

Arun B Taly

List of Publications by Year in descending order

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Version: 2024-02-01

114
papers

1,761
citations

331670

21
h-index

377865

34
g-index

114
all docs

114
docs citations

114
times ranked

2193
citing authors

#	ARTICLE	IF	CITATIONS
1	Wilson Disease. <i>Medicine (United States)</i> , 2007, 86, 112-121.	1.0	224
2	Sleep-related disorders among a healthy population in South India. <i>Neurology India</i> , 2012, 60, 68.	0.4	98
3	Sleep disturbances in juvenile myoclonic epilepsy: A sleep questionnaire-based study. <i>Epilepsy and Behavior</i> , 2012, 23, 305-309.	1.7	61
4	Autonomic dysfunction in Wilson's disease - a clinical and electrophysiological study. <i>Clinical Autonomic Research</i> , 2002, 12, 185-189.	2.5	51
5	Mortality in mechanically ventilated patients of Guillain Barré Syndrome. <i>Annals of Indian Academy of Neurology</i> , 2011, 14, 262.	0.5	46
6	Cognitive profile and structural findings in Wilson's disease: A neuropsychological and MRI-based study. <i>Neurology India</i> , 2010, 58, 708.	0.4	42
7	Palatal tremor, progressive multiple cranial nerve palsies, and cerebellar ataxia: A case report and review of literature of palatal tremors in neurodegenerative disease. <i>Movement Disorders</i> , 1999, 14, 689-693.	3.9	38
8	Sleep disorders in children with cerebral palsy and its correlation with sleep disturbance in primary caregivers and other associated factors. <i>Annals of Indian Academy of Neurology</i> , 2014, 17, 473.	0.5	34
9	Bodyweight-supported treadmill training for retraining gait among chronic stroke survivors: A randomized controlled study. <i>Annals of Physical and Rehabilitation Medicine</i> , 2016, 59, 235-241.	2.3	34
10	Diffusion Tensor Imaging (DTI) and its clinical correlates in drug naïve Wilson's disease. <i>Metabolic Brain Disease</i> , 2013, 28, 455-462.	2.9	32
11	Transient computerised tomographic (CT) abnormalities following partial seizures. <i>Acta Neurologica Scandinavica</i> , 1985, 72, 26-29.	2.1	30
12	Prognosis of patients with Guillain-Barré syndrome requiring mechanical ventilation. <i>Neurology India</i> , 2011, 59, 707.	0.4	30
13	Magnetic resonance imaging correlates of genetically characterized patients with mitochondrial disorders: A study from south India. <i>Mitochondrion</i> , 2015, 25, 6-16.	3.4	28
14	Functional outcome following rehabilitation in chronic severe traumatic brain injury patients: A prospective study. <i>Annals of Indian Academy of Neurology</i> , 2012, 15, 120.	0.5	27
15	Mitochondrial leukoencephalopathies: A border zone between acquired and inherited white matter disorders in children?. <i>Multiple Sclerosis and Related Disorders</i> , 2018, 20, 84-92.	2.0	27
16	Contribution of nuclear and mitochondrial gene mutations in mitochondrial encephalopathy, lactic acidosis, and stroke-like episodes (MELAS) syndrome. <i>Journal of Neurology</i> , 2021, 268, 2192-2207.	3.6	27
17	Clinical Features, Therapeutic Response, and Follow-Up in Pediatric Anti-N-Methyl-d-Aspartate Receptor Encephalitis: Experience from a Tertiary Care University Hospital in India. <i>Neuropediatrics</i> , 2016, 47, 024-032.	0.6	26
18	Effect of valproate on the sleep microstructure of juvenile myoclonic epilepsy patients – a cross-sectional CAP based study. <i>Sleep Medicine</i> , 2016, 17, 129-133.	1.6	26

