Martina C Cornel

List of Publications by Year
in descending order

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220 papers	7,112 citations	61977 43 h-index	88628 70 g-index
246 all docs	246 docs citations	246 times ranked	7280 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	Teratogenic Effects of Antiepileptic Drugs: Use of an International Database on Malformations and Drug Exposure (MADRE). Epilepsia, 2000, 41, 1436-1443.	5.1	186
5	Genetic testing in asymptomatic minors. European Journal of Human Genetics, 2009, 17, 711-719.	2.8	167
6	2020 APHRS/HRS expert consensus statement on the investigation of decedents with sudden unexplained death and patients with sudden cardiac arrest, and of their families. Heart Rhythm, 2021, 18, e1-e50.	0.7	151
7	The interface between assisted reproductive technologies and genetics: technical, social, ethical and legal issues. European Journal of Human Genetics, 2006, 14, 588-645.	2.8	137
8	Preconceptional genetic carrier testing and the commercial offer directly-to-consumers. Human Reproduction, 2011, 26, 972-977.	0.9	124
9	Current issues in medically assisted reproduction and genetics in Europe: research, clinical practice, ethics, legal issues and policy. European Journal of Human Genetics, 2013, 21, S1-S21.	2.8	120
10	Population Screening for Genetic Disorders in the 21st Century: Evidence, Economics, and Ethics. Public Health Genomics, 2010, 13, 106-115.	1.0	114
11	Genetic educational needs and the role of genetics in primary care: a focus group study with multiple perspectives. BMC Family Practice, 2011, 12, 5.	2.9	106
12	Public attitudes towards genetic testing revisited: comparing opinions between 2002 and 2010. European Journal of Human Genetics, 2013, 21, 793-799.	2.8	103
13	Where are you going, where have you been: a recent history of the direct-to-consumer genetic testing market. Journal of Community Genetics, 2010, 1, 101-106.	1.2	97
14	Accuracy of family history of cancer: clinical genetic implications. European Journal of Human Genetics, 2000, 8, 181-186.	2.8	91
15	Newborn screening programmes in Europe; arguments and efforts regarding harmonization. Part 2 â€“ From screening laboratory results to treatment, followâ€up and quality assurance. Journal of Inherited Metabolic Disease, 2012, 35, 613-625.	3.6	88
16	Whole-genome sequencing in newborn screening? A statement on the continued importance of targeted approaches in newborn screening programmes. European Journal of Human Genetics, 2015, 23, 1593-1600.	2.8	87
17	Towards a European consensus for reporting incidental findings during clinical NGS testing. European Journal of Human Genetics, 2015, 23, 1601-1606.	2.8	85
18	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

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19	Genomic newborn screening: public health policy considerations and recommendations. BMC Medical Genomics, 2017, 10, 9.	1.5	78
20	European recommendations integrating genetic testing into multidisciplinary management of sudden cardiac death. European Journal of Human Genetics, 2019, 27, 1763-1773.	2.8	78
21	Recent developments in genetics and medically assisted reproduction: from research to clinical applications. European Journal of Human Genetics, 2018, 26, 12-33.	2.8	76
22	Opportunistic genomic screening. Recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2021, 29, 365-377.	2.8	76
23	Improvement of drug exposure data in a registration of congenital anomalies. Pilot-study: Pharmacist and mother as sources for drug exposure data during pregnancy. Teratology, 1999, 60, 33-36.	1.6	75
24	Genetic testing and common disorders in a public health framework: how to assess relevance and possibilities. European Journal of Human Genetics, 2011, 19, S6-S44.	2.8	75
25	Time trends in neural tube defects prevalence in relation to preventive strategies: an international study. Journal of Epidemiology and Community Health, 1999, 53, 630-635.	3.7	68
