

# Annemieke J M H Verkerk

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5995634/publications.pdf>

Version: 2024-02-01

13  
papers

3,654  
citations

932766

10  
h-index

1199166

12  
g-index

13  
all docs

13  
docs citations

13  
times ranked

4163  
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of a gene (FMR-1) containing a CGG repeat coincident with a breakpoint cluster region exhibiting length variation in fragile X syndrome. <i>Cell</i> , 1991, 65, 905-914.	13.5	3,285
2	Mutation in the AP4M1 Gene Provides a Model for Neuroaxonal Injury in Cerebral Palsy. <i>American Journal of Human Genetics</i> , 2009, 85, 40-52.	2.6	156
3	Allelic imbalance of expression and epigenetic regulation within the alpha-synuclein wild-type and p.Ala53Thr alleles in Parkinson disease. <i>Human Mutation</i> , 2010, 31, 685-691.	1.1	53
4	A comparison of genotyping arrays. <i>European Journal of Human Genetics</i> , 2021, 29, 1611-1624.	1.4	43
5	Reduced penetrance of pathogenic ACMG variants in a deeply phenotyped cohort study and evaluation of ClinVar classification over time. <i>Genetics in Medicine</i> , 2020, 22, 1812-1820.	1.1	24
6	Boston type craniosynostosis: Report of a second mutation in <i>MSX2</i> . <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 2626-2633.	0.7	23
7	Linkage analysis and whole exome sequencing identify a novel candidate gene in a Dutch multiple sclerosis family. <i>Multiple Sclerosis Journal</i> , 2019, 25, 909-917.	1.4	19
8	Complex craniosynostosis is associated with the 2p15p16.1 microdeletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 244-253.	0.7	16
9	The Genetics of Atypical Femur Fractures – a Systematic Review. <i>Current Osteoporosis Reports</i> , 2021, 19, 123-130.	1.5	15
10	CXorf56, a dendritic neuronal protein, identified as a new candidate gene for X-linked intellectual disability. <i>European Journal of Human Genetics</i> , 2018, 26, 552-560.	1.4	12
11	Unbalanced der(5)t(5;20) translocation associated with megalencephaly, perisylvian polymicrogyria, polydactyly and hydrocephalus. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 1488-1497.	0.7	6
12	Congenital hypopituitarism in two brothers with a duplication of the <i>acrogigantism gene</i> ™ GPR101: clinical findings and review of the literature. <i>Pituitary</i> , 2021, 24, 229-241.	1.6	2
13	Reply: PRKAR1B mutations are a rare cause of FUS negative neuronal intermediate filament inclusion disease. <i>Brain</i> , 2015, 138, e358-e358.	3.7	0