

Works Cited

“‘Elimination’ of Down Syndrome Is a ‘Great Hate Crime,’ Says Holy See Conference at the U.N.” *America Magazine*, 5 Apr. 2018,
www.americamagazine.org/politics-society/2018/03/21/elimination-down-syndrome-great-hate-crime-says-holy-see-conference-un.

Caplan, Arthur L. “Chloe’s Law: A Powerful Legislative Movement Challenging a Core Ethical Norm of Genetic Testing.” *PLOS Biology*, vol. 13, no. 8, 2015,
doi:10.1371/journal.pbio.1002219.

Chapman, Robin S. “Language Development in Children and Adolescents with Down Syndrome.” *The Handbook of Child Language*, 2019, pp. 641–663.,
doi:10.1111/b.9780631203124.1996.00027.x.

Constantine, MI, et al. “Imperfect Informed Consent for Prenatal Screening: Lessons from the Quad Screen.” *Clinical Ethics*, vol. 9, no. 1, 2013, pp. 17–27.,
doi:10.1177/1477750913511339.

Crombag, Neeltje Mth, et al. “Explaining Variation in Down’s Syndrome Screening Uptake: Comparing the Netherlands with England and Denmark Using Documentary Analysis and Expert Stakeholder Interviews.” *BMC Health Services Research*, vol. 14, no. 1, 2014, doi:10.1186/1472-6963-14-437.

Deutsche Welle. “Germany to Make Down Syndrome Blood Test Free under ‘Strict’ Conditions: DW: 19.09.2019.” *DW.COM*,

www.dw.com/en/germany-to-make-down-syndrome-blood-test-free-under-strict-condition

“Eugenics, Genetics and the Family. Volume I, Scientific Papers of the Second International Congress of Eugenics.” *JAMA: The Journal of the American Medical Association*, vol. 80, no. 20, 1923, p. 1480., doi:10.1001/jama.1923.02640470058043.

Garland-Thomson, Rosemarie. “Building a World with Disability in It.” *Culture - Theory - Disability*, 2017, doi:10.14361/9783839425336-006.

Garrard, Eve, and Stephen Wilkinson. “Selecting Disability and the Welfare of the Child.” *Monist*, vol. 89, no. 4, 2006, pp. 482–504., doi:10.5840/monist20068944.

Graaf, Gert De, et al. “Estimates of the Live Births, Natural Losses, and Elective Terminations with Down Syndrome in the United States.” *American Journal of Medical Genetics Part A*, vol. 167, no. 4, 2015, pp. 756–767., doi:10.1002/ajmg.a.37001.

Greely, Henry T. *The End of Sex and the Future of Human Reproduction*. Harvard University Press, 2016.

“Growth Charts for Children With Down Syndrome in the United States.” *Pediatrics*, 2015, doi:10.1542/peds.2015-1652d.

Guedj, Fayçal, et al. “Prenatal Treatment of Down Syndrome.” *Current Opinion in Obstetrics and Gynecology*, vol. 26, no. 2, 2014, pp. 92–103., doi:10.1097/gco.0000000000000056.

Hagard, S, and F A Carter. “Preventing the Birth of Infants with Down's Syndrome: a

Cost-Benefit Analysis.” *Bmj*, vol. 1, no. 6012, 1976, pp. 753–756.,
doi:10.1136/bmj.1.6012.753.

Hall, Sue, et al. “Undergoing Prenatal Screening for Down's Syndrome: Presentation of Choice and Information in Europe and Asia.” *European Journal of Human Genetics*, vol. 15, no. 5, 2007, pp. 563–569., doi:10.1038/sj.ejhg.5201790.

Iltis, Ana S. “Prenatal Screening and Prenatal Diagnosis: Contemporary Practices in Light of the Past.” *Journal of Medical Ethics*, vol. 42, no. 6, 2016, pp. 334–339.,
doi:10.1136/medethics-2016-103623.

Irving, Claire A, and Milind P Chaudhari. “Cardiovascular Abnormalities in Down's Syndrome: Spectrum, Management and Survival over 22 Years.” *Archives of Disease in Childhood*, vol. 97, no. 4, 2011, pp. 326–330., doi:10.1136/adc.2010.210534.

Jensen, K. M., and M. M. Davis. “Health Care in Adults with Down Syndrome: a Longitudinal Cohort Study.” *Journal of Intellectual Disability Research*, vol. 57, no. 10, 2012, pp. 947–958., doi:10.1111/j.1365-2788.2012.01589.x.

Jj, Dominguez-Cruz, and Bueno Delgado. “Dermatological Manifestations of Down Syndrome.” *Genetics and Etiology of Down Syndrome*, 2011, doi:10.5772/19321.