26	Heterogeneity of neural tube defects in europe: The significance of site of defect and presence of other major anomalies in relation to geographic differences in prevalence. Teratology, 1991, 44, 547-559.	1.6	66
27	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
28	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
29	Developing a policy for paediatric biobanks: principles for good practice. European Journal of Human Genetics, 2013, 21, 2-7.	2.8	63
30	Comparison of national policies on periconceptional use of folic acid to prevent spina bifida and anencephaly (SBA). , 1997, 55, 134-137.		60
31	International perspectives on the implementation of reproductive carrier screening. Prenatal Diagnosis, 2020, 40, 301-310.	2.3	60
32	Deficient knowledge of genetics relevant for daily practice among medical students nearing graduation. Genetics in Medicine, 2005, 7, 295-301.	2.4	58
33	Current issues in medically assisted reproduction and genetics in Europe: research, clinical practice, ethics, legal issues and policy. Human Reproduction, 2014, 29, 1603-1609.	0.9	57
34	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
35	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
36	Limb defects associated with major congenital anomalies: Clinical and epidemiological study from the International Clearinghouse for Birth Defects Monitoring Systems. American Journal of Medical Genetics Part A, 2000, 93, 110-116.	2.4	51

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37	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
38	Genetic horoscopes: is it all in the genes? Points for regulatory control of direct-to-consumer genetic testing. European Journal of Human Genetics, 2009, 17, 857-859.	2.8	50
39	Prioritization of future genetics education for general practitioners: a Delphi study. Genetics in Medicine, 2012, 14, 323-329.	2.4	49
40	A framework to start the debate on neonatal screening policies in the EU: an Expert Opinion Document. European Journal of Human Genetics, 2014, 22, 12-17.	2.8	49
41	Folic acidâ€"the scientific debate as a base for public health policy. Reproductive Toxicology, 2005, 20, 411-415.	2.9	47
42	Genetic testing and common disorders in a public health framework. European Journal of Human Genetics, 2011, 19, 377-381.	2.8	46
43	Three-month follow-up of Western and non-Western participants in a study on preconceptional ancestry-based carrier couple screening for cystic fibrosis and hemoglobinopathies in the Netherlands. Genetics in Medicine, 2008, 10, 820-830.	2.4	45
44	Reflecting on Earlier Experiences with Unsolicited Findings: Points to Consider for Nextâ€Generation Sequencing and Informed Consent in Diagnostics. Human Mutation, 2013, 34, 1322-1328.	2.5	45
45	The challenges of the expanded availability of genomic information: an agenda-setting paper. Journal of Community Genetics, 2018, 9, 103-116.	1.2	45
46	The Dutch 'Folic Acid Campaign'-have the goals been achieved?. Paediatric and Perinatal Epidemiology, 2000, 14, 111-117.	1.7	44
47	Neonatal Screening for Treatable and Untreatable Disorders: Prospective Parentsâ€™ Opinions. Pediatrics, 2010, 125, e99-e106.	2.1	44
48	What Do Parents of Children with Down Syndrome Think about Nonâ€Invasive Prenatal Testing (NIPT)?. Journal of Genetic Counseling, 2017, 26, 522-531.	1.6	43
49	Preconceptional ancestry-based carrier couple screening for cystic fibrosis and haemoglobinopathies: what determines the intention to participate or not and actual participation?. European Journal of Human Genetics, 2009, 17, 999-1009.	2.8	42
50	Genome-wide sequencing in acutely ill infants: genomic medicineâ€™s critical application?. Genetics in Medicine, 2019, 21, 498-504.	2.4	42
51	Sustained effects of online genetics education: a randomized controlled trial on oncogenetics. European Journal of Human Genetics, 2014, 22, 310-316.	2.8	40
52	Common Elements in Rare Kidney Diseases: Conclusions from a Kidney Disease: Improving Global Outcomes (KDIGO) Controversies Conference. Kidney International, 2017, 92, 796-808.	5.2	40
