Maarten J A Van Den Bossche

List of Publications by Year
in descending order

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17 papers	463 citations	933447 10 h-index	996975 15 g-index
17 all docs	17 docs citations	17 times ranked	1472 citing authors

#	Article	IF	Citations
1	A paleo-neurologic investigation of the social brain hypothesis in frontotemporal dementia. Cerebral Cortex, 2023, 33, 622-633.	2.9	2
2	Management Approaches for Behavioural and Psychological Symptoms of Dementia. , 2021, , 129-153.		0
3	How non-rapid eye movement sleep and Alzheimer pathology are linked. World Journal of Psychiatry, 2021, 11, 1027-1038.	2.7	4
4	Hippocampal volume change following ECT is mediated by rs699947 in the promotor region of VEGF. Translational Psychiatry, 2019, 9, 191.	4.8	17
5	Points to consider regarding DSM alternatives. Lancet Psychiatry,the, 2019, 6, 894-895.	7.4	0
6	Clinical validation of the Psychotic Depression Assessment Scale (PDAS) against independent global severity ratings in older adults. Acta Neuropsychiatrica, 2018, 30, 203-208.	2.1	4
7	Longitudinal Evaluation of the Psychomotor Syndrome in Schizophrenia. Journal of Neuropsychiatry and Clinical Neurosciences, 2014, 26, 359-368.	1.8	14
8	The nature of the relationship of psychomotor slowing with negative symptomatology in schizophrenia. Cognitive Neuropsychiatry, 2014, 19, 36-46.	1.3	39
9	Identification of rare copy number variants in high burden schizophrenia families. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 273-282.	1.7	23
10	Multiplex amplicon quantification screening the ABCA13 gene for copy number variation in schizophrenia and bipolar disorder. Psychiatric Genetics, 2012, 22, 269-270.	1.1	8
11	Optimized filtering reduces the error rate in detecting genomic variants by short-read sequencing. Nature Biotechnology, 2012, 30, 61-68.	17.5	211
12	Less Cognitive and Neurological Deficits in Schizophrenia Patients Carrying Risk Variant in <i>ZNF804A</i>. Neuropsychobiology, 2012, 66, 158-166.	1.9	15
13	Co-occurrence of Marfan syndrome and schizophrenia: What can be learned?. European Journal of Medical Genetics, 2012, 55, 252-255.	1.3	11
14	Rare copy number variants in neuropsychiatric disorders: Specific phenotype or not?. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 812-822.	1.7	34
15	Identification of a CACNA2D4 deletion in late onset bipolar disorder patients and implications for the involvement of voltageâ€dependent calcium channels in psychiatric disorders. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 465-475.	1.7	27
16	Sequencing of DISC1 Pathway Genes Reveals Increased Burden of Rare Missense Variants in Schizophrenia Patients from a Northern Swedish Population. PLoS ONE, 2011, 6, e23450.	2.5	46
17	PCM1 and schizophrenia: A replication study in the Northern Swedish population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 1240-1243.	1.7	8