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19	Genetic Analysis of PLA2G6 in 22 Indian Families with Infantile Neuroaxonal Dystrophy, Atypical Late-Onset Neuroaxonal Dystrophy and Dystonia Parkinsonism Complex. PLoS ONE, 2016, 11, e0155605.	2.5	26
20	Effect of pranayama and meditation as an add-on therapy in rehabilitation of patients with Guillain-Barré syndrome—a randomized control pilot study. Disability and Rehabilitation, 2013, 35, 57-62.	1.8	24
21	Altered polysomnographic profile in juvenile myoclonic epilepsy. Epilepsy Research, 2014, 108, 459-467.	1.6	22
22	Central Pontine Signal Changes in Wilson's Disease: Distinct MRI Morphology and Sequential Changes with Deferrioxamine Therapy. Journal of Neuroimaging, 2007, 17, 286-291.	2.0	20
23	Depression and sleep disturbances in patients with multiple sclerosis and correlation with associated fatigue. Journal of Neurosciences in Rural Practice, 2013, 04, 387-391.	0.8	20
24	Clinical and magnetic resonance imaging findings in patients with Leigh syndrome and SURF1 mutations. Brain and Development, 2014, 36, 807-812.	1.1	20
25	Alterations in Polysomnographic (PSG) profile in drug-naïve Parkinson's disease. Annals of Indian Academy of Neurology, 2014, 17, 287.	0.5	19
26	Role of ankle foot orthosis in improving locomotion and functional recovery in patients with stroke: A prospective rehabilitation study. Journal of Neurosciences in Rural Practice, 2016, 7, 544-549.	0.8	19
27	Comparing sleep profiles between patients with juvenile myoclonic epilepsy and symptomatic partial epilepsy: Sleep questionnaire-based study. Epilepsy and Behavior, 2017, 66, 34-38.	1.7	19
28	Leber's Hereditary Optic Neuropathy—Specific Mutation m.11778G>A Exists on Diverse Mitochondrial Haplogroups in India. , 2017, 58, 3923.		19
29	Guillain-Barre Syndrome—rehabilitation outcome, residual deficits and requirement of lower limb orthosis for locomotion at 1 year follow-up. Disability and Rehabilitation, 2010, 32, 1897-1902.	1.8	18
30	Audiological manifestations in mitochondrial encephalomyopathy lactic acidosis and stroke like episodes (MELAS) syndrome. Clinical Neurology and Neurosurgery, 2016, 148, 17-21.	1.4	18
31	Neuropathy in elderly: lessons learnt from nerve biopsy. Age and Ageing, 2015, 44, 312-317.	1.6	17
32	Effect of carbamazepine on the sleep microstructure of temporal lobe epilepsy patients: a cyclic alternating pattern-based study. Sleep Medicine, 2016, 27-28, 80-85.	1.6	17
33	Mitochondrial oxidative phosphorylation disorders in children: Phenotypic, genotypic and biochemical correlations in 85 patients from South India. Mitochondrion, 2017, 32, 42-49.	3.4	17
34	Outcome of epilepsy in patients with mitochondrial disorders: Phenotype genotype and magnetic resonance imaging correlations. Clinical Neurology and Neurosurgery, 2018, 164, 182-189.	1.4	17
35	Child Neurology: Sjögren-Larsson syndrome. Neurology, 2017, 88, e1-e4.	1.1	16
36	IL-23/IL-17 immune axis in Guillain Barré Syndrome: Exploring newer vistas for understanding pathobiology and therapeutic implications. Cytokine, 2018, 103, 77-82.	3.2	16

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37	Cognitive and Functional Outcomes following Inpatient Rehabilitation in Patients with Acquired Brain Injury: A Prospective Follow-up Study. <i>Journal of Neurosciences in Rural Practice</i> , 2017, 08, 357-363.	0.8	15
38	Anti-NMDA receptor encephalitis presenting as postpartum psychosis—a clinical description and review. <i>Archives of Women's Mental Health</i> , 2018, 21, 465-469.	2.6	15
39	Comprehensive cytokine profiling provides evidence for a multi-lineage Th responses in Guillain Barré Syndrome. <i>Cytokine</i> , 2018, 110, 58-62.	3.2	15
40	Comparing the efficacy of sodium valproate and levetiracetam following initial lorazepam in elderly patients with generalized convulsive status epilepticus (GCSE): A prospective randomized controlled pilot study. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 65, 111-117.	2.0	15
41	Stroke with supernumerary phantom limb: case study, review of literature and pathogenesis. <i>Acta Neuropsychiatrica</i> , 2008, 20, 256-264.	2.1	13
42	Seasonal variation in the clinical recovery of patients with Guillain Barré syndrome requiring mechanical ventilation. <i>Neurology India</i> , 2013, 61, 349.	0.4	13
43	Child Neurology: Molybdenum cofactor deficiency. <i>Neurology</i> , 2015, 85, e175-8.	1.1	13
44	Lack of heart rate variability during apnea in patients with juvenile myoclonic epilepsy (JME). <i>Sleep and Breathing</i> , 2015, 19, 1175-1183.	1.7	13
45	Diagnosis of myasthenia gravis: Comparison of anti-nicotinic acetyl choline receptor antibodies, repetitive nerve stimulation and Neostigmine tests at a tertiary neuro care centre in India, a ten year study. <i>Journal of Neuroimmunology</i> , 2016, 292, 81-84.	2.3	13
46	Child Neurology: Ethylmalonic encephalopathy. <i>Neurology</i> , 2020, 94, e1336-e1339.	1.1	13
47	Tropical ataxic neuropathy – A century old enigma. <i>Neurology India</i> , 2016, 64, 1151.	0.4	13
48	Prevalence of fatigue in Guillain-Barre syndrome in neurological rehabilitation setting. <i>Annals of Indian Academy of Neurology</i> , 2014, 17, 331.	0.5	12
49	Complications in mechanically ventilated patients of Guillain-Barre syndrome and their prognostic value. <i>Journal of Neurosciences in Rural Practice</i> , 2017, 08, 068-073.	0.8	12
50	Th17 pathway signatures in a large Indian cohort of Guillain Barré syndrome. <i>Journal of Neuroimmunology</i> , 2018, 323, 125-130.	2.3	12
51	Rehabilitation interventions to improve locomotor outcome in chronic stroke survivors: A prospective, repeated-measure study. <i>Neurology India</i> , 2015, 63, 347.	0.4	11
52	Leukodystrophies and Genetic Leukoencephalopathies in Children Specified by Exome Sequencing in an Expanded Gene Panel. <i>Journal of Child Neurology</i> , 2020, 35, 433-441.	1.4	11
53	Antecedent infections in Guillain-Barré syndrome patients from south India. <i>Journal of the Peripheral Nervous System</i> , 2021, 26, 298-306.	3.1	11
54	MG-QoL-15 scores in treated myasthenia gravis: Experience from a university hospital in India. <i>Neurology India</i> , 2016, 64, 405.	0.4	11