53	Three Years after the Dutch Folic Acid Campaign: Growing Socioeconomic Differences. Preventive Medicine, 2002, 35, 65-69.	3.4	39
54	Italian appeal court: a genetic predisposition to commit murder?. European Journal of Human Genetics, 2010, 18, 519-521.	2.8	39

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55	The Dutch national summit on preconception care: a summary of definitions, evidence and recommendations. Journal of Community Genetics, 2015, 6, 107-115.	1.2	39
56	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
57	Periconceptional folic acid intake in the northern Netherlands. Lancet, The, 1999, 353, 1187.	13.7	38
58	Implementation of Pharmacogenetics in Primary Care: A Multi-Stakeholder Perspective. Frontiers in Genetics, 2020, 11, 10.	2.3	38
59	Stakeholder perspectives on the implementation of genetic carrier screening in a changing landscape. BMC Health Services Research, 2017, 17, 146.	2.2	36
60	Systematic Review of N-of-1 Studies in Rare Genetic Neurodevelopmental Disorders. Neurology, 2021, 96, 529-540.	1.1	36
61	Informed consent for exome sequencing in diagnostics: exploring first experiences and views of professionals and patients. Clinical Genetics, 2014, 85, 417-422.	2.0	35
62	Preconceptional Cystic Fibrosis Carrier Screening: Attitudes and Intentions of the Target Population. Genetic Testing and Molecular Biomarkers, 2004, 8, 80-89.	1.7	34
63	Consanguineous marriage and reproductive risk: attitudes and understanding of ethnic groups practising consanguinity in Western society. European Journal of Human Genetics, 2014, 22, 452-457.	2.8	34
64	Implementation of preconceptional carrier screening for cystic fibrosis and haemoglobinopathies: A sociotechnical analysis. Health Policy, 2007, 83, 277-286.	3.0	33
65	Recreational genomics? Dreams and fears on genetic susceptibility screening. European Journal of Human Genetics, 2008, 16, 403-404.	2.8	33
66	Developing a framework for implementation of genetic services: learning from examples of testing for monogenic forms of common diseases. Journal of Community Genetics, 2014, 5, 337-347.	1.2	33
67	Effectiveness of oncogenetics training on general practitionersâ€™ consultation skills: a randomized controlled trial. Genetics in Medicine, 2014, 16, 45-52.	2.4	32
68	Inability to detect plasma etretinate and acitretin is a poor predictor of the absence of these teratogens in tissue after stopping acitretin treatment.. British Journal of Clinical Pharmacology, 1994, 38, 229-235.	2.4	31
69	Factors for successful implementation of population-based expanded carrier screening: learning from existing initiatives: Table 1. European Journal of Public Health, 2016, 27, ckw110.	0.3	31
70	DPD Testing Before Treatment With Fluoropyrimidines in the Amsterdam UMCs: An Evaluation of Current Pharmacogenetic Practice. Frontiers in Pharmacology, 2019, 10, 1609.	3.5	31
71	Down syndrome: effects of demographic factors and prenatal diagnosis on the future livebirth prevalence. Human Genetics, 1993, 92, 163-168.	3.8	30
72	Possibilities and barriers in the implementation of a preconceptional screening programme for cystic fibrosis carriers: a focus group study. Public Health, 2003, 117, 396-403.	2.9	30

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73	Attitudes of Potential Providers Towards Preconceptional Cystic Fibrosis Carrier Screening. Journal of Genetic Counseling, 2004, 13, 31-44.	1.6	30
74	Human germline gene editing: Recommendations of ESHG and ESHRE. European Journal of Human Genetics, 2018, 26, 445-449.	2.8	30
75	Neonatal and carrier screening for rare diseases: how innovation challenges screening criteria worldwide. Journal of Community Genetics, 2021, 12, 257-265.	1.2	30
76	Effect of Comprehensive Oncogenetics Training Interventions for General Practitioners, Evaluated at Multiple Performance Levels. PLoS ONE, 2015, 10, e0122648.	2.5	29
77	Key Implications of Data Sharing in Pediatric Genomics. JAMA Pediatrics, 2018, 172, 476.	6.2	29
