Martje E Van Egmond

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8449093/publications.pdf

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25 papers 844 citations

759233 12 h-index 25 g-index

25 all docs

25 docs citations

25 times ranked

1208 citing authors

#	Article	IF	CITATIONS
1	Dystonia in children and adolescents: a systematic review and a new diagnostic algorithm. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 774-781.	1.9	124
2	Ramsay hunt syndrome: Clinical characterization of progressive myoclonus ataxia caused by <i>GOSR2</i> mutation. Movement Disorders, 2014, 29, 139-143.	3.9	113
3	Van Buchem disease: Clinical, biochemical, and densitometric features of patients and disease carriers. Journal of Bone and Mineral Research, 2013, 28, 848-854.	2.8	102
4	Lossâ€ofâ€Function Variants in <scp>HOPS</scp> Complex Genes <scp><i>VPS16</i></scp> and <scp><i>VPS41</i></scp> Cause Early Onset Dystonia Associated with Lysosomal Abnormalities. Annals of Neurology, 2020, 88, 867-877.	5.3	70
5	A novel diagnostic approach to patients with myoclonus. Nature Reviews Neurology, 2015, 11, 687-697.	10.1	67
6	A post hoc study on gene panel analysis for the diagnosis of dystonia. Movement Disorders, 2017, 32, 569-575.	3.9	59
7	Efficacy of hematopoietic cell transplantation in metachromatic leukodystrophy: the Dutch experience. Blood, 2016, 127, 3098-3101.	1.4	56
8	Improvement of White Matter Changes on Neuroimaging Modalities After Stem Cell Transplant in Metachromatic Leukodystrophy. JAMA Neurology, 2013, 70, 779.	9.0	44
9	Toward adaptive deep brain stimulation for dystonia. Neurosurgical Focus, 2018, 45, E3.	2.3	38
10	Non-motor effects of deep brain stimulation in dystonia: A systematic review. Parkinsonism and Related Disorders, 2018, 55, 26-44.	2.2	22
11	Myoclonus in childhood-onset neurogenetic disorders: The importance of early identification and treatment. European Journal of Paediatric Neurology, 2015, 19, 726-729.	1.6	20
12	Bilateral Pallidotomy for Dystonia: A Systematic Review. Movement Disorders, 2021, 36, 547-557.	3.9	19
13	Dystoniaâ€deafness syndrome caused by a βâ€actin gene mutation and response to deep brain stimulation. Movement Disorders, 2017, 32, 162-165.	3.9	13
14	Variable Interpretation of the Dystonia Consensus Classification Items Compromises Its Solidity. Movement Disorders, 2019, 34, 317-320.	3.9	12
15	Challenges in Clinicogenetic Correlations: One Phenotype – Many Genes. Movement Disorders Clinical Practice, 2021, 8, 311-321.	1.5	12
16	The efficacy of the modified Atkins diet in North Sea Progressive Myoclonus Epilepsy: an observational prospective open-label study. Orphanet Journal of Rare Diseases, 2017, 12, 45.	2.7	11
17	Diagnostic approach to paediatric movement disorders: a clinical practice guide. Developmental Medicine and Child Neurology, 2021, 63, 252-258.	2.1	11
18	Crossing barriers: a multidisciplinary approach to children and adults with young-onset movement disorders. Journal of Clinical Movement Disorders, 2018, 5, 3.	2.2	10

#	Article	IF	CITATION
19	A detailed description of the phenotypic spectrum of North Sea Progressive Myoclonus Epilepsy in a large cohort of seventeen patients. Parkinsonism and Related Disorders, 2020, 72, 44-48.	2.2	9
20	Reversal of Status Dystonicus after Relocation of Pallidal Electrodes in DYT6 Generalized Dystonia. Tremor and Other Hyperkinetic Movements, 2018, 8, 530.	2.0	9
21	Cortical Myoclonus in a Young Boy with <i><scp>GOSR</scp>2</i> Mutation Mimics Chorea. Movement Disorders Clinical Practice, 2015, 2, 61-63.	1.5	7
22	Diaphragmatic weakness caused by neuroborreliosis. Clinical Neurology and Neurosurgery, 2011, 113, 153-155.	1.4	6
23	The Effectiveness of Deep Brain Stimulation in Dystonia: A Patient-Centered Approach. Tremor and Other Hyperkinetic Movements, 2020, 10, 2.	2.0	5
24	A novel diagnostic approach for patients with adult-onset dystonia. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 1039-1048.	1.9	3
25	Are we on the right track in DBS surgery for dystonic head tremor? Polymyography is a promising answer. Parkinsonism and Related Disorders, 2021, 93, 74-76.	2.2	2