Jong, Antina De, and Guido M.w.r. De Wert. “Prenatal Screening: An Ethical Agenda for the Near Future.” *Bioethics*, vol. 29, no. 1, 2014, pp. 46–55., doi:10.1111/bioe.12122.

Krahn, T. “P46 ‘Seriousness’ and the Regulation of Preimplantation Genetic Diagnosis: the Case of Down's Syndrome.” *Reproductive BioMedicine Online*, vol. 20, 2010,

doi:10.1016/s1472-6483(10)62362-0.

Lawson, Karen L. “Expectations of the Parenting Experience and Willingness to Consider Selective Termination for Down Syndrome.” *Journal of Reproductive and Infant Psychology*, vol. 24, no. 1, 2006, pp. 43–59., doi:10.1080/02646830500475351.

Lewis, Celine, et al. “Women’s Experiences and Preferences for Service Delivery of Non-Invasive Prenatal Testing for Aneuploidy in a Public Health Setting: A Mixed Methods Study.” *Plos One*, vol. 11, no. 4, 2016, doi:10.1371/journal.pone.0153147.

Livermore, Roy. “Ups and Downs.” *Oxford Scholarship Online*, 2018, doi:10.1093/oso/9780198717867.003.0011.

Mahy, J., et al. “Identifying Facilitators and Barriers to Physical Activity for Adults with Down Syndrome.” *Journal of Intellectual Disability Research*, vol. 54, no. 9, 2010, pp. 795–805., doi:10.1111/j.1365-2788.2010.01308.x.

Minor, Jessica. “Revised Model of Informed Consent.” *Informed Consent in Predictive Genetic Testing*, 2015, pp. 109–163., doi:10.1007/978-3-319-17416-7_4.

Mégarbané, André, et al. “The 50th Anniversary of the Discovery of Trisomy 21: The Past, Present, and Future of Research and Treatment of Down Syndrome.” *Genetics in Medicine*, vol. 11, no. 9, 2009, pp. 611–616., doi:10.1097/gim.0b013e3181b2e34c.

Noble, Ray, et al. “Pandora's Box: Ethics of PGD for Inherited Risk of Late-Onset Disorders.” *Reproductive BioMedicine Online*, vol. 17, 2008, pp. 55–60., doi:10.1016/s1472-6483(10)60332-x.

Patterson, David, and Alberto C. S. Costa. “Down Syndrome and Genetics — a Case of

Linked Histories.” *Nature Reviews Genetics*, vol. 6, no. 2, 2005, pp. 137–147.,
doi:10.1038/nrg1525.

Petersen, T S. “Just Diagnosis? Preimplantation Genetic Diagnosis and Injustices to Disabled People.” *Journal of Medical Ethics*, vol. 31, no. 4, 2005, pp. 231–234.,

Reinders, Johannes Sjoerd. *The Future of the Disabled in Liberal Society: an Ethical Analysis*. University of Notre Dame Press, 2009.

Reynolds, T M. “Down's Syndrome Screening Is Unethical: Views of Today's Research Ethics Committees.” *Journal of Clinical Pathology*, vol. 56, no. 4, 2003, pp. 268–270.,
doi:10.1136/jcp.56.4.268.

Rubeis, Giovanni, and Florian Steger. “A Burden from Birth? Non-Invasive Prenatal Testing and the Stigmatization of People with Disabilities.” *Bioethics*, vol. 33, no. 1, 2018, pp. 91–97., doi:10.1111/bioe.12518.

Scott, R. “Choosing between Possible Lives: Legal and Ethical Issues in Preimplantation Genetic Diagnosis.” *Oxford Journal of Legal Studies*, vol. 26, no. 1, 2006, pp. 153–178., doi:10.1093/ojls/gqi048.

Scully, Jackie Leach. *Disability Bioethics: Moral Bodies, Moral Difference*. Rowman & Littlefield Publishers, 2008.

Smith-Bindman, Rebecca, et al. “Prenatal Screening for Down Syndrome in England and Wales and Population-Based Birth Outcomes.” *American Journal of Obstetrics and Gynecology*, vol. 189, no. 4, 2003, pp. 980–985., doi:10.1067/s0002-9378(03)00721-x.

Taneja, Alankrita, et al. "Abortion of Fetus with Down's Syndrome: India Joins the Worldwide Controversy Surrounding Abortion Laws." *Science and Engineering Ethics*, 2017, doi:10.1007/s11948-017-9926-y.

Underwood, E. "Can Down Syndrome Be Treated?" *Science*, vol. 343, no. 6174, 2014, pp. 964–967., doi:10.1126/science.343.6174.964.

Wayne, Deborah O'mullan, and Sheama Krishnagiri. "Parents' Leisure: the Impact of Raising a Child with down Syndrome." *Occupational Therapy International*, vol. 12, no. 3, 2005, pp. 180–194., doi:10.1002/oti.4.