78	The ethics of clinical applications of germline genome modification: a systematic review of reasons. Human Reproduction, 2018, 33, 1777-1796.	0.9	29
79	Association between holoprosencephaly and exposure to topical retinoids: results of the EUROCAT survey. Paediatric and Perinatal Epidemiology, 1991, 5, 445-447.	1.7	28
80	Influence of educational level on determinants of folic acid use. Paediatric and Perinatal Epidemiology, 2003, 17, 256-263.	1.7	28
81	Changing to NIPT as a first-tier screening test and future perspectives: opinions of health professionals. Prenatal Diagnosis, 2015, 35, 1316-1323.	2.3	28
82	Increasing awareness of and behaviour towards periconceptional folic acid consumption in The Netherlands from 1994 to 1995. European Journal of Clinical Pharmacology, 1998, 54, 329-331.	1.9	27
83	Review of the Reported Measures of Clinical Validity and Clinical Utility as Arguments for the Implementation of Pharmacogenetic Testing: A Case Study of Statin-Induced Muscle Toxicity. Frontiers in Pharmacology, 2017, 8, 555.	3.5	27
84	Community genetics. Its definition 2010. Journal of Community Genetics, 2010, 1, 19-22.	1.2	26
85	Points to consider for prioritizing clinical genetic testing services: a European consensus process oriented at accountability for reasonableness. European Journal of Human Genetics, 2015, 23, 729-735.	2.8	26
86	Ephedrine treatment for autoimmune myasthenia gravis. Neuromuscular Disorders, 2017, 27, 259-265.	0.6	26
87	Some epidemiological data on oral clefts in the northern Netherlands, 1981â€“1988. Journal of Cranio-Maxillo-Facial Surgery, 1992, 20, 147-152.	1.7	25
88	The challenge of implementing genetic tests with clinical utility while avoiding unsound applications. Journal of Community Genetics, 2014, 5, 7-12.	1.2	25
89	Maternal Plasma DNA and RNA Sequencing for Prenatal Testing. Advances in Clinical Chemistry, 2016, 74, 63-102.	3.7	25
90	Uptake of fetal aneuploidy screening after the introduction of the nonâ€invasive prenatal test: A national populationâ€based register study. Acta Obstetricia Et Gynecologica Scandinavica, 2021, 100, 1265-1272.	2.8	25

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91	Current practice and future interest of GPs and prospective parents in pre-conception care in The Netherlands. Family Practice, 2004, 21, 307-309.	1.9	24
92	Lower limb deficient children in the Netherlands. Prosthetics and Orthotics International, 2000, 24, 13-18.	1.0	23
93	Factors associated with not using folic acid supplements preconceptionally. Public Health Nutrition, 2014, 17, 2344-2350.	2.2	23
94	A Demographic Approach to the Assessment of Down Syndrome Screening Performance. , 1997, 17, 717-724.		22
95	Periconceptional folic acid in The Netherlands in 1995. Socioeconomic differences. Journal of Epidemiology and Community Health, 1998, 52, 826-827.	3.7	22
96	Genetic screening and democracy: lessons from debating genetic screening criteria in the Netherlands. Journal of Community Genetics, 2012, 3, 79-89.	1.2	22
97	Is There an Association between Maternal Carbamazepine Use during Pregnancy and Eye Malformations in the Child?. Epilepsia, 2002, 43, 929-931.	5.1	21
98	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
99	Raising awareness of carrier testing for hereditary haemoglobinopathies in high-risk ethnic groups in the Netherlands: a pilot study among the general public and primary care providers. BMC Public Health, 2009, 9, 338.	2.9	20
100	NEONATAL SCREENING FOR CYSTIC FIBROSIS. Lancet, The, 1986, 327, 802-803.	13.7	19
101	Consanguinity sans reproche. Human Genetics, 1991, 86, 295-296.	3.8	19
102	The expansion of newborn screening: is reproductive benefit an appropriate pursuit?. Nature Reviews Genetics, 2009, 10, 666-667.	16.3	19
103	Public support for neonatal screening for Pompe disease, a broad-phenotype condition. Orphanet Journal of Rare Diseases, 2012, 7, 15.	2.7	19
104	A decade of molecular genetic testing for MODY: a retrospective study of utilization in The Netherlands. European Journal of Human Genetics, 2015, 23, 29-33.	2.8	19
