Carla G Van El

List of Publications by Year
in descending order

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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	Whole-genome sequencing in health care. European Journal of Human Genetics, 2013, 21, 580-584.	2.8	330
2	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
3	Responsible implementation of expanded carrier screening. European Journal of Human Genetics, 2016, 24, e1-e12.	2.8	240
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	European recommendations integrating genetic testing into multidisciplinary management of sudden cardiac death. European Journal of Human Genetics, 2019, 27, 1763-1773.	2.8	78
6	Opportunistic genomic screening. Recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2021, 29, 365-377.	2.8	76
7	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
8	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
9	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
10	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
11	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
12	Genetic testing and common disorders in a public health framework. European Journal of Human Genetics, 2011, 19, 377-381.	2.8	46
13	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
14	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
15	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
16	Human germline gene editing: Recommendations of ESHG and ESHRE. European Journal of Human Genetics, 2018, 26, 445-449.	2.8	30
17	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
18	Barriers and Facilitating Factors for Implementation of Genetic Services: A Public Health Perspective. Frontiers in Public Health, 2017, 5, 195.	2.7	19

#	Article	IF	Citations
19	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
20	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
21	Newborn screening for pompe disease? a qualitative study exploring professional views. BMC Pediatrics, 2014, 14, 203.	1.7	11
22	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
23	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
24	Systematic scoping review of the concept of â€˜genetic identityâ€™ and its relevance for germline modification. PLoS ONE, 2020, 15, e0228263.	2.5	8
25	Risk and the politics of boundary work: preserving autonomous midwifery in the Netherlands. Health, Risk and Society, 2018, 20, 379-407.	1.7	6
26	Neuroimaging in the Courtroom: Normative Frameworks and Consensual Practices. AJOB Neuroscience, 2014, 5, 37-39.	1.1	5
27	Parental perspectives on retention and secondary use of neonatal dried bloodspots: a Dutch mixed methods study. BMC Pediatrics, 2019, 19, 230.	1.7	4
28	Value-based genomic screening: exploring genomic screening for chronic diseases using triple value principles. BMC Health Services Research, 2019, 19, 823.	2.2	4
29	Human germline gene editing. Recommendations of ESHG and ESHREâ€â€¡. Human Reproduction Open, 2018, 2018, hox025.	5.4	3
30	ESHG PPPC Comments on postmortem use of genetic data for research purposes. European Journal of Human Genetics, 2020, 28, 144-146.	2.8	3
31	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
32	Reply to Bombard and Mighton. European Journal of Human Genetics, 2019, 27, 507-508.	2.8	0