Prashanth Lingappa Kukkle

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4603028/publications.pdf

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37	1,067	471061	433756
papers	citations	17 h-index	31 g-index

39	39	39	1095
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical Study of 668 Indian Subjects with Juvenile, Young, and Early Onset Parkinsonâ€™s Disease. Canadian Journal of Neurological Sciences, 2022, 49, 93-101.	0.3	8
2	Genomeâ€Wide Polygenic Score Predicts Large Number of High Risk Individuals in Monogenic Undiagnosed Young Onset Parkinson's Disease Patients from India. Advanced Biology, 2022, 6, .	1.4	5
3	Reply to: Fatal Familial Insomnia: A Rare Disease with Unique Clinicoâ€Neurophysiological Features. Movement Disorders Clinical Practice, 2021, 8, 164-165.	0.8	2
4	Does Apomorphine Have an Effect on Body Weight? An Observational Study. Canadian Journal of Neurological Sciences, 2021, 48, 281-284.	0.3	1
5	Huntingtonâ€™s disease: The Indian perspective. Annals of Movement Disorders, 2021, 4, 4.	0.3	0
6	Chorea-acanthocytosis: 3 new families with novel genetic and metabolic findings. Annals of Indian Academy of Neurology, 2021, 24, 452.	0.2	1
7	Hypoparathyroidism Masquerading as Corticobasal Syndrome. Movement Disorders Clinical Practice, 2021, 8, 600-603.	0.8	1
8	Cross-sectional analysis of the Parkinsonâ€™s disease Non-motor International Longitudinal Study baseline non-motor characteristics, geographical distribution and impact on quality of life. Scientific Reports, 2021, 11, 9611.	1.6	21
9	The spectrum of movement disorders in tertiary care centers in India: A tale of three cities. Annals of Indian Academy of Neurology, 2021, 24, 721.	0.2	1
10	Genetic Architecture of Parkinson's Disease in the Indian Population: Harnessing Genetic Diversity to Address Critical Gaps in Parkinson's Disease Research. Frontiers in Neurology, 2020, 11, 524.	1.1	23
11	A Case of Autosomal Dominant Ataxia with Vocal Cord Palsy Attributed to a Mutation in the <scp>PRNP</scp> Gene. Movement Disorders Clinical Practice, 2020, 7, 688-692.	0.8	4
12	Apomorphine: The initial indian experience in relation to response tests and pumps. Annals of Indian Academy of Neurology, 2020, 23, 20.	0.2	13
13	Subcutaneous apomorphine in advanced Parkinsonâ€™s disease and its use in Indian population. Annals of Movement Disorders, 2020, 3, 145.	0.3	2
14	Wilson's Disease: Clinical Practice Guidelines of the Indian National Association for Study of the Liver, the Indian Society of Pediatric Gastroenterology, Hepatology and Nutrition, and the Movement Disorders Society of India. Journal of Clinical and Experimental Hepatology, 2019, 9, 74-98.	0.4	63
15	Surgical treatment of dystonia. Expert Review of Neurotherapeutics, 2018, 18, 477-492.	1.4	36
16	Effects of trunk proprioceptive neuromuscular facilitation on dynamic balance, mobility and quality of life in Parkinson's disease. Parkinsonism and Related Disorders, 2016, 22, e68-e69.	1.1	3
17	First Report of Kuforâ€Rakeb Syndrome (<scp>PARK</scp> 9) from India, and a Novel Nonsense Mutation in <scp>ATP</scp>13A2 Gene. Movement Disorders Clinical Practice, 2015, 2, 326-327.	0.8	11
18	Minimal change multiple system atrophy: An aggressive variant?. Movement Disorders, 2015, 30, 960-967.	2.2	45

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19	Over-the-counter self-medication leading to intracranial hypertension in a young lady. Journal of Neurosciences in Rural Practice, 2014, 05, 384-386.	0.3	6
20	Amantadine improves gait in PD patients with STN stimulation. Parkinsonism and Related Disorders, 2013, 19, 316-319.	1.1	24
21	Reply to letter: multiple system atrophy-parkinsonism with slow progression and prolonged survival: A diagnostic catch. Movement Disorders, 2013, 28, 408-408.	2.2	0
22	A randomized trial of varenicline (chantix) for the treatment of spinocerebellar ataxia type 3. Neurology, 2012, 79, 2218-2218.	1.5	8
23	Psychogenic facial movement disorders: Clinical features and associated conditions. Movement Disorders, 2012, 27, 1544-1551.	2.2	93
24	Extreme task specificity: Is it dystonia or another form of motor programming abnormality?. Movement Disorders, 2012, 27, 1202-1203.	2.2	2
25	Multiple system atrophyâ€“parkinsonism with slow progression and prolonged survival: A diagnostic catch. Movement Disorders, 2012, 27, 1186-1190.	2.2	164
26	l-Dopa-Induced Dyskinesiaâ€”Clinical Presentation, Genetics, and Treatment. International Review of Neurobiology, 2011, 98, 31-54.	0.9	46
27	Wilsonâ€™s disease: 31P and 1H MR spectroscopy and clinical correlation. Neuroradiology, 2010, 52, 977-985.	1.1	18
28	Do MRI features distinguish Wilson's disease from other early onset extrapyramidal disorders? An analysis of 100 cases. Movement Disorders, 2010, 25, 672-678.	2.2	78
29	Spectrum of epilepsy in Wilson's disease with electroencephalographic, MR imaging and pathological correlates. Journal of the Neurological Sciences, 2010, 291, 44-51.	0.3	50
30	Wilsonâ€²s disease: An Indian perspective. Neurology India, 2009, 57, 528.	0.2	43
31	Dominant psychiatric manifestations in Wilson's disease: A diagnostic and therapeutic challenge!. Journal of the Neurological Sciences, 2008, 266, 104-108.	0.3	96
32	Psychiatric Manifestations in Wilsonâ€™s Disease: A Cross-Sectional Analysis. Journal of Neuropsychiatry and Clinical Neurosciences, 2008, 20, 81-85.	0.9	48
33	Quality of life in Wilsonâ€²s disease. Annals of Indian Academy of Neurology, 2008, 11, 37.	0.2	16
34	Sequential MRI changes in Wilson's disease with de-coppering therapy: a study of 50 patients. British Journal of Radiology, 2007, 80, 744-749.	1.0	74
35	Subacute Sclerosing Panencephalitis (SSPE): An Insight Into the Diagnostic Errors From a Tertiary Care University Hospital. Journal of Child Neurology, 2007, 22, 683-688.	0.7	34
36	Central Pontine Signal Changes in Wilson's Disease: Distinct MRI Morphology and Sequential Changes with Deâ€Coppering Therapy. Journal of Neuroimaging, 2007, 17, 286-291.	1.0	20

#	Article	IF	Citations
37	Oculogyric Crisis Phenotype of Levodopaâ€Induced Ocular Dyskinesia. Movement Disorders Clinical Practice, 0, , .	0.8	1