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55	Ambulation following spinal cord injury and its correlates. <i>Annals of Indian Academy of Neurology</i> , 2015, 18, 167.	0.5	10
56	Peripheral neuropathy in genetically characterized patients with mitochondrial disorders: A study from south India. <i>Mitochondrion</i> , 2016, 27, 1-5.	3.4	10
57	Hypersomnolence-hyperkinetic movement disorder in a child with compound heterozygous mutation in 4-aminobutyrate aminotransferase (ABAT) gene. <i>Brain and Development</i> , 2017, 39, 161-165.	1.1	10
58	Genetic analysis of ATP7B in 102 south Indian families with Wilson disease. <i>PLoS ONE</i> , 2019, 14, e0215779.	2.5	10
59	Tangier's disease: An uncommon cause of facial weakness and non-length dependent demyelinating neuropathy. <i>Annals of Indian Academy of Neurology</i> , 2016, 19, 137.	0.5	10
60	Non-Wilsonian hepatolenticular degeneration: Clinical and MRI observations in four families from south India. <i>Journal of Clinical Neuroscience</i> , 2016, 27, 91-94.	1.5	9
61	Urodynamic profile in acute transverse myelitis patients: Its correlation with neurological outcome. <i>Journal of Neurosciences in Rural Practice</i> , 2017, 08, 044-048.	0.8	9
62	Ganglioside complex antibodies in an Indian cohort of Guillain-Barré syndrome. <i>Muscle and Nerve</i> , 2020, 62, 728-734.	2.2	9
63	Evidence of altered Th17 pathway signatures in the cerebrospinal fluid of patients with Guillain Barré Syndrome. <i>Journal of Clinical Neuroscience</i> , 2020, 75, 176-180.	1.5	9
64	Sleep profile and Polysomnography in patients with drug-resistant temporal lobe epilepsy (TLE) due to hippocampal sclerosis (HS) and the effect of epilepsy surgery on sleep-a prospective cohort study. <i>Sleep Medicine</i> , 2021, 80, 176-183.	1.6	9
65	Serum fibroblast growth factor 21 and growth differentiation factor 15: Two sensitive biomarkers in the diagnosis of mitochondrial disorders. <i>Mitochondrion</i> , 2021, 60, 170-177.	3.4	9
66	Neuropsychiatric Manifestations of Pediatric NMDA Receptor Autoimmune Encephalitis. primary care companion for CNS disorders, <i>The</i> , 2017, 19, .	0.6	9
67	Pediatric opsoclonus-myoclonus-ataxia syndrome: Experience from a tertiary care university hospital. <i>Neurology India</i> , 2018, 66, 1332.	0.4	9
68	The "Double Panda" Sign in Leigh Disease. <i>Journal of Child Neurology</i> , 2014, 29, 980-982.	1.4	8
69	Audiological findings in Infantile Refsum disease. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2015, 79, 1366-1369.	1.0	8
70	Palatal Tremor Revisited: Disorder with Nosological Diversity and Etiological Heterogeneity. <i>Canadian Journal of Neurological Sciences</i> , 2018, 45, 243-247.	0.5	8
71	Clinical profile and treatment response in patients with CASPR2 antibody-associated neurological disease. <i>Annals of Indian Academy of Neurology</i> , 2021, 24, 178.	0.5	8
72	Clinical, hematological, and imaging observations in a 25-year-old woman with abetalipoproteinemia. <i>Annals of Indian Academy of Neurology</i> , 2014, 17, 113.	0.5	7

