

# Lidewij Henneman

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

126  
papers

3,004  
citations

32  
h-index

50  
g-index

136  
ext. papers

3,718  
ext. citations

4  
avg, IF

4.98  
L-index

#	Paper	IF	Citations
126	Noninvasive Prenatal Test Results Indicative of Maternal Malignancies: A Nationwide Genetic and Clinical Follow-Up Study.. <i>Journal of Clinical Oncology</i> , <b>2022</b> , JCO2102260	2.2	2
125	Medical costs of children admitted to the neonatal intensive care unit: The role and possible economic impact of WES in early diagnosis.. <i>European Journal of Medical Genetics</i> , <b>2022</b> , 104467	2.6	0
124	Nationwide implementation of the non-invasive prenatal test: Evaluation of a blended learning program for counselors.. <i>PLoS ONE</i> , <b>2022</b> , 17, e0267865	3.7	1
123	Clinical impact of additional findings detected by genome-wide non-invasive prenatal testing: Follow-up results of the TRIDENT-2 study. <i>American Journal of Human Genetics</i> , <b>2022</b> , 109, 1140-1152	11	2
122	Economic evaluations of exome and genome sequencing in pediatric genetics: considerations towards a consensus strategy.. <i>Journal of Medical Economics</i> , <b>2021</b> , 24, 60-70	2.4	1
121	Expanding Neonatal Bloodspot Screening: A Multi-Stakeholder Perspective. <i>Frontiers in Pediatrics</i> , <b>2021</b> , 9, 706394	3.4	0
120	Preconception risk assessment for thalassaemia, sickle cell disease, cystic fibrosis and Tay-Sachs disease. <i>The Cochrane Library</i> , <b>2021</b> , 10, CD010849	5.2	1
119	Preconception expanded carrier screening: a focus group study with relatives of mucopolysaccharidosis type III patients and the general population. <i>Journal of Community Genetics</i> , <b>2021</b> , 12, 311-323	2.5	1
118	Clinical and community genetics services in the Dutch Caribbean. <i>Journal of Community Genetics</i> , <b>2021</b> , 12, 497-501	2.5	0
117	CouplesSexperiences with expanded carrier screening: evaluation of a university hospital screening offer. <i>European Journal of Human Genetics</i> , <b>2021</b> , 29, 1252-1258	5.3	0
116	How will new genetic technologies, such as gene editing, change reproductive decision-making? Views of high-risk couples. <i>European Journal of Human Genetics</i> , <b>2021</b> , 29, 39-50	5.3	1
115	Preconception expanded carrier screening: Impact of information presented by text or video on genetic knowledge and attitudes. <i>Journal of Genetic Counseling</i> , <b>2021</b> , 30, 457-469	2.5	5
114	Should germline genome editing be allowed? The effect of treatment characteristics on public acceptability. <i>Human Reproduction</i> , <b>2021</b> , 36, 465-478	5.7	3
113	In memoriam Prof. Dr. Leo P. ten Kate. <i>Journal of Community Genetics</i> , <b>2021</b> , 12, 1-3	2.5	
112	Neonatal and carrier screening for rare diseases: how innovation challenges screening criteria worldwide. <i>Journal of Community Genetics</i> , <b>2021</b> , 12, 257-265	2.5	8
111	ParentsSPerspectives and Societal Acceptance of Implementation of Newborn Screening for SCID in the Netherlands. <i>Journal of Clinical Immunology</i> , <b>2021</b> , 41, 99-108	5.7	8
110	Informing Parents about Newborn Screening: A European Comparison Study. <i>International Journal of Neonatal Screening</i> , <b>2021</b> , 7,	2.6	2