105	Barriers and Facilitating Factors for Implementation of Genetic Services: A Public Health Perspective. Frontiers in Public Health, 2017, 5, 195.	2.7	19
106	Folate prophylaxis in pregnancy. Lancet, The, 1995, 346, 1227-1228.	13.7	18
107	CFTR Mutations in Turkish and North African Cystic Fibrosis Patients in Europe: Implications for Screening. Genetic Testing and Molecular Biomarkers, 2008, 12, 25-35.	1.7	18
108	Developing and optimizing a decisional instrument using self-reported ancestry for carrier screening in a multi-ethnic society. Genetics in Medicine, 2006, 8, 502-509.	2.4	17

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109	Newborn Screening Programmes in Europe, Arguments and Efforts Regarding Harmonisation: Focus on Organic Acidurias. JIMD Reports, 2016, 32, 105-115.	1.5	17
110	The Gen-Equip Project: evaluation and impact of genetics e-learning resources for primary care in six European languages. Genetics in Medicine, 2019, 21, 718-726.	2.4	17
111	2020 APHRS/HRS expert consensus statement on the investigation of decedents with sudden unexplained death and patients with sudden cardiac arrest, and of their families. Journal of Arrhythmia, 2021, 37, 481-534.	1.2	17
112	Ensuring best practice in genomics education and evaluation: reporting item standards for education and its evaluation in genomics (RISE2 Genomics). Genetics in Medicine, 2021, 23, 1356-1365.	2.4	17
113	Population-based birth-defect and risk-factor surveillance: data from the Northern Netherlands. International Journal of Risk and Safety in Medicine, 1996, 8, 197-209.	0.6	16
114	How Should Preconceptional Cystic Fibrosis Carrier Screening Be Provided? Opinions of Potential Providers and the Target Population. Public Health Genomics, 2003, 6, 157-165.	1.0	16
115	Translational Research in Genomics of Alzheimer's Disease: A Review of Current Practice and Future Perspectives. Journal of Alzheimer's Disease, 2010, 20, 967-980.	2.6	16
116	Autosomal recessive disease in children of consanguineous parents: inferences from the proportion of compound heterozygotes. Journal of Community Genetics, 2010, 1, 37-40.	1.2	16
117	Lay perceptions of predictive testing for diabetes based on DNA test results versus family history assessment: a focus group study. BMC Public Health, 2011, 11, 535.	2.9	16
118	Current and Best Practices of Genetic Testing for Maturity Onset Diabetes of the Young: Views of Professional Experts. Public Health Genomics, 2015, 18, 52-59.	1.0	16
119	Implementing genetic education in primary care: the Gen-Equip programme. Journal of Community Genetics, 2017, 8, 147-150.	1.2	16
120	Policy Making in Newborn Screening Needs a Structured and Transparent Approach. Frontiers in Public Health, 2017, 5, 53.	2.7	16
121	Ethical and Social Issues in Pharmacogenomics Testing. Current Pharmaceutical Design, 2010, 16, 245-252.	1.9	15
122	Do consanguineous parents of a child affected by an autosomal recessive disease have more DNA identical-by-descent than similarly-related parents with healthy offspring? Design of a case-control study. BMC Medical Genetics, 2010, 11, 113.	2.1	15
123	Family communication as strategy in diabetes prevention: An observational study in families with Dutch and Surinamese South-Asian ancestry. Patient Education and Counseling, 2012, 87, 23-29.	2.2	15
124	Stakeholder Views on Active Cascade Screening for Familial Hypercholesterolemia. Healthcare (Switzerland), 2018, 6, 108.	2.0	15
125	TypeÂ2 diabetes and inheritance: what information do diabetes organizations provide on the Internet?. Diabetic Medicine, 2006, 23, 1233-1238.	2.3	14
126	Effects of a simple educational intervention in well-baby clinics on womenâ€™s knowledge about and intake of folic acid supplements in the periconceptional period: a controlled trial. Public Health Nutrition, 2015, 18, 1119-1126.	2.2	14