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73	Clinical and Neuroimaging Features in Two Children with Mutations in the Mitochondrial ND5 Gene. <i>Neuropediatrics</i> , 2015, 46, 277-281.	0.6	7
74	NREM Sleep and Antiepileptic Medications Modulate Epileptiform Activity by Altering Cortical Synchrony. <i>Clinical EEG and Neuroscience</i> , 2018, 49, 417-424.	1.7	7
75	PMP22 Gene-associated Neuropathies: Phenotypic Spectrum in a Cohort from India. <i>Journal of Molecular Neuroscience</i> , 2020, 70, 778-789.	2.3	7
76	Neurogenic bladder following myelopathies: Has it any correlation with neurological and functional recovery?. <i>Journal of Neurosciences in Rural Practice</i> , 2014, 05, S013-S016.	0.8	6
77	Oculomotor Apraxia in Gaucher Disease. <i>Pediatric Neurology</i> , 2015, 52, 468-469.	2.1	6
78	Giant Axonal Neuropathy. <i>Journal of Child Neurology</i> , 2015, 30, 912-915.	1.4	6
79	Role of pulse methylprednisolone in epileptic encephalopathy: A retrospective observational analysis. <i>Epilepsy Research</i> , 2021, 173, 106611.	1.6	6
80	Clinico-pathological and Molecular Spectrum of Mitochondrial Polymerase β Mutations in a Cohort from India. <i>Journal of Molecular Neuroscience</i> , 2021, 71, 2219-2228.	2.3	6
81	Serial macro-architectural alterations with levodopa in Parkinson's disease: Polysomnography (PSC)-based analysis. <i>Annals of Indian Academy of Neurology</i> , 2015, 18, 309.	0.5	6
82	Variations within Toll-like receptor (TLR) and TLR signaling pathway-related genes and their synergistic effects on the risk of Guillain-Barré syndrome. <i>Journal of the Peripheral Nervous System</i> , 2022, 27, 131-143.	3.1	6
83	Guillain-Barré Syndrome in Postpartum Period: Rehabilitation Issues and Outcome – Three Case Reports. <i>Journal of Neurosciences in Rural Practice</i> , 2017, 08, 475-477.	0.8	5
84	Exome sequencing in adult neurology practice: Challenges and rewards in a mixed resource setting. <i>Clinical Neurology and Neurosurgery</i> , 2018, 174, 48-56.	1.4	5
85	Autoantibodies in acquired myasthenia gravis: Clinical phenotype and immunological correlation. <i>Acta Neurologica Scandinavica</i> , 2019, 139, 428-437.	2.1	5
86	Urinary symptoms in patients with Parkinson's disease and progressive supranuclear palsy: Urodynamic findings and management of bladder dysfunction. <i>Annals of Indian Academy of Neurology</i> , 2019, 22, 432.	0.5	5
87	Role of altered IL-33/ST2 immune axis in the immunobiology of Guillain-Barré syndrome. <i>European Journal of Neurology</i> , 2022, 29, 2074-2083.	3.3	5
88	Pitfalls in the diagnosis of leprosy neuropathy: Lessons learnt from a University hospital in an endemic zone. <i>Journal of the Neurological Sciences</i> , 2015, 357, 252-256.	0.6	4
89	Pulmonary Involvement in Patients with Guillain-Barré Syndrome in Subacute Phase. <i>Journal of Neurosciences in Rural Practice</i> , 2017, 08, 412-416.	0.8	4
90	Heightened Background Cortical Synchrony in Patients With Epilepsy: EEG Phase Synchrony Analysis During Awake and Sleep Stages Using Novel Ensemble Measure. <i>Clinical EEG and Neuroscience</i> , 2018, 49, 177-186.	1.7	4

