Mirjam Plantinga

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8784797/publications.pdf

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17 papers	396 citations	10 10 h-index	993246 17 g-index
17	17	17	649
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Parental experiences of rapid exome sequencing in cases with major ultrasound anomalies during pregnancy. Prenatal Diagnosis, 2022, 42, 762-774.	1.1	17
2	Couple-based expanded carrier screening provided by general practitioners to couples in the Dutch general population: psychological outcomes and reproductive intentions. Genetics in Medicine, 2021, 23, 1761-1768.	1.1	10
3	GP-provided couple-based expanded preconception carrier screening in the Dutch general population: who accepts the test-offer and why?. European Journal of Human Genetics, 2020, 28, 182-192.	1.4	15
4	Cognitive and affective outcomes of genetic counselling in the Netherlands at group and individual level: a personalized approach seems necessary. European Journal of Human Genetics, 2020, 28, 1187-1195.	1.4	8
5	Practical Barriers and Facilitators Experienced by Patients, Pharmacists and Physicians to the Implementation of Pharmacogenomic Screening in Dutch Outpatient Hospital Care—An Explorative Pilot Study. Journal of Personalized Medicine, 2020, 10, 293.	1.1	15
6	A validated PROM in genetic counselling: the psychometric properties of the Dutch version of the Genetic Counselling Outcome Scale. European Journal of Human Genetics, 2019, 27, 681-690.	1.4	12
7	Informing relatives at risk of inherited cardiac conditions: experiences and attitudes of healthcare professionals and counselees. European Journal of Human Genetics, 2019, 27, 1341-1350.	1.4	10
8	Feasibility of couple-based expanded carrier screening offered by general practitioners. European Journal of Human Genetics, 2019, 27, 691-700.	1.4	48
9	Expanded carrier screening for autosomal recessive conditions in health care: Arguments for a coupleâ€based approach and examination of couples' views. Prenatal Diagnosis, 2019, 39, 369-378.	1.1	20
10	The phenotypic spectrum of proximal 6q deletions based on a large cohort derived from social media and literature reports. European Journal of Human Genetics, 2018, 26, 1478-1489.	1.4	31
11	Expanded carrier screening: what determines intended participation and can this be influenced by message framing and narrative information?. European Journal of Human Genetics, 2017, 25, 793-800.	1.4	17
12	Population-based preconception carrier screening: how potential users from the general population view a test for 50 serious diseases. European Journal of Human Genetics, 2016, 24, 1417-1423.	1.4	70
13	Assessment and Treatment of Pain during Treatment of Buruli Ulcer. PLoS Neglected Tropical Diseases, 2015, 9, e0004076.	1.3	8
14	Is there a duty to recontact in light of new genetic technologies? A systematic review of the literature. Genetics in Medicine, 2015, 17, 668-678.	1.1	77
15	Maximising the efficiency of clinical screening programmes: balancing predictive genetic testing with a right not to know. European Journal of Human Genetics, 2015, 23, 1124-1128.	1.4	6
16	Training healthcare professionals as moral case deliberation facilitators: evaluation of a Dutch training programme. Journal of Medical Ethics, 2012, 38, 630-635.	1.0	19
17	Choosing Whether to Buy or Make: The Contracting Out of Employment Reintegration Services by Dutch Municipalities. Social Policy and Administration, 2011, 45, 245-263.	2.1	13