109	Uptake of fetal aneuploidy screening after the introduction of the non-invasive prenatal test: A national population-based register study. <i>Acta Obstetrica Et Gynecologica Scandinavica</i> , <b>2021</b> , 100, 1265-1272	3.8	8
108	Association between low fetal fraction in cell-free DNA testing and adverse pregnancy outcome: A systematic review. <i>Prenatal Diagnosis</i> , <b>2021</b> , 41, 1287-1295	3.2	2
107	The Emergence and Global Spread of Noninvasive Prenatal Testing. <i>Annual Review of Genomics and Human Genetics</i> , <b>2021</b> , 22, 309-338	9.7	6
106	Routinization of prenatal screening with the non-invasive prenatal test: pregnant women's perspectives. <i>European Journal of Human Genetics</i> , <b>2021</b> ,	5.3	1
105	Non-invasive prenatal test uptake in socioeconomically disadvantaged neighborhoods. <i>Prenatal Diagnosis</i> , <b>2021</b> , 41, 1395-1400	3.2	2
104	Low fetal fraction in cell-free DNA testing is associated with adverse pregnancy outcome: Analysis of a subcohort of the TRIDENT-2 study. <i>Prenatal Diagnosis</i> , <b>2021</b> , 41, 1296-1304	3.2	0
103	Who ever heard of 16p11.2 deletion syndrome? Parents' perspectives on a susceptibility copy number variation syndrome. <i>European Journal of Human Genetics</i> , <b>2020</b> , 28, 1196-1204	5.3	2
102	Attitudes of relatives of mucopolysaccharidosis type III patients toward preconception expanded carrier screening. <i>European Journal of Human Genetics</i> , <b>2020</b> , 28, 1331-1340	5.3	3
101	Do Preferred Risk Formats Lead to Better Understanding? A Multicenter Controlled Trial on Communicating Familial Breast Cancer Risks Using Different Risk Formats. <i>Patient Preference and Adherence</i> , <b>2020</b> , 14, 333-342	2.4	2
100	Benefit vs potential harm of genome-wide prenatal cfDNA testing requires further investigation and should not be dismissed based on current data. <i>Ultrasound in Obstetrics and Gynecology</i> , <b>2020</b> , 55, 695-696	5.8	2
99	Effect of education and attitude on health professionals' knowledge on prenatal screening. <i>European Journal of Midwifery</i> , <b>2020</b> , 4, 38	0.7	1
98	Genetic Health Care Before Conception <b>2020</b> , 35-52		1
97	International perspectives on the implementation of reproductive carrier screening. <i>Prenatal Diagnosis</i> , <b>2020</b> , 40, 301-310	3.2	29
96	Women's experiences of monitoring the small-for-gestational age fetus by ultrasound: A qualitative study. <i>PLoS ONE</i> , <b>2019</b> , 14, e0216052	3.7	0
95	Informing relatives at risk of inherited cardiac conditions: experiences and attitudes of healthcare professionals and counselees. <i>European Journal of Human Genetics</i> , <b>2019</b> , 27, 1341-1350	5.3	8
94	The Gen-Equip Project: evaluation and impact of genetics e-learning resources for primary care in six European languages. <i>Genetics in Medicine</i> , <b>2019</b> , 21, 718-726	8.1	11
93	TRIDENT-2: National Implementation of Genome-wide Non-invasive Prenatal Testing as a First-Tier Screening Test in the Netherlands. <i>American Journal of Human Genetics</i> , <b>2019</b> , 105, 1091-1101	11	96
92	Dilemma of Reporting Incidental Findings in Newborn Screening Programs for SCID: Parents' Perspective on Ataxia Telangiectasia. <i>Frontiers in Immunology</i> , <b>2019</b> , 10, 2438	8.4	12

