## Abbas Tafakhori

List of Publications by Year in descending order

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279798 361022 1,663 106 23 35 citations h-index g-index papers 112 112 112 2937 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Safety and Efficacy of Nanocurcumin as Add-On Therapy to Riluzole in Patients With Amyotrophic Lateral Sclerosis: A Pilot Randomized Clinical Trial. Neurotherapeutics, 2018, 15, 430-438.	4.4	90
2	The synergistic effects of ω-3 fatty acids and nano-curcumin supplementation on tumor necrosis factor (TNF)-α gene expression and serum level in migraine patients. Immunogenetics, 2017, 69, 371-378.	2.4	75
3	Efficacy and safety of oral ketamine versus diclofenac to alleviate mild to moderate depression in chronic pain patients: A double-blind, randomized, controlled trial. Journal of Affective Disorders, 2016, 204, 1-8.	4.1	70
4	A Clinical and Molecular Genetic Study of 50 Families with Autosomal Recessive Parkinsonism Revealed Known and Novel Gene Mutations. Molecular Neurobiology, 2018, 55, 3477-3489.	4.0	67
5	Neurological manifestations as the predictors of severity and mortality in hospitalized individuals with COVID-19: a multicenter prospective clinical study. BMC Neurology, 2021, 21, 116.	1.8	58
6	Memantine for Prophylactic Treatment of Migraine Without Aura: A Randomized Doubleâ€Blind Placeboâ€Controlled Study. Headache, 2016, 56, 95-103.	3.9	57
7	A Novel Combination of I‰-3 Fatty Acids and Nano-Curcumin Modulates Interleukin-6 Gene Expression and High Sensitivity C-reactive Protein Serum Levels in Patients with Migraine: A Randomized Clinical Trial Study. CNS and Neurological Disorders - Drug Targets, 2018, 17, 430-438.	1.4	53
8	The Neuromodulatory Effects of ï‰-3 Fatty Acids and Nano-Curcumin on the COX-2/ iNOS Network in Migraines: A Clinical Trial Study from Gene Expression to Clinical Symptoms. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2019, 19, 874-884.	1.2	53
9	Transplantation of olfactory ensheathing cells on functional recovery and neuropathic pain after spinal cord injury; systematic review and meta-analysis. Scientific Reports, 2018, 8, 325.	3.3	49
10	Bi-allelic variants in RNF170 are associated with hereditary spastic paraplegia. Nature Communications, 2019, 10, 4790.	12.8	39
11	Sensing of Alzheimer's Disease and Multiple Sclerosis Using Nano-Bio Interfaces. Journal of Alzheimer's Disease, 2017, 59, 1187-1202.	2.6	38
12	<i>PTRHD1</i> (C2orf79) mutations lead to autosomal-recessive intellectual disability and parkinsonism. Movement Disorders, 2017, 32, 287-291.	3.9	38
13	Identification of a large homozygous <i>VPS13C</i> deletion in a patient with earlyâ€onset Parkinsonism. Movement Disorders, 2018, 33, 1968-1970.	3.9	38
14	Identification of a LargeDNAJB2Deletion in a Family with Spinal Muscular Atrophy and Parkinsonism. Human Mutation, 2016, 37, 1180-1189.	2.5	36
15	Validation of the Persian Version of the Brief Pain Inventory (BPI-P) in Chronic Pain Patients. Journal of Pain and Symptom Management, 2017, 54, 132-138.e2.	1.2	35
16	The Combined Effects of i‰ -3 Fatty Acids and Nano-Curcumin Supplementation on Intercellular Adhesion Molecule-1 (ICAM-1) Gene Expression and Serum Levels in Migraine Patients. CNS and Neurological Disorders - Drug Targets, 2018, 16, 1120-1126.	1.4	35
17	RIT2 Polymorphisms: Is There a Differential Association?. Molecular Neurobiology, 2017, 54, 2234-2240.	4.0	31
18	Bi-allelic HPDL Variants Cause a Neurodegenerative Disease Ranging from Neonatal Encephalopathy to Adolescent-Onset Spastic Paraplegia. American Journal of Human Genetics, 2020, 107, 364-373.	6.2	30