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127	Does nonâ€invasive prenatal testing affect the livebirth prevalence of Down syndrome in the Netherlands? A populationâ€based register study. Prenatal Diagnosis, 2021, 41, 1351-1359.	2.3	14
128	Comparison of couples referred and not referred for genetic counseling in a genetic clinic after the birth of a child with a congenital anomaly: A study in a population in the northeastern Netherlands. American Journal of Medical Genetics Part A, 1992, 42, 387-392.	2.4	13
129	The need for interaction between assisted reproduction technology and genetics. European Journal of Human Genetics, 2006, 14, 509-511.	2.8	13
130	Governing biological material at the intersection of care and research: the use of dried blood spots for biobanking. Croatian Medical Journal, 2012, 53, 390-397.	0.7	13
131	Severely impaired health status at diagnosis of Pompe disease: A cross-sectional analysis to explore the potential utility of neonatal screening. Molecular Genetics and Metabolism, 2012, 107, 448-455.	1.1	13
132	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
133	A genetic diagnosis of maturityâ€onset diabetes of the young (<scp>MODY</scp>): experiences of patients and family members. Diabetic Medicine, 2015, 32, 1385-1392.	2.3	13
134	Mainstreaming informed consent for genomic sequencing: A call for action. European Journal of Cancer, 2021, 148, 405-410.	2.8	13
135	Couplesâ€™ experiences with expanded carrier screening: evaluation of a university hospital screening offer. European Journal of Human Genetics, 2021, 29, 1252-1258.	2.8	13
136	Expanding Neonatal Bloodspot Screening: A Multi-Stakeholder Perspective. Frontiers in Pediatrics, 2021, 9, 706394.	1.9	13
137	Prevalence of congenital heart disease in patients with phenylketonuria. Journal of Pediatrics, 1991, 119, 282-283.	1.8	12
138	Acardius acephalus after induced ovulation: A case report. Teratology, 1993, 47, 257-262.	1.6	12
139	Preconception Cystic Fibrosis Carrier Screening: Costs and Consequences. Genetic Testing and Molecular Biomarkers, 2005, 9, 158-166.	1.7	12
140	Validation of self-reported folic acid use in a multiethnic population: results of the Amsterdam Born Children and their Development study. Public Health Nutrition, 2011, 14, 2022-2028.	2.2	12
141	A case study of haemoglobinopathy screening in the Netherlands: witnessing the past, lessons for the future. Ethnicity and Health, 2012, 17, 217-239.	2.5	12
142	'A morass of considerations': exploring attitudes towards ethnicity-based haemoglobinopathy-carrier screening in primary care. Family Practice, 2013, 30, 604-610.	1.9	12
143	Call for Prudence in Whole-Genome Testing. Science, 2013, 341, 958-959.	12.6	12
144	Illness representations of type 2 diabetes patients are associated with perceptions of diabetes threat in relatives. Journal of Health Psychology, 2014, 19, 358-368.	2.3	12

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145	Neonatal diagnosis of <scp>D</scp>own syndrome in the <scp>N</scp>etherlands: suspicion and communication with parents. Journal of Intellectual Disability Research, 2014, 58, 953-961.	2.0	12
146	Screening for Familial Hypercholesterolemia in Children: What Can We Learn From Adult Screening Programs?. Healthcare (Switzerland), 2015, 3, 1018-1030.	2.0	12
147	Genomics for all in the 21st century?. Journal of Community Genetics, 2017, 8, 249-251.	1.2	12
148	Association between ovulation stimulation, in vitro fertilisation, and neural tube defects?. Teratology, 1990, 42, 201-203.	1.6	11
149	Newborn screening for pompe disease? a qualitative study exploring professional views. BMC Pediatrics, 2014, 14, 203.	1.7	11
150	First steps in exploring prospective exome sequencing of consanguineous couples. European Journal of Medical Genetics, 2014, 57, 613-616.	1.3	11
151	Ephedrine as add-on therapy for patients with myasthenia gravis: protocol for a series of randomised, placebo-controlled n-of-1 trials. BMJ Open, 2015, 5, e007863.	1.9	11
152	Recent developments in genetics and medically-assisted reproduction: from research to clinical applicationsâ€â€¡. Human Reproduction Open, 2017, 2017, hox015.	5.4	11