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91	Ethylmalonic encephalopathy ETHE1 p. D165H mutation alters the mitochondrial function in human skeletal muscle proteome. <i>Mitochondrion</i> , 2021, 58, 64-71.	3.4	4
92	Prevalence of depression, fatigue, and sleep disturbances in patients with myelopathy: Their relation with functional and neurological recovery. <i>Journal of Spinal Cord Medicine</i> , 2016, 39, 620-626.	1.4	3
93	Management of Anti-N-Methyl-D-Aspartate (NMDA) Receptor Encephalitis in Children. <i>Journal of Child Neurology</i> , 2017, 32, 513-514.	1.4	3
94	Ankle-Foot Orthosis in Duchenne Muscular Dystrophy: A 4-Year Experience in a Multidisciplinary Neuromuscular Disorders Clinic. <i>Indian Journal of Pediatrics</i> , 2017, 84, 211-215.	0.8	3
95	Spectrum and evolution of EEG changes in Anti-NMDAR encephalitis. <i>Annals of Indian Academy of Neurology</i> , 2021, 24, 396.	0.5	3
96	Electrophysiological observations in critically ill Guillain-Barré syndrome. <i>Neurology India</i> , 2016, 64, 914.	0.4	3
97	Differential improvement of the sleep quality among patients with juvenile myoclonic epilepsy with valproic acid: A longitudinal sleep questionnaire-based study. <i>Annals of Indian Academy of Neurology</i> , 2015, 18, 403-7.	0.5	3
98	Case Report: Chronic Fungal Meningitis Masquerading as Tubercular Meningitis. <i>American Journal of Tropical Medicine and Hygiene</i> , 2020, 103, 1473-1479.	1.4	3
99	Child Neurology: Hereditary Folate Malabsorption. <i>Neurology</i> , 2021, 97, 40-43.	1.1	2
100	Whole exome sequencing reveals a homozygous C1QBP deletion as the cause of progressive external ophthalmoplegia and multiple mtDNA deletions. <i>Neuromuscular Disorders</i> , 2021, 31, 859-864.	0.6	2
101	An uncommon cause of bifacial weakness and non-length-dependent demyelinating neuropathy. <i>Annals of Indian Academy of Neurology</i> , 2015, 18, 445-8.	0.5	2
102	Cholecysto-cardiac link: The heart of the matter. <i>Neurology India</i> , 2019, 67, 391.	0.4	2
103	Speech-Language and swallowing manifestations and rehabilitation in an 11-year-old girl with MELAS syndrome. <i>Journal of Pediatric Neurosciences</i> , 2015, 10, 31-4.	0.3	2
104	Genetically Established Familial Amyloidotic Polyneuropathy from India: Narrating the Diagnostic "Odyssey" and a Mini Review. <i>Neurology India</i> , 2020, 68, 1084.	0.4	2
105	Ictal Generalized EEG Attenuation (IGEAA) and hypopnea in a child with occipital type 1 cortical dysplasia - Is it a biomarker for SUDEP?. <i>Annals of Indian Academy of Neurology</i> , 2014, 18, 103-7.	0.5	1
106	Etiologic Spectrum of Biopsy-Proven Peripheral Neuropathies in Childhood from a Resource-Poor Setting. <i>Journal of Child Neurology</i> , 2015, 30, 707-715.	1.4	1
107	Clinical Reasoning: West syndrome, pontocerebellar hypoplasia, and hypomyelination in a 6-month-old boy. <i>Neurology</i> , 2018, 91, e1652-e1656.	1.1	1
108	Vogt-koyanagi-harada syndrome - A neurologist's perspective. <i>Annals of Indian Academy of Neurology</i> , 2021, 24, 405.	0.5	1

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109	Leukodystrophy due to eIF2B mutations in adults. Canadian Journal of Neurological Sciences, 2021, , 1-11.	0.5	1
110	Is Perls Prussian Blue Stain for Hemosiderin a Useful Adjunct in the Diagnosis of Vasculitic Neuropathies?. Neurology India, 2021, 69, 140.	0.4	1
111	Vasculitic neuropathy in elderly: A study from a tertiary care university hospital in South India. Annals of Indian Academy of Neurology, 2016, 19, 323.	0.5	1
112	Reply to Letter to the Editor: Hearing impairment in m.3243A>G carriers requires comprehensive work- and follow-up. Clinical Neurology and Neurosurgery, 2016, 150, 198-199.	1.4	0
113	Enigmas in immunobiology of Guillain-Barré syndrome: Ganglioside antibodies and beyond!. Neurology India, 2017, 65, 973.	0.4	0
114	Modulatory effects of vitamin D on IL-33/ST2 immune axis in Guillain-Barré syndrome. Quo Vadis? European Journal of Neurology, 2022, , .	3.3	0