## Edward J Cupler

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7640333/publications.pdf

Version: 2024-02-01

44 papers

1,247 citations

686830 13 h-index 34 g-index

46 all docs 46 docs citations

46 times ranked

 $\begin{array}{c} 2029 \\ \text{citing authors} \end{array}$ 

#	Article	IF	CITATIONS
1	Pattern reversal visual evoked potentials (prVEPs) in autosomal recessive hereditary spastic paraplegia with thin corpus callosum (ARHSPTCC) patients with SPG 11 mutations in Saudi Arabia, cross section hospital base study. Journal of the Neurological Sciences, 2022, 434, 120144.	0.3	o
2	Real-world effectiveness and safety profile of teriflunomide in the management of multiple sclerosis in the Gulf Cooperation Council countries: An expert consensus narrative review. Multiple Sclerosis Journal - Experimental, Translational and Clinical, 2022, 8, 205521732210771.	0.5	2
3	Clinical, Neurophysiological, Radiological, Pathological, and Genetic Features of Dysferlinopathy in Saudi Arabia. Frontiers in Neuroscience, 2022, 16, 815556.	1.4	5
4	SARS-COV-2 Triggers the Development of Class I and Class II HLA Antibodies in Recovered Convalescent Plasma Donors. Intervirology, 2022, 65, 230-235.	1.2	1
5	A Multidisciplinary Perspective Addressing the Diagnostic Challenges of Late-Onset Pompe Disease in the Arabian Peninsula Region Developed From an Expert Group Meeting. Journal of Neuromuscular Diseases, 2022, 9, 661-673.	1.1	2
6	The disease course of multiple sclerosis before and during COVID-19 pandemic: A retrospective five-year study. Multiple Sclerosis and Related Disorders, 2022, 65, 103985.	0.9	5
7	Further delineation of <scp><i>MYO18B</i></scp> â€related autosomal recessive <scp>Klippelâ€Feil</scp> syndrome with myopathy and facial dysmorphism. American Journal of Medical Genetics, Part A, 2021, 185, 370-376.	0.7	4
8	Family Planning for People with Multiple Sclerosis in Saudi Arabia: an Expert Consensus. Multiple Sclerosis International, 2021, 2021, 1-8.	0.4	2
9	A Prospective Multicenter Study for Assessing MusiQoL Validity among Arabic-Speaking MS Patients Treated with Subcutaneous Interferon $\hat{I}^{2}$ -1a. Multiple Sclerosis International, 2021, 2021, 1-7.	0.4	2
10	Managing multiple sclerosis in the Covid19 era: a review of the literature and consensus report from a panel of experts in Saudi Arabia. Multiple Sclerosis and Related Disorders, 2021, 51, 102925.	0.9	11
11	Prevalence of Pediatric Onset Multiple Sclerosis in Saudi Arabia. Multiple Sclerosis International, 2021, 2021, 1-6.	0.4	3
12	Acute reversible rhabdomyolysis during direct-acting antiviral hepatitis C virus treatment: a case report. Journal of Medical Case Reports, 2021, 15, 627.	0.4	1
13	Amyotrophic lateral sclerosis care in Saudi Arabia: A survey of providers' perceptions. Brain and Behavior, 2020, 10, e01795.	1.0	1
14	Cladribine Tablets and Relapsing–Remitting Multiple Sclerosis: A Pragmatic, Narrative Review of What Physicians Need to Know. Neurology and Therapy, 2020, 9, 11-23.	1.4	6
15	Abstract P4-09-13: Spectrum analysis of the inherited mutations of BRCA1/-2 genes that associated with high risk breast cancer in Saudi population. , 2020, , .		0
16	Current management of Duchenne muscular dystrophy in the Middle East: expert report. Neurodegenerative Disease Management, 2019, 9, 123-133.	1.2	15
17	Lessons Learned from Large-Scale, First-Tier Clinical Exome Sequencing in a Highly Consanguineous Population. American Journal of Human Genetics, 2019, 104, 1182-1201.	2.6	184
18	Autosomal recessive ADCY5-Related dystonia and myoclonus: Expanding the genetic spectrum of ADCY5-Related movement disorders. Parkinsonism and Related Disorders, 2019, 64, 145-149.	1.1	18