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19	Application of Ultrasonography and Radiography in Detection of Hemothorax; a Systematic Review and Meta-Analysis. Emergency, 2016, 4, 116-26.	0.6	29
20	Diagnostic Accuracy of Ultrasonography and Radiography in Detection of Pulmonary Contusion; a Systematic Review and Meta-Analysis. Emergency, 2015, 3, 127-36.	0.6	28
21	Early versus late spinal decompression surgery in treatment of traumatic spinal cord injuries; a systematic review and meta-analysis. Emergency, 2017, 5, e37.	0.6	27
22	Variation in the miRNA-433 binding site of FGF20 is a risk factor for Parkinson's disease in Iranian population. Journal of the Neurological Sciences, 2015, 355, 72-74.	0.6	25
23	A genetic variant in miRNA binding site of glutamate receptor 4, metabotropic (GRM4) is associated with increased risk of major depressive disorder. Journal of Affective Disorders, 2017, 208, 218-222.	4.1	25
24	A Preliminary Study Evaluating the Safety and Efficacy of Bumetanide, an NKCC1 Inhibitor, in Patients with Drug-Resistant Epilepsy. CNS Drugs, 2019, 33, 283-291.	5.9	24
25	Evaluation of 99mTc-TRODAT-1 SPECT in the diagnosis of Parkinson's disease versus other progressive movement disorders. Annals of Nuclear Medicine, 2016, 30, 153-162.	2.2	23
26	Neurological Symptoms, Comorbidities, and Complications of COVID-19: A Literature Review and Meta-Analysis of Observational Studies. European Neurology, 2021, 84, 307-324.	1.4	23
27	The analysis of association between SNCA, HUSEYO and CSMD1 gene variants and Parkinson's disease in Iranian population. Neurological Sciences, 2016, 37, 731-736.	1.9	20
28	Copy Number Variants in Patients with Autism and Additional Clinical Features: Report of VIPR2 Duplication and a Novel Microduplication Syndrome. Molecular Neurobiology, 2017, 54, 7019-7027.	4.0	20
29	Biphasic effect of sumatriptan on PTZ-induced seizures in mice: Modulation by 5-HT1B/D receptors and NOS/NO pathway. European Journal of Pharmacology, 2018, 824, 140-147.	3.5	19
30	Functional improvement and immune-inflammatory cytokines profile of ischaemic stroke patients after treatment with boswellic acids: a randomized, double-blind, placebo-controlled, pilot trial. Inflammopharmacology, 2019, 27, 1101-1112.	3.9	19
31	Genetic Analysis of the <i>ZNF512B</i> , <i>SLC41A1,</i> and <i>ALDH2</i> Polymorphisms in Parkinson's Disease in the Iranian Population. Genetic Testing and Molecular Biomarkers, 2016, 20, 629-632.	0.7	18
32	Evaluating executive function in patients with temporal lobe epilepsy using the frontal assessment battery. Epilepsy Research, 2017, 133, 22-27.	1.6	18
33	Plasma Neurofilament Light Chain Levels Are Associated With Cortical Hypometabolism in Alzheimer Disease Signature Regions. Journal of Neuropathology and Experimental Neurology, 2019, 78, 709-716.	1.7	18
34	A novel <i>PUS7</i> mutation causes intellectual disability with autistic and aggressive behaviors. Neurology: Genetics, 2019, 5, e356.	1.9	18
35	Relationship between Sleep Quality and Quality of Life in Patients with Multiple Sclerosis. International Journal of Preventive Medicine, 2014, 5, 1582-6.	0.4	17
36	The omega-3 and Nano-curcumin effects on vascular cell adhesion molecule (VCAM) in episodic migraine patients: a randomized clinical trial. BMC Research Notes, 2021, 14, 283.	1.4	16

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37	SIPA1L2, MIR4697, GCH1 and VPS13C loci and risk of Parkinson's diseases in Iranian population: A case-control study. Journal of the Neurological Sciences, 2016, 369, 1-4.	0.6	15
38	Assessment of Factors Predicting Inadequate Pain Management in Chronic Pain Patients. Anesthesiology and Pain Medicine, 2019, In Press, e97229.	1.3	15
39	Attentional bias towards and away from fearful faces is modulated by developmental amygdala damage. Cortex, 2016, 81, 24-34.	2.4	14
40	Associations of Serum S100B and S100P With the Presence and Classification of Diabetic Peripheral Neuropathy in Adults With Type 2 Diabetes: A Case-Cohort Study. Canadian Journal of Diabetes, 2019, 43, 336-344.e2.	0.8	14
41	Molecular mechanisms of omega-3 fatty acids in the migraine headache. Iranian Journal of Neurology, 2017, 16, 210-217.	0.5	13
42	Granulocyte Colony-Stimulating Factor for Amyotrophic Lateral Sclerosis: A Randomized,		