91	Attitudes of the general population towards preconception expanded carrier screening for autosomal recessive disorders including inborn errors of metabolism. <i>Molecular Genetics and Metabolism</i> , <b>2019</b> , 126, 14-22	3.7	19
90	Direct-to-consumer carrier screening for cystic fibrosis via a hospital website: a 6-year evaluation. <i>Journal of Community Genetics</i> , <b>2019</b> , 10, 249-257	2.5	3
89	Origin and clinical relevance of chromosomal aberrations other than the common trisomies detected by genome-wide NIPS: results of the TRIDENT study. <i>Genetics in Medicine</i> , <b>2018</b> , 20, 480-485	8.1	54
88	Preconception carrier screening for multiple disorders: evaluation of a screening offer in a Dutch founder population. <i>European Journal of Human Genetics</i> , <b>2018</b> , 26, 166-175	5.3	10
87	Experiences of a High-Risk Population with Prenatal Hemoglobinopathy Carrier Screening in a Primary Care Setting: a Qualitative Study. <i>Journal of Genetic Counseling</i> , <b>2018</b> , 27, 635-646	2.5	2
86	Introduction of non-invasive prenatal testing as a first-tier aneuploidy screening test: A survey among Dutch midwives about their role as counsellors. <i>Midwifery</i> , <b>2018</b> , 56, 1-8	2.8	17
85	Recontacting in light of new genetic diagnostic techniques for patients with intellectual disability: Feasibility and parental perspectives. <i>European Journal of Medical Genetics</i> , <b>2018</b> , 61, 213-218	2.6	10
84	Cell-Free DNA-Based Noninvasive Prenatal Testing and Society <b>2018</b> , 235-249		0
83	Women's decision making regarding prenatal screening for fetal aneuploidy: A qualitative comparison between 2003 and 2016. <i>Midwifery</i> , <b>2018</b> , 64, 93-100	2.8	4
82	What do people want to know about NIPT? Content analysis of questions emailed to national NIPT information websites. <i>Prenatal Diagnosis</i> , <b>2017</b> , 37, 412-415	3.2	7
81	Stakeholder perspectives on the implementation of genetic carrier screening in a changing landscape. <i>BMC Health Services Research</i> , <b>2017</b> , 17, 146	2.9	18
80	With expanded carrier screening, founder populations run the risk of being overlooked. <i>Journal of Community Genetics</i> , <b>2017</b> , 8, 327-333	2.5	5
79	Implementing non-invasive prenatal testing for aneuploidy in a national healthcare system: global challenges and national solutions. <i>BMC Health Services Research</i> , <b>2017</b> , 17, 670	2.9	38
78	Women's Experience with Non-Invasive Prenatal Testing and Emotional Well-being and Satisfaction after Test-Results. <i>Journal of Genetic Counseling</i> , <b>2017</b> , 26, 1348-1356	2.5	26
77	Uptake of prenatal diagnostic testing for retinoblastoma compared to other hereditary cancer syndromes in the Netherlands. <i>Familial Cancer</i> , <b>2017</b> , 16, 271-277	3	9
76	What Do Parents of Children with Down Syndrome Think about Non-Invasive Prenatal Testing (NIPT)?. <i>Journal of Genetic Counseling</i> , <b>2017</b> , 26, 522-531	2.5	37
75	Factors for successful implementation of population-based expanded carrier screening: learning from existing initiatives. <i>European Journal of Public Health</i> , <b>2017</b> , 27, 372-377	2.1	25
74	Attitudes of cystic fibrosis patients and parents toward carrier screening and related reproductive issues. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, 506-12	5.3	21

