

Aad Tibben

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4578702/publications.pdf>

Version: 2024-02-01

62
papers

2,675
citations

201385

27
h-index

189595

50
g-index

66
all docs

66
docs citations

66
times ranked

2277
citing authors

#	ARTICLE	IF	CITATIONS
1	Long-Term Psychological Impact of Carrying a BRCA1/2 Mutation and Prophylactic Surgery: A 5-Year Follow-Up Study. <i>Journal of Clinical Oncology</i> , 2003, 21, 3867-3874.	0.8	268
2	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. <i>European Journal of Human Genetics</i> , 2015, 23, 1438-1450.	1.4	260
3	Psychological impact of receiving aBRCA1/BRCA2 test result. <i>American Journal of Medical Genetics Part A</i> , 2001, 98, 15-24.	2.4	151
4	Three-year follow-up after presymptomatic testing for Huntington's disease in tested individuals and partners.. <i>Health Psychology</i> , 1997, 16, 20-35.	1.3	120
5	The counselees' view of an unclassified variant in BRCA1/2: recall, interpretation, and impact on life. <i>Psycho-Oncology</i> , 2008, 17, 822-830.	1.0	112
6	Predictive testing for Huntington's disease. <i>Brain Research Bulletin</i> , 2007, 72, 165-171.	1.4	105
7	On attitudes and appreciation 6 months after predictive DNA testing for huntington disease in the Dutch program. <i>American Journal of Medical Genetics Part A</i> , 1993, 48, 103-111.	2.4	101
8	Adverse Effects of Predictive Testing for Huntington Disease Underestimated: Long-Term Effects 7-10 Years After the Test.. <i>Health Psychology</i> , 2004, 23, 189-197.	1.3	98
9	Cross-Sectional Study on Prevalences of Psychiatric Disorders in Mutation Carriers of Huntington's Disease Compared With Mutation-Negative First-Degree Relatives. <i>Journal of Clinical Psychiatry</i> , 2008, 69, 1804-1810.	1.1	90
10	DNA-Testing for Huntington's disease in The Netherlands: A retrospective study on psychosocial effects. <i>American Journal of Medical Genetics Part A</i> , 1992, 44, 94-99.	2.4	84
11	Presymptomatic DNA testing for Huntington disease: Identifying the need for psychological intervention. <i>American Journal of Medical Genetics Part A</i> , 1993, 48, 137-144.	2.4	81
12	Presymptomatic DNA-testing for Huntington disease: Pretest attitudes and expectations of applicants and their partners in the Dutch program. <i>American Journal of Medical Genetics Part A</i> , 1993, 48, 10-16.	2.4	76
13	Distress in individuals facing predictive DNA testing for autosomal dominant late-onset disorders: Comparing questionnaire results with in-depth interviews. , 1998, 75, 62-74.		62
14	Opening the psychological black box in genetic counseling. The psychological impact of DNA testing is predicted by the counselees' perception, the medical impact by the pathogenic or uninformative BRCA1/2 result. <i>Psycho-Oncology</i> , 2012, 21, 29-42.	1.0	60
15	A Hereditary Disorder In the Family and the Family Life Cycle: Huntington Disease as a Paradigm. <i>Family Process</i> , 2002, 41, 677-692.	1.4	59
16	Psychological distress in women at increased risk for breast cancer: the role of risk perception. <i>European Journal of Cancer</i> , 2004, 40, 2056-2063.	1.3	56
17	What's the message? Interpretation of an uninformative BRCA1/2 test result for women at risk of familial breast cancer. <i>Genetics in Medicine</i> , 2005, 7, 239-245.	1.1	55
18	Comparison of individuals opting for BRCA1/2 or HNPCC genetic susceptibility testing with regard to coping, illness perceptions, illness experiences, family system characteristics and hereditary cancer distress. <i>Patient Education and Counseling</i> , 2007, 65, 58-68.	1.0	55

