Carla G Van El

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8234892/publications.pdf

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32 papers	1,676 citations	430874 18 h-index	434195 31 g-index
33 all docs	33 docs citations	33 times ranked	2360 citing authors

#	Article	IF	Citations
1	The use of polygenic risk scores in pre-implantation genetic testing: an unproven, unethical practice. European Journal of Human Genetics, 2022, 30, 493-495.	2.8	38
2	Opportunistic genomic screening. Recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2021, 29, 365-377.	2.8	76
3	ESHG PPPC Comments on postmortem use of genetic data for research purposes. European Journal of Human Genetics, 2020, 28, 144-146.	2.8	3
4	The use of PROMs and shared decisionâ€making in medical encounters with patients: An opportunity to deliver valueâ€based health care to patients. Journal of Evaluation in Clinical Practice, 2020, 26, 524-540.	1.8	82
5	Web-based return of BRCA2 research results: one-year genetic counselling experience in Iceland. European Journal of Human Genetics, 2020, 28, 1656-1661.	2.8	12
6	Systematic scoping review of the concept of â€˜genetic identityâ€™ and its relevance for germline modification. PLoS ONE, 2020, 15, e0228263.	2.5	8
7	Parental perspectives on retention and secondary use of neonatal dried bloodspots: a Dutch mixed methods study. BMC Pediatrics, 2019, 19, 230.	1.7	4
8	European recommendations integrating genetic testing into multidisciplinary management of sudden cardiac death. European Journal of Human Genetics, 2019, 27, 1763-1773.	2.8	78
9	Reply to Bombard and Mighton. European Journal of Human Genetics, 2019, 27, 507-508.	2.8	0
10	How Should Decision Aids Be Used During Counseling to Help Patients Who Are â€œGenetically at Riskâ€?. AMA Journal of Ethics, 2019, 21, E865-872.	0.7	2
11	Value-based genomic screening: exploring genomic screening for chronic diseases using triple value principles. BMC Health Services Research, 2019, 19, 823.	2.2	4
12	How to Integrate Personalized Medicine into Prevention? Recommendations from the Personalized Prevention of Chronic Diseases (PRECeDI) Consortium. Public Health Genomics, 2019, 22, 208-214.	1.0	21
13	Recontacting patients in clinical genetics services: recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2019, 27, 169-182.	2.8	65
14	Recontacting or not recontacting? A survey of current practices in clinical genetics centres in Europe. European Journal of Human Genetics, 2018, 26, 946-954.	2.8	33
15	Human germline gene editing. Recommendations of ESHG and ESHREâ€â€¡. Human Reproduction Open, 2018, 2018, hox025.	5.4	3
16	Human germline gene editing: Recommendations of ESHG and ESHRE. European Journal of Human Genetics, 2018, 26, 445-449.	2.8	30
17	Responsible innovation in human germline gene editing: Background document to the recommendations of ESHG and ESHRE. European Journal of Human Genetics, 2018, 26, 450-470.	2.8	39
18	One small edit for humans, one giant edit for humankind? Points and questions to consider for a responsible way forward for gene editing in humans. European Journal of Human Genetics, 2018, 26, 1-11.	2.8	55

#	Article	IF	Citations
19	Responsible innovation in human germline gene editing. Background document to the recommendations of ESHG and ESHREâ€â€¡. Human Reproduction Open, 2018, 2018, hox024.	5.4	9
20	Risk and the politics of boundary work: preserving autonomous midwifery in the Netherlands. Health, Risk and Society, 2018, 20, 379-407.	1.7	6
21	A response to the forensic genetics policy initiativeâ€™s report â€œEstablishing Best Practice for Forensic DNA Databasesâ€• Forensic Science International: Genetics, 2018, 36, e19-e21.	3.1	11
22	Barriers and Facilitating Factors for Implementation of Genetic Services: A Public Health Perspective. Frontiers in Public Health, 2017, 5, 195.	2.7	19
23	Implementing non-invasive prenatal testing for aneuploidy in a national healthcare system: global challenges and national solutions. BMC Health Services Research, 2017, 17, 670.	2.2	55
24	Responsible implementation of expanded carrier screening. European Journal of Human Genetics, 2016, 24, e1-e12.	2.8	240
25	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. Summary and recommendations. European Journal of Human Genetics, 2015, , .	2.8	13
26	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. European Journal of Human Genetics, 2015, 23, 1438-1450.	2.8	260
27	Public attitudes towards preventive genomics and personal interest in genetic testing to prevent disease: a survey study. European Journal of Public Health, 2014, 24, 768-775.	0.3	51
28	Neuroimaging in the Courtroom: Normative Frameworks and Consensual Practices. AJOB Neuroscience, 2014, 5, 37-39.	1.1	5
29	Newborn screening for pompe disease? a qualitative study exploring professional views. BMC Pediatrics, 2014, 14, 203.	1.7	11
30	Whole-genome sequencing in health care. European Journal of Human Genetics, 2013, 21, 580-584.	2.8	330
31	Whole-genome sequencing in health care. Recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2013, 21 Suppl 1, S1-5.	2.8	66
32	Genetic testing and common disorders in a public health framework. European Journal of Human Genetics, 2011, 19, 377-381.	2.8	46