73	Psychological factors associated with the intention to choose for risk-reducing mastectomy in family cancer clinic attendees. <i>Breast</i> , <b>2016</b> , 30, 66-72	3.6	18
72	Trial by Dutch laboratories for evaluation of non-invasive prenatal testing. Part I-clinical impact. <i>Prenatal Diagnosis</i> , <b>2016</b> , 36, 1083-1090	3.2	92
71	Trial by Dutch laboratories for evaluation of non-invasive prenatal testing. Part II-women's perspectives. <i>Prenatal Diagnosis</i> , <b>2016</b> , 36, 1091-1098	3.2	38
70	Mothers' Views on Longer Storage of Neonatal Dried Blood Spots for Specific Secondary Uses. <i>Public Health Genomics</i> , <b>2016</b> , 19, 25-33	1.9	4
69	Opinions of maternity care professionals and other stakeholders about integration of maternity care: a qualitative study in the Netherlands. <i>BMC Pregnancy and Childbirth</i> , <b>2016</b> , 16, 188	3.2	22
68	Present to future: what the reasons for declining first-trimester combined testing tell us about accepting or declining cell-free DNA testing. <i>Prenatal Diagnosis</i> , <b>2016</b> , 36, 587-90	3.2	9
67	Responsible implementation of expanded carrier screening. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, e1-e12	5.3	167
66	Preferences for prenatal tests for Down syndrome: an international comparison of the views of pregnant women and health professionals. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, 968-75	5.3	38
65	Users evaluate a detailed familial risk questionnaire as valuable and no more time consuming than a simple enquiry in a web-based diabetes risk assessment tool. <i>Public Health</i> , <b>2016</b> , 130, 87-90	4	1
64	Do people from the Jewish community prefer ancestry-based or pan-ethnic expanded carrier screening?. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, 171-7	5.3	17
63	Advantages of expanded universal carrier screening: what is at stake?. <i>European Journal of Human Genetics</i> , <b>2016</b> , 25, 17-21	5.3	32
62	Maternal Plasma DNA and RNA Sequencing for Prenatal Testing. <i>Advances in Clinical Chemistry</i> , <b>2016</b> , 74, 63-102	5.8	17
61	Genetic Screening for Disease <b>2015</b> , 926-931		
60	Current and best practices of genetic testing for maturity onset diabetes of the young: views of professional experts. <i>Public Health Genomics</i> , <b>2015</b> , 18, 52-9	1.9	12
59	Targeted carrier screening for four recessive disorders: high detection rate within a founder population. <i>European Journal of Medical Genetics</i> , <b>2015</b> , 58, 123-8	2.6	28
58	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. Summary and recommendations. <i>European Journal of Human Genetics</i> , <b>2015</b> ,	5.3	9
57	NIPT-based screening for Down syndrome and beyond: what do pregnant women think?. <i>Prenatal Diagnosis</i> , <b>2015</b> , 35, 598-604	3.2	49
56	A genetic diagnosis of maturity-onset diabetes of the young (MODY): experiences of patients and family members. <i>Diabetic Medicine</i> , <b>2015</b> , 32, 1385-92	3.5	7

55	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. <i>European Journal of Human Genetics</i> , <b>2015</b> , 23, 1438-50	5.3	189
54	Determining the genome-wide kinship coefficient seems unhelpful in distinguishing consanguineous couples with a high versus low risk for adverse reproductive outcome. <i>BMC Medical Genetics</i> , <b>2015</b> , 16, 50	2.1	1
53	A decade of molecular genetic testing for MODY: a retrospective study of utilization in The Netherlands. <i>European Journal of Human Genetics</i> , <b>2015</b> , 23, 29-33	5.3	14
52	Changing to NIPT as a first-tier screening test and future perspectives: opinions of health professionals. <i>Prenatal Diagnosis</i> , <b>2015</b> , 35, 1316-23	3.2	22
51	Attitudes of cystic fibrosis patients and their parents towards direct-to-consumer genetic testing for carrier status. <i>Personalized Medicine</i> , <b>2015</b> , 12, 99-107	2.2	6
50	Screening for Familial Hypercholesterolemia in Children: What Can We Learn From Adult Screening Programs?. <i>Healthcare (Switzerland)</i> , <b>2015</b> , 3, 1018-30	3.4	10
49	Do Health Professionals Need Additional Competencies for Stratified Cancer Prevention Based on Genetic Risk Profiling?. <i>Journal of Personalized Medicine</i> , <b>2015</b> , 5, 191-212	3.6	9
48	Effect of comprehensive oncogenetics training interventions for general practitioners, evaluated at multiple performance levels. <i>PLoS ONE</i> , <b>2015</b> , 10, e0122648	3.7	24
47	Effectiviteit van nascholing over (onco)genetica. <i>Huisarts En Wetenschap</i> , <b>2014</b> , 57, 294-297	0.1	
46	First steps in exploring prospective exome sequencing of consanguineous couples. <i>European Journal of Medical Genetics</i> , <b>2014</b> , 57, 613-6	2.6	10
45	Developing a framework for implementation of genetic services: learning from examples of testing for monogenic forms of common diseases. <i>Journal of Community Genetics</i> , <b>2014</b> , 5, 337-47	2.5	20
44	Public attitudes towards preventive genomics and personal interest in genetic testing to prevent disease: a survey study. <i>European Journal of Public Health</i> , <b>2014</b> , 24, 768-75	2.1	39
43	Consanguinity and endogamy in the Netherlands: demographic and medical genetic aspects. <i>Human Heredity</i> , <b>2014</b> , 77, 161-6	1.1	7
42	Sustained effects of online genetics education: a randomized controlled trial on oncogenetics. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22, 310-6	5.3	32
41	Effectiveness of oncogenetics training on general practitioners: consultation skills: a randomized controlled trial. <i>Genetics in Medicine</i> , <b>2014</b> , 16, 45-52	8.1	29
40	Attitudes of pregnant women and male partners towards non-invasive prenatal testing and widening the scope of prenatal screening. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22, 1345-50	5.3	77
39	Consanguineous marriage and reproductive risk: attitudes and understanding of ethnic groups practising consanguinity in Western society. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22, 452-7	5.3	27
38	Comment on Gialluisi et al. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22, 157	5.3	1

