	Founding Member , Cystic Fibrosis Patient and Family Research Advisory Committee Collaborated with eight co-founders to launch “Insight CF,” the Cystic Fibrosis Foundation’s first patient-driven research initiative. Provided training, solicited, and reviewed 441 research questions from cystic fibrosis patients and relatives. Engaged 1795 CF community members in prioritizing 22 final research questions, representing the Cystic Fibrosis Foundation’s most successful engagement effort. Findings illustrate that patients are interested in research involvement beyond the traditional role of research subject.
2015-17	Principal Investigator (PI) , Stanford Medical School Initiated and conducted an international cross-sectional study on social support in cystic fibrosis. Completed Human Subjects training through Stanford’s Research Compliance Office, received approval through Stanford’s Institutional Review Board (IRB) and Madrid’s hospital ethics committees. Worked full-time in summer of 2016 to consent and enroll 110 patients and analyze data. Presented work at the North American Cystic Fibrosis Conference in coordination with 5 cystic fibrosis researchers.
2013	Intern , Cystic Fibrosis Foundation’s Public Policy Department Organized Teen Advocacy Day attended by 55 teenagers from 23 states to meet with more than 200 members of Congress. Compiled evidence of variability in healthcare coverage offered to CF patients across state boundaries. Successfully lobbied for the passage of the Ensuring Access to Clinical Trials Act.

LEADERSHIP AND SERVICE:

2016-17	Bridge Peer Counselor to Stanford students in mental health crises
2015-17	Latinos Unidos Academic Committee Chair
2016	EPASA Tutor to middle school students in East Palo Alto
2016	SLE Writing Tutor to Stanford freshmen
2016	Latino Community Orientation Volunteer
2014-15	Hope House Scholars Tutor to women in rehabilitation for drug abuse

ADDITIONAL INFORMATION: Native Spanish speaker (intermediate proficiency), QPR Certified for Suicide Prevention, CITI trained for work with human subjects