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55	Evaluation of regulatory T lymphocytes and IL2Ra and FOXP3 gene expression in peripheral mononuclear cells from patients with amyotrophic lateral sclerosis. Irish Journal of Medical Science, 2018, 187, 1065-1071.	1.5	8
56	Conformational change and GTPase activity of human tubulin: A comparative study on Alzheimer's disease and healthy brain. Journal of Neurochemistry, 2020, 155, 207-224.	3.9	8
57	Phenotypic and genotypic characterization of families with complex intellectual disability identified pathogenic genetic variations in known and novel disease genes. Scientific Reports, 2020, 10, 968.	3.3	8
58	The Efficacy of Sphenopalatine Ganglion Block and Radiofrequency Denervation in the Treatment of Cluster Headache: A Case Series. Anesthesiology and Pain Medicine, 2020, 10, e104466.	1.3	8
59	Investigating the possible association between <scp>NLRP3</scp> gene polymorphisms and myasthenia gravis. Muscle and Nerve, 2021, 63, 730-736.	2.2	7
60	<scp><i>ANXA1</i></scp> with Antiâ€Inflammatory Properties Might Contribute to Parkinsonism. Annals of Neurology, 2021, 90, 319-323.	5.3	7
61	PRDM12 Is Transcriptionally Active and Required for Nociceptor Function Throughout Life. Frontiers in Molecular Neuroscience, 2021, 14, 720973.	2.9	7
62	The Influence of Chronic Pain on Number Sense and Numeric Rating Scale: A prospective Cohort Study. Anesthesiology and Pain Medicine, 2020, 10, e103532.	1.3	7
63	Clinical and molecular spectrum of $P/Q$ type calcium channel Cav2.1 in epileptic patients. Orphanet Journal of Rare Diseases, 2021, 16, 461.	2.7	7
64	Analysis of Copy Number Variations in Patients with Autism Using Cytogenetic and MLPA Techniques: Report of 16p13.1p13.3 and 10q26.3 Duplications. International Journal of Molecular and Cellular Medicine, 2016, 5, 236-245.	1.1	7
65	A Case Based-Shared Teaching Approach in Undergraduate Medical Curriculum: A Way for Integration in Basic and Clinical Sciences. Acta Medica Iranica, 2017, 55, 259-264.	0.8	7
66	Extraintestinal Involvement of Rotavirus Infection in Children. Archives of Iranian Medicine, 2015, 18, 604-5.	0.6	7
67	Potential diagnostic value of 131 I-MIBG myocardial scintigraphy in discrimination between Alzheimer disease and dementia with Lewy bodies. Clinical Neurology and Neurosurgery, 2017, 163, 163-166.	1.4	6
68	Molecular characterization of <i>PRKN</i> structural variations identified through wholeâ€genome sequencing. Molecular Genetics & Enough Community (Senomic Medicine, 2018, 6, 1243-1248.	1.2	6
69	Paracetamol 325Âmg/tramadol 37.5Âmg effect on pain during needle electromyography: a double-blind crossover clinical trial. Acta Neurologica Belgica, 2016, 116, 599-604.	1.1	5
70	Tongue Protrusion Dystonia in Pantothenate Kinase-Associated Neurodegeneration. Pediatric Neurology, 2020, 103, 76-78.	2.1	5
71	Safety and efficacy of memantine for multiple sclerosis-related fatigue: A pilot randomized, double-blind placebo-controlled trial. Journal of the Neurological Sciences, 2020, 414, 116844.	0.6	5
72	Cognitive Impairment in Opium Use Disorder. Behavioural Neurology, 2021, 2021, 1-12.	2.1	5

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73	From a simple chronic headache to neurobrucellosis: a case report. Medical Journal of the Islamic Republic of Iran, 2014, 28, 12.	0.9	5
74	Idiopathic superficial siderosis of the central nervous system. Cerebellum and Ataxias, 2021, 8, 9.	1.9	4
75	The phenotypic spectrum of PCDH12 associated disorders - Five new cases and review of the literature. European Journal of Paediatric Neurology, 2022, 36, 7-13.	1.6	4
76	Saffron () versus duloxetine for treatment of patients with fibromyalgia: A randomized double-blind clinical trial. Avicenna Journal of Phytomedicine, 2018, 8, 513-523.	0.2	4
77	Mutation in TWINKLE in a Large Iranian Family with Progressive External Ophthalmoplegia, Myopathy, Dysphagia and Dysphonia, and Behavior Change. Archives of Iranian Medicine, 2016, 19, 87-91.	0.6	4
78	Treatment of Myoclonus–dystonia with carbamazepine. Parkinsonism and Related Disorders, 2018, 53, 116-117.	2.2	3
79	Evaluating Executive Functions in Patients with Juvenile Myoclonic Epilepsy Using Frontal Assessment Battery. Behavioural Neurology, 2020, 2020, 1-10.	2.1	3
80	Recessive <i>COL4A2</i> Mutation Leads to Intellectual Disability, Epilepsy, and Spastic Cerebral Palsy. Neurology: Genetics, 2021, 7, e583.	1.9	3
81	An interictal measurement of cerebral oxygen extraction fraction in MRI-negative refractory epilepsy using quantitative susceptibility mapping. Physica Medica, 2021, 85, 87-97.	0.7	3
82	Emery-Dreifuss Muscular Dystrophy: a Report of a Large Family with 11 Affected Individuals. International Journal of Molecular and Cellular Medicine, 2016, 5, 196-198.	1.1	3
83	A systematic review of resting-state and task-based fmri in juvenile myoclonic epilepsy. Brain Imaging and Behavior, 2022, 16, 1465-1494.	2.1	3
84	Adie's pupil during migraine attack: case report and review of literature. Acta Neurologica Belgica, 2011, 111, 66-8.	1.1	3
85	Homozygous Mutation in TWNK Cases Ataxia, Sensorineural Hearing Loss and Optic Nerve Atrophy. Archives of Iranian Medicine, 2019, 22, 728-730.	0.6	3
86	Clinical use of 99mTc-HMPAO-labeled platelets in cerebral sinus thrombosis imaging. Acta Neurologica Belgica, 2019, 119, 549-553.	1.1	2
87	An Unusual Location for Sphenopalatine Ganglion in the Pterygopalatine Fossa Which May Facilitate Radiofrequency Neurolysis: A Case Report. Archives of Neuroscience, 2016, 3, .	0.3	2
88	Nutritional Aspects of Treatment in Epileptic Patients. Iranian Journal of Child Neurology, 2016, 10, 1-12.	0.3	2
89	Leu226Trp CACNA1A variant associated with juvenile myoclonic epilepsy with and without intellectual disability. Clinical Neurology and Neurosurgery, 2022, 213, 107108.	1.4	2
90	Evaluation of partial epilepsy in Iran: role of video-EEG, EEG, and MRI with epilepsy protocol. Iranian Journal of Neurology, 2011, 10, 9-15.	0.5	1