#	ARTICLE	IF	CITATIONS
19	A whisper-game perspective on the family communication of DNA-test results: a retrospective study on the communication process of BRCA1/2-test results between proband and relatives. <i>Familial Cancer</i> , 2011, 10, 87-96.	0.9	48
20	Men at risk of being a mutation carrier for hereditary breast/ovarian cancer: an exploration of attitudes and psychological functioning during genetic testing. <i>European Journal of Human Genetics</i> , 2001, 9, 492-500.	1.4	47
21	The common sense model of self-regulation and psychological adjustment to predictive genetic testing: a prospective study. <i>Psycho-Oncology</i> , 2007, 16, 1121-1129.	1.0	44
22	The counselees' self-reported request for psychological help in genetic counseling for hereditary breast/ovarian cancer: not only psychopathology matters. <i>Psycho-Oncology</i> , 2013, 22, 902-910.	1.0	35
23	Family communication matters: The impact of telling relatives about unclassified variants and uninformative DNA-test results. <i>Genetics in Medicine</i> , 2011, 13, 333-341.	1.1	33
24	Attachment in families with Huntington's disease. <i>Patient Education and Counseling</i> , 2006, 63, 246-254.	1.0	32
25	Coordinated multidisciplinary care for Huntington's disease. An outpatient department. <i>Brain Research Bulletin</i> , 2009, 80, 192-195.	1.4	32
26	Exploring the short-term impact of DNA-testing in breast cancer patients: The counselees'™ perception matters, but the actual BRCA1/2 result does not. <i>Patient Education and Counseling</i> , 2012, 86, 239-251.	1.0	31
27	Mothers'™ psychological adaptation to Duchenne/Becker muscular dystrophy. <i>European Journal of Human Genetics</i> , 2016, 24, 633-637.	1.4	31
28	Barriers and facilitators to clinical trial participation among parents of children with pediatric neuromuscular disorders. <i>Clinical Trials</i> , 2018, 15, 139-148.	0.7	30
29	Long-term psychological distress in women at risk for hereditary breast cancer adhering to regular surveillance: a risk profile. <i>Psycho-Oncology</i> , 2013, 22, 598-604.	1.0	27
30	Genetic testing in familial melanoma: uptake and implications. <i>Psycho-Oncology</i> , 2008, 17, 790-796.	1.0	25
31	A counselee-oriented perspective on risk communication in genetic counseling: Explaining the inaccuracy of the counselees' risk perception shortly after BRCA1/2 test result disclosure. <i>Genetics in Medicine</i> , 2011, 13, 800-811.	1.1	25
32	Offering a choice between NIPT and invasive PND in prenatal genetic counseling: the impact of clinician characteristics on patients'™ test uptake. <i>European Journal of Human Genetics</i> , 2019, 27, 235-243.	1.4	25
33	Whole-exome sequencing in pediatrics: parents'™ considerations toward return of unsolicited findings for their child. <i>European Journal of Human Genetics</i> , 2016, 24, 1681-1687.	1.4	22
34	Attitudes of Dutch general practitioners towards presymptomatic DNA-testing for Huntington disease. <i>Clinical Genetics</i> , 1993, 43, 63-68.	1.0	20
35	The impact of social and personal resources on psychological distress in women at risk for hereditary breast cancer. <i>Psycho-Oncology</i> , 2012, 21, 153-160.	1.0	20
36	The Psychological Challenges of Replacing Conventional Karyotyping with Genomic SNP Array Analysis in Prenatal Testing. <i>Journal of Clinical Medicine</i> , 2014, 3, 713-723.	1.0	20