37	Informed consent for exome sequencing in diagnostics: exploring first experiences and views of professionals and patients. <i>Clinical Genetics</i> , <b>2014</b> , 85, 417-22	4	35
36	Retinoblastoma and Reproductive Decision-Making. <i>Tumors of the Central Nervous System</i> , <b>2014</b> , 75-82		
35	Ethnicity, educational level and attitudes contribute to parental intentions about genetic testing for child obesity. <i>Journal of Community Genetics</i> , <b>2013</b> , 4, 243-50	2.5	3
34	The effectiveness of a graphical presentation in addition to a frequency format in the context of familial breast cancer risk communication: a multicenter controlled trial. <i>BMC Medical Informatics and Decision Making</i> , <b>2013</b> , 13, 55	3.6	9
33	Reflecting on earlier experiences with unsolicited findings: points to consider for next-generation sequencing and informed consent in diagnostics. <i>Human Mutation</i> , <b>2013</b> , 34, 1322-8	4.7	44
32	Public attitudes towards genetic testing revisited: comparing opinions between 2002 and 2010. <i>European Journal of Human Genetics</i> , <b>2013</b> , 21, 793-9	5.3	77
31	PS6 - 5. Towards best practice guidelines for genetic testing for Maturity-Onset Diabetes of the Young. <i>Nederlands Tijdschrift Voor Diabetologie</i> , <b>2013</b> , 11, 151-152	0	
30	A morass of considerations exploring attitudes towards ethnicity-based haemoglobinopathy-carrier screening in primary care. <i>Family Practice</i> , <b>2013</b> , 30, 604-10	1.9	4
29	Genetic testing and implications for personalized medicine: changes in public and healthcare professional perspectives. <i>Personalized Medicine</i> , <b>2013</b> , 10, 217-219	2.2	2
28	Challenges in the care for consanguineous couples: an exploratory interview study among general practitioners and midwives. <i>BMC Family Practice</i> , <b>2012</b> , 13, 105	2.6	7
27	Prioritization of future genetics education for general practitioners: a Delphi study. <i>Genetics in Medicine</i> , <b>2012</b> , 14, 323-9	8.1	36
26	Attitudes of general practitioners and midwives towards ethnicity-based haemoglobinopathy-carrier screening. <i>European Journal of Human Genetics</i> , <b>2012</b> , 20, 1112-7	5.3	9
25	Genetic educational needs and the role of genetics in primary care: a focus group study with multiple perspectives. <i>BMC Family Practice</i> , <b>2011</b> , 12, 5	2.6	86
24	Lay perceptions of predictive testing for diabetes based on DNA test results versus family history assessment: a focus group study. <i>BMC Public Health</i> , <b>2011</b> , 11, 535	4.1	14
23	A low risk is still a risk exploring women's attitudes towards genetic testing for breast cancer susceptibility in order to target disease prevention. <i>Public Health Genomics</i> , <b>2011</b> , 14, 238-47	1.9	34
22	Preconceptional genetic carrier testing and the commercial offer directly-to-consumers. <i>Human Reproduction</i> , <b>2011</b> , 26, 972-7	5.7	110
21	Autosomal recessive disease in children of consanguineous parents: inferences from the proportion of compound heterozygotes. <i>Journal of Community Genetics</i> , <b>2010</b> , 1, 37-40	2.5	15
20	Using family history information to promote healthy lifestyles and prevent diseases; a discussion of the evidence. <i>BMC Public Health</i> , <b>2010</b> , 10, 248	4.1	74