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91	Neglected Alkaptonuric Patient Presenting with Steppage Gait. Archives of Bone and Joint Surgery, 2016, 4, 188-91.	0.2	1
92	Molecular changes in obese and depressive patients are similar to neurodegenerative disorders. Iranian Journal of Neurology, $2017, 16, 192-200$ .	0.5	1
93	Plasma Cytokines Profile in Subjects with Alzheimer's Disease: Interleukin 1 Alpha as a Candidate for Target Therapy. Galen, 2021, 10, e1974.	0.6	1
94	Seroprevalence of NMO-IgG Antibody in Neuromyelitis optica (NMO) and Its Specificity in Differentiating NMO from Other Demyelinating Diseases with Overlap Symptoms: An Iranian Experience. Iranian Journal of Allergy, Asthma and Immunology, 2015, 14, 98-104.	0.4	1
95	99m Technetium-HMPAO-labeled platelet scan in practice: Preparation, quality control, and biodistribution studies. Brazilian Journal of Pharmaceutical Sciences, 0, 58, .	1.2	1
96	Boswellia serrata extract shows cognitive benefits in a double-blind, randomized, placebo-controlled pilot clinical trial in individuals who suffered traumatic brain injury. Brain Injury, 2022, , 1-7.	1.2	1
97	Curable episodic ataxia? History taking versus advanced diagnostic instruments. Journal of the Neurological Sciences, 2014, 336, 295.	0.6	O
98	Novel <i>ABCD1</i> gene mutations in Iranian pedigrees with X-linked adrenoleukodystrophy. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 1207-1215.	0.9	0
99	Seizure-Control Effect of Levatiracetam on Juvenile Myoclonic Epilepsy and Other Epileptic Syndromes: Literature Review of Recent Studies. Iranian Journal of Child Neurology, 2015, 9, 1-8.	0.3	0
100	Hashimoto Encephalopathy in Case of Progressive Cognitive Impairment; a Case Report. Emergency, 2014, 2, 144-6.	0.6	0
101	Imaging of Clot by Tc-HMPAO Labeled Platelet in Animal Model Induced Thrombosis. Iranian Journal of Pharmaceutical Research, 2020, 19, 76-84.	0.5	0
102	The Effect of Oral Simvastatin on the Clinical Outcome of Patients with Severe Traumatic Brain Injury: A Randomized Clinical Trial. Ethiopian Journal of Health Sciences, 2021, 31, 807-816.	0.4	0
103	The Phenotypic Spectrum of PCDH12-Associated Disorders: Five New Cases and Review of the Literature. , 2021, 52, .		0
104	Psychometric evaluation of Persian version of Seizure Severity Questionnaire. Epilepsy and Behavior, 2022, 128, 108506.	1.7	0
105	Clinical, Laboratory and Imaging Characteristics of Hospitalized COVID-19 Patients with Neurologic Involvement; a Cross-Sectional Study Archives of Academic Emergency Medicine, 2022, 10, e10.	0.4	0
106	Association of Depression with Migraine without Aura; A Cross-Sectional Study , 2019, 16, 182-186.		0