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19	To report trans-cranial magnetic stimulation to describe the NORMAL LATENCIES OF major nerve with central conduction time among Saudi Arabian population; a hospital based study. Journal of the Neurological Sciences, 2019, 405, 359-360.	0.3	O
20	Alazami syndrome due to LARP7 gene mutation: Expanding the phenotype. Journal of the Neurological Sciences, 2019, 405, 272-273.	0.3	1
21	Non-convulsive seizures and electroencephalography findings as predictors of clinical outcomes at a tertiary intensive care unit in Saudi Arabia. Clinical Neurology and Neurosurgery, 2018, 171, 95-99.	0.6	2
22	The landscape of genetic diseases in Saudi Arabia based on the first 1000 diagnostic panels and exomes. Human Genetics, 2017, 136, 921-939.	1.8	209
23	Improving Reports Turnaround Time: An Essential Healthcare Quality Dimension. Studies in Health Technology and Informatics, 2016, 226, 205-8.	0.2	2
24	Neuromyelitis optica (NMO) antibody positive disorder: a case series from Saudi Arabia. Journal of the Neurological Sciences, 2015, 357, e296.	0.3	0
25	Diagnosis and treatment of late-onset Pompe disease in the Middle East and North Africa region: consensus recommendations from an expert group. BMC Neurology, 2015, 15, 205.	0.8	28
26	Congenital absence of gluteal muscles, optic nerve hypoplasia, and central nervous system hamartomas. Clinical Dysmorphology, 2012, 21, 106-108.	0.1	0
27	Consensus treatment recommendations for lateâ€onset Pompe disease. Muscle and Nerve, 2012, 45, 319-333.	1.0	130
28	A novel syndromic form of sensory-motor polyneuropathy is linked to chromosome 22q13.31-q13.33. Clinical Genetics, 2011, 79, 193-195.	1.0	7
29	The Heterogeneity of Pompe Disease: Early Data on Genotype From the Pompe Registry. Clinical Therapeutics, 2011, 33, S27-S28.	1.1	0
30	Anterior interosseous nerve syndrome following peripheral catheterization: Magnetic resonance imaging and electromyography correlation. Muscle and Nerve, 2011, 43, 758-760.	1.0	1
31	Prolonged Improvement After Rituximab: Two Cases of Resistant Muscle-Specific Receptor Tyrosine Kinase + Myasthenia Gravis. Journal of Clinical Neuromuscular Disease, 2010, 12, 85-87.	0.3	13
32	Nephrogenic systemic fibrosis presenting as myopathy: A case report with histopathologic correlation. Neuromuscular Disorders, 2010, 20, 411-413.	0.3	8
33	Macrophagic Myofasciitis in Children Is a Localized Reaction to Vaccination. Journal of Child Neurology, 2008, 23, 614-619.	0.7	30
34	Adult Myopathies with Tubular Aggregates: Clinical Correlates. FASEB Journal, 2008, 22, 708.22.	0.2	0
35	Miliary tuberculomas of the brain: case report. Clinical Neurology and Neurosurgery, 2006, 108, 411-414.	0.6	14
36	Neurotrophin 4/5 immunoassay: identification of sources of errors for the quantification of neurotrophins. Journal of Neuroscience Methods, 2000, 99, 119-127.	1.3	9

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37	Miyoshi myopathy in Saudi Arabia: clinical, electrophysiological, histopathological and radiological features. Neuromuscular Disorders, 1998, 8, 321-326.	0.3	35
38	Miyoshi myopathy in Saudi Arabia. Neuromuscular Disorders, 1997, 7, 461-462.	0.3	1
39	A controlled study of intravenous immunoglobulin in demyelinating neuropathy with IgM gammopathy. Annals of Neurology, 1996, 40, 792-795.	2.8	200
40	Acetylcholine receptor antibodies as a marker of treatable fatigue in HIV-1 infected individuals. , 1996, 19, 1186-1188.		9
41	Inclusion body myositis in HIV-1 and HTLV-1 infected patients. Brain, 1996, 119, 1887-1893.	3.7	134
42	Early features of zidovudine-associated myopathy: histopathological findings and clinical correlations. Acta Neuropathologica, 1995, 90, 1-6.	3.9	70
43	Exacerbation of peripheral neuropathy by lamivudine. Lancet, The, 1995, 345, 460-461.	6.3	30
44	Effect of High-Dose Intravenous Immunoglobulin on Amyotrophic Lateral Sclerosis and Multifocal Motor Neuropathy. Archives of Neurology, 1994, 51, 861-864.	4.9	52