#	ARTICLE	IF	CITATIONS
37	Genetic counseling does not fulfill the counselees' need for certainty in hereditary breast/ovarian cancer families: an explorative assessment. <i>Psycho-Oncology</i> , 2013, 22, 1167-1176.	1.0	19
38	Implementing non-invasive prenatal testing (NIPT) in the Netherlands: An interview study exploring opinions about and experiences with societal pressure, reimbursement, and an expanding scope. <i>Journal of Genetic Counseling</i> , 2020, 29, 112-121.	0.9	19
39	Do Attachment Style and Emotion Regulation Strategies Indicate Distress in Predictive Testing?. <i>Journal of Genetic Counseling</i> , 2015, 24, 862-871.	0.9	18
40	“Watching time tick by” – Decision making for Duchenne muscular dystrophy trials. <i>Contemporary Clinical Trials</i> , 2016, 46, 1-6.	0.8	18
41	Nature and Development of Huntington Disease in a Nursing Home Population. <i>Cognitive and Behavioral Neurology</i> , 2005, 18, 215-222.	0.5	17
42	Euthanasia and Advance Directives in Huntington's Disease: Qualitative Analysis of Interviews with Patients. <i>Journal of Huntington's Disease</i> , 2013, 2, 323-330.	0.9	17
43	Most women recover from psychological distress after postoperative complications following implant or DIEP flap breast reconstruction: A prospective long-term follow-up study. <i>PLoS ONE</i> , 2017, 12, e0174455.	1.1	14
44	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> , 2015, , .	1.4	13
45	Contributors to and consequences of burnout among clinical genetic counselors in the United States. <i>Journal of Genetic Counseling</i> , 2022, 31, 269-278.	0.9	11
46	The Effect of Predictive Testing in Adult-Onset Neurodegenerative Diseases on Social and Personal Life. <i>Journal of Genetic Counseling</i> , 2018, 27, 947-954.	0.9	10
47	“Be an ambassador for change that you would like to see” a call to action to all stakeholders for co-creation in healthcare and medical research to improve quality of life of people with a neuromuscular disease. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 126.	1.2	10
48	Validation of a scale for assessing attitudes towards outcomes of genetic cancer testing among primary care providers and breast specialists. <i>PLoS ONE</i> , 2017, 12, e0178447.	1.1	9
49	Perhaps the subject of the questionnaire was too sensitive: Do we expect too much too soon?. <i>Journal of Huntington's Disease</i> , 2014, 3, 229-232.	0.9	7
50	Offspring of a parent with genetic disease: Childhood experiences and adult psychological characteristics.. <i>Health Psychology</i> , 2014, 33, 1445-1453.	1.3	7
51	The Position of Neuromuscular Patients in Shared Decision Making. Report from the 235th ENMC Workshop: Milan, Italy, January 19-20, 2018. <i>Journal of Neuromuscular Diseases</i> , 2019, 6, 161-172.	1.1	7
52	Patients' and surgeons' experiences after failed breast reconstruction: A qualitative study. <i>Journal of Plastic, Reconstructive and Aesthetic Surgery</i> , 2021, 74, 1480-1485.	0.5	7
53	Parents, their children, whole exome sequencing and unsolicited findings: growing towards the child's future autonomy. <i>European Journal of Human Genetics</i> , 2021, 29, 911-919.	1.4	7
54	Risk Assessment for Huntington's Disease for (Future) Offspring Requires Offering Preconceptional CAG Analysis to Both Partners. <i>Journal of Huntington's Disease</i> , 2019, 8, 71-78.	0.9	7

#	ARTICLE	IF	CITATIONS
55	General Practitioners and Breast Surgeons in France, Germany, Netherlands and the UK show variable breast cancer risk communication profiles. BMC Cancer, 2015, 15, 243.	1.1	6
56	Psychosocial Needs and Facilitators of Mothers Caring for Children with Duchenne/Becker Muscular Dystrophy. Journal of Genetic Counseling, 2018, 27, 197-203.	0.9	6
57	Aesthetic outcome after implant and DIEP flap breast reconstruction: An exploratory, prospective comparison of 25 cases. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2015, 68, 1018-1019.	0.5	5
58	Mindfulness-Based Stress Reduction in Pre-symptomatic Genetic Frontotemporal Dementia: A Pilot Study. Frontiers in Psychiatry, 2022, 13, 864391.	1.3	1
59	Reply to Oliver W. Quarrell et al.: Letter in response to Tibben et al., Risk Assessment for Huntington's Disease for (Future) Offspring Requires Offering Preconceptional CAG Analysis to Both Partners. Journal of Huntington's Disease, 2019, , 1-2.	0.9	0
60	Reply to Oliver W Quarrell et al.: "Letter in response to Tibben et al., Risk Assessment for Huntington's Disease for (Future) Offspring Requires Offering Preconceptional CAG Analysis to Both Partners" Journal of Huntington's Disease, 2019, 8, 361-362.	0.9	0
61	Genetic Testing Expanded. , 2019, , 1-16.		0
62	Uncertainties in Genome Sequencing. , 2019, , 75-88.		0