19	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. <i>BMC Medical Genetics</i> , <b>2010</b> , 11, 113	2.1	14
18	Impact of communicating familial risk of diabetes on illness perceptions and self-reported behavioral outcomes: a randomized controlled trial. <i>Diabetes Care</i> , <b>2009</b> , 32, 597-9	14.6	53
17	Preconceptional ancestry-based carrier couple screening for cystic fibrosis and haemoglobinopathies: what determines the intention to participate or not and actual participation?. <i>European Journal of Human Genetics</i> , <b>2009</b> , 17, 999-1009	5.3	34
16	Family history of diabetes: exploring perceptions of people at risk in the Netherlands. <i>Preventing Chronic Disease</i> , <b>2009</b> , 6, A54	3.7	20
15	Design of the BRISC study: a multicentre controlled clinical trial to optimize the communication of breast cancer risks in genetic counselling. <i>BMC Cancer</i> , <b>2008</b> , 8, 283	4.8	15
14	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. <i>Genetics in Medicine</i> , <b>2008</b> , 10, 820-30	8.1	40
13	Clinical geneticists and genetic counselors' views on the communication of genetic risks: a qualitative study. <i>Patient Education and Counseling</i> , <b>2008</b> , 73, 42-9	3.1	25
12	Presenting health risk information in different formats: the effect on participants' cognitive and emotional evaluation and decisions. <i>Patient Education and Counseling</i> , <b>2008</b> , 73, 443-7	3.1	66
11	Public attitudes toward genetic testing: perceived benefits and objections. <i>Genetic Testing and Molecular Biomarkers</i> , <b>2006</b> , 10, 139-45		59
10	Developing and optimizing a decisional instrument using self-reported ancestry for carrier screening in a multi-ethnic society. <i>Genetics in Medicine</i> , <b>2006</b> , 8, 502-9	8.1	15
9	Deficiency of knowledge of genetics and genetic tests among general practitioners, gynecologists, and pediatricians: a global problem. <i>Genetics in Medicine</i> , <b>2005</b> , 7, 605-10	8.1	165
8	Deficient knowledge of genetics relevant for daily practice among medical students nearing graduation. <i>Genetics in Medicine</i> , <b>2005</b> , 7, 295-301	8.1	53
7	Public experiences, knowledge and expectations about medical genetics and the use of genetic information. <i>Public Health Genomics</i> , <b>2004</b> , 7, 33-43	1.9	91
6	Attitudes of potential providers towards preconceptional cystic fibrosis carrier screening. <i>Journal of Genetic Counseling</i> , <b>2004</b> , 13, 31-44	2.5	25
5	Preconceptional cystic fibrosis carrier screening: attitudes and intentions of the target population. <i>Genetic Testing and Molecular Biomarkers</i> , <b>2004</b> , 8, 80-9		31
4	Offering preconceptional cystic fibrosis carrier couple screening in the absence of established preconceptional care services. <i>Public Health Genomics</i> , <b>2003</b> , 6, 5-13	1.9	33
3	How should preconceptional cystic fibrosis carrier screening be provided? Opinions of potential providers and the target population. <i>Public Health Genomics</i> , <b>2003</b> , 6, 157-65	1.9	14
2	Evaluation of cystic fibrosis carrier screening programs according to genetic screening criteria. <i>Genetics in Medicine</i> , <b>2002</b> , 4, 241-9	8.1	29



1 Genetic Carrier Testing1-4