## Wolfram Demaerel

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4633260/publications.pdf

Version: 2024-02-01

1684188 1720034 6 180 5 7 citations g-index h-index papers 7 7 7 537 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Genetic contributors to risk of schizophrenia in the presence of a 22q11.2 deletion. Molecular Psychiatry, 2021, 26, 4496-4510.	7.9	87
2	Pathogenic variants in CDC45 on the remaining allele in patients with a chromosome 22q11.2 deletion result in a novel autosomal recessive condition. Genetics in Medicine, 2020, 22, 326-335.	2.4	17
3	The 22q11 low copy repeats are characterized by unprecedented size and structural variability. Genome Research, 2019, 29, 1389-1401.	5 <b>.</b> 5	39
4	Atypical chromosome 22q11.2 deletions are complex rearrangements and have different mechanistic origins. Human Molecular Genetics, 2019, 28, 3724-3733.	2.9	7
5	Deletion size analysis of 1680 22q11.2DS subjects identifies a new recombination hotspot on chromosome 22q11.2. Human Molecular Genetics, 2018, 27, 1150-1163.	2.9	22
6	Reciprocal 22q11.2 Deletion and Duplication in Siblings with Karyotypically Normal Parents. Cytogenetic and Genome Research, 2016, 148, 1-5.	1.1	2