153	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
154	Evidence-Based Genetic Education of Non-Genetic-Expert Physicians: Experiences Over Three Decades in Amsterdam. Frontiers in Genetics, 2019, 10, 712.	2.3	11
155	Moving towards a cure in genetics: what is needed to bring somatic gene therapy to the clinic?. European Journal of Human Genetics, 2019, 27, 484-487.	2.8	11
156	Consanguinity and Endogamy in the Netherlands: Demographic and Medical Genetic Aspects. Human Heredity, 2014, 77, 161-166.	0.8	10
157	Older mothers and increased impact of prenatal screening: stable livebirth prevalence of trisomy 21 in the Netherlands for the period 2000â€“2013. European Journal of Human Genetics, 2018, 26, 157-165.	2.8	10
158	Changes in opinions about human germline gene editing as a result of the Dutch DNA-dialogue project. European Journal of Human Genetics, 2023, 31, 409-416.	2.8	10
159	Variation in prenatal cytogenetic diagnosis: Policies in 13 european countries, 1989â€“1991. Prenatal Diagnosis, 1994, 14, 337-344.	2.3	9
160	Renal defects and limb deficiencies in 197 infants: Is it possible to define the â€œacrorenal syndromeâ€?. , 2004, 129A, 149-155.		9
161	Connective tissue: Cancer patientsâ€™ attitudes towards medical research using excised (tumour) tissue. BioSocieties, 2011, 6, 466-486.	1.3	9
162	Attitudes of general practitioners and midwives towards ethnicity-based haemoglobinopathy-carrier screening. European Journal of Human Genetics, 2012, 20, 1112-1117.	2.8	9

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163	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
164	Moving somatic gene editing to the clinic: routes to market access and reimbursement in Europe. European Journal of Human Genetics, 2021, 29, 1477-1484.	2.8	9
165	Ad hoc tracing of a cohort of patients exposed to acitretine (NeotigasonÂ®) on a nation-wide scale. European Journal of Clinical Pharmacology, 1992, 42, 555-557.	1.9	8
166	Additional information from parental questionnaires and pharmacy records for registration of birth defects. EuroMAP-group. European Journal of Epidemiology, 2000, 16, 329-336.	5.7	8
167	Wealth and health in relation to birth defects mortality. Journal of Epidemiology and Community Health, 2000, 54, 644-644.	3.7	8
168	On the symmetry of limb deficiencies among childrenwith multiple congenital anomalies. Annales De GÃ©nÃ©tique, 2001, 44, 19-24.	0.4	8
169	The potential of the European network of congenital anomaly registers (EUROCAT) for drug safety surveillance: a descriptive study. Pharmacoepidemiology and Drug Safety, 2006, 15, 675-682.	1.9	8
170	Patientsâ€™ intentions to inform relatives about TypeÂ2 diabetes risk: the role of worry in the process of family risk disclosure. Diabetic Medicine, 2012, 29, e461-7.	2.3	8
171	What do people want to know about NIPT? Content analysis of questions emailed to national NIPT information websites. Prenatal Diagnosis, 2017, 37, 412-415.	2.3	8
172	Aggregated N-of-1 trials for unlicensed medicines for small populations: an assessment of a trial with ephedrine for myasthenia gravis. Orphanet Journal of Rare Diseases, 2017, 12, 88.	2.7	8
173	Contentious ethical issues in community genetics: letâ€™s talk about them. Journal of Community Genetics, 2020, 11, 5-6.	1.2	8
174	Systematic scoping review of the concept of â€˜genetic identityâ€™ and its relevance for germline modification. PLoS ONE, 2020, 15, e0228263.	2.5	8
175	Ovulation-inducing drugs: a drug utilization and risk study in the Dutch population. International Journal of Risk and Safety in Medicine, 1992, 3, 99-111.	0.6	7
176	Communicating a drug alert. European Journal of Clinical Pharmacology, 1994, 47, 125-132.	1.9	7
177	Letter to the Editor: Folic acid prevents more than neural tube defects: A registry-based study in the northern Netherlands. European Journal of Epidemiology, 2002, 18, 279-280.	5.7	7
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