# 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 | 1,247 | 686830 13 | 377514 34 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 46 | 46 | 46 | 2029 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | 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 |
| 2 | A controlled study of intravenous immunoglobulin in demyelinating neuropathy with IgM gammopathy. Annals of Neurology, 1996, 40, 792-795. | 2.8 | 200 |
| 3 | 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 |
| 4 | Inclusion body myositis in HIV-1 and HTLV-1 infected patients. Brain, 1996, 119, 1887-1893. | 3.7 | 134 |
| 5 | Consensus treatment recommendations for lateâ€onset Pompe disease. Muscle and Nerve, 2012, 45, 319-333. | 1.0 | 130 |
| 6 | Early features of zidovudine-associated myopathy: histopathological findings and clinical correlations. Acta Neuropathologica, 1995, 90, 1-6. | 3.9 | 70 |
| 7 | Effect of High-Dose Intravenous Immunoglobulin on Amyotrophic Lateral Sclerosis and Multifocal Motor Neuropathy. Archives of Neurology, 1994, 51, 861-864. | 4.9 | 52 |
| 8 | Miyoshi myopathy in Saudi Arabia: clinical, electrophysiological, histopathological and radiological features. Neuromuscular Disorders, 1998, 8, 321-326. | 0.3 | 35 |
| 9 | Exacerbation of peripheral neuropathy by lamivudine. Lancet, The, 1995, 345, 460-461. | 6.3 | 30 |
| 10 | Macrophagic Myofasciitis in Children Is a Localized Reaction to Vaccination. Journal of Child Neurology, 2008, 23, 614-619. | 0.7 | 30 |
| 11 | 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 |
| 12 | 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 |
| 13 | Current management of Duchenne muscular dystrophy in the Middle East: expert report. Neurodegenerative Disease Management, 2019, 9, 123-133. | 1.2 | 15 |
| 14 | Miliary tuberculomas of the brain: case report. Clinical Neurology and Neurosurgery, 2006, 108, 411-414. | 0.6 | 14 |
| 15 | 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 |
| 16 | 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 |
| 17 | Acetylcholine receptor antibodies as a marker of treatable fatigue in HIV-1 infected individuals. , 1996, 19, 1186-1188. | | 9 |
| 18 | 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Nephrogenic systemic fibrosis presenting as myopathy: A case report with histopathologic correlation. Neuromuscular Disorders, 2010, 20, 411-413. | 0.3 | 8 |
| 20 | 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 |
| 21 | Cladribine Tablets and Relapsingâ€“Remitting Multiple Sclerosis: A Pragmatic, Narrative Review of What Physicians Need to Know. Neurology and Therapy, 2020, 9, 11-23. | 1.4 | 6 |
| 22 | Clinical, Neurophysiological, Radiological, Pathological, and Genetic Features of Dysferlinopathy in Saudi Arabia. Frontiers in Neuroscience, 2022, 16, 815556. | 1.4 | 5 |
| 23 | 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 |
| 24 | Further delineation of <scp>MYO18B</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 |
| 25 | Prevalence of Pediatric Onset Multiple Sclerosis in Saudi Arabia. Multiple Sclerosis International, 2021, 2021, 1-6. | 0.4 | 3 |
| 26 | 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 |
| 27 | Family Planning for People with Multiple Sclerosis in Saudi Arabia: an Expert Consensus. Multiple Sclerosis International, 2021, 2021, 1-8. | 0.4 | 2 |
| 28 | A Prospective Multicenter Study for Assessing MusiQoL Validity among Arabic-Speaking MS Patients Treated with Subcutaneous Interferon Î²-1a. Multiple Sclerosis International, 2021, 2021, 1-7. | 0.4 | 2 |
| 29 | Improving Reports Turnaround Time: An Essential Healthcare Quality Dimension. Studies in Health Technology and Informatics, 2016, 226, 205-8. | 0.2 | 2 |
| 30 | 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 |
| 31 | 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 |
| 32 | Miyoshi myopathy in Saudi Arabia. Neuromuscular Disorders, 1997, 7, 461-462. | 0.3 | 1 |
| 33 | Anterior interosseous nerve syndrome following peripheral catheterization: Magnetic resonance imaging and electromyography correlation. Muscle and Nerve, 2011, 43, 758-760. | 1.0 | 1 |
| 34 | Alazami syndrome due to LARP7 gene mutation: Expanding the phenotype. Journal of the Neurological Sciences, 2019, 405, 272-273. | 0.3 | 1 |
| 35 | Amyotrophic lateral sclerosis care in Saudi Arabia: A survey of providersâ€™ perceptions. Brain and Behavior, 2020, 10, e01795. | 1.0 | 1 |
| 36 | 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 |

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|---|---|---|---|
| 37 | 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 |
| 38 | The Heterogeneity of Pompe Disease: Early Data on Genotype From the Pompe Registry. Clinical Therapeutics, 2011, 33, S27-S28. | 1.1 | 0 |
| 39 | Congenital absence of gluteal muscles, optic nerve hypoplasia, and central nervous system hamartomas. Clinical Dysmorphology, 2012, 21, 106-108. | 0.1 | 0 |
| 40 | Neuromyelitis optica (NMO) antibody positive disorder: a case series from Saudi Arabia. Journal of the Neurological Sciences, 2015, 357, e296. | 0.3 | 0 |
| 41 | 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 | 0 |
| 42 | Adult Myopathies with Tubular Aggregates: Clinical Correlates. FASEB Journal, 2008, 22, 708.22. | 0.2 | 0 |
| 43 | 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 |
| 44 | 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 | 0 |