Hussein A Algahtani

List of Publications by Year
in descending order

119	894	566801	580395
papers	citations	15 h-index	25 g-index

122	122	122	1559
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Hyperferritinemia with iron deposition in the basal ganglia and tremor as the initial manifestation of follicular lymphoma. International Journal of Neuroscience, 2023, 133, 896-900.	0.8	1
2	Autosomal Recessive Spastic Paraplegia Type 78 Associated with a Homozygous Variant in the <scp>ATP13A2</scp> Gene. Movement Disorders Clinical Practice, 2022, 9, 997-1002.	0.8	1
3	Perception of the general population towards migraine in Jeddah, Saudi Arabia. Egyptian Journal of Neurology, Psychiatry and Neurosurgery, 2022, 58, .	0.4	3
4	Botulinum toxin injection in the management of chronic migraine: the Saudi experience with a proposal for a new protocol. Acta Neurologica Belgica, 2021, 121, 1783-1787.	0.5	0
5	A novel mutation in ATM gene in a Saudi female with ataxia telangiectasia. International Journal of Neuroscience, 2021, 131, 206-211.	0.8	2
6	Perceptions and attitudes of different healthcare professionals and students toward interprofessional education in Saudi Arabia: a cross-sectional survey. Journal of Interprofessional Care, 2021, 35, 476-481.	0.8	10
7	A Novel Variant in CWF19L1 Gene in a Family with Late-Onset Autosomal Recessive Cerebellar Ataxia 17. Neurological Research, 2021, 43, 141-147.	0.6	6
8	Decompressive craniectomy as a lifesaving intervention for acute disseminated encephalomyelitis (ADEM). Multiple Sclerosis and Related Disorders, 2021, 47, 102612.	0.9	1
9	Clinical and Radiological Characteristics of Neuromyelitis Optica Spectrum Disorder: The Experience from Saudi Arabia. Multiple Sclerosis and Related Disorders, 2021, 47, 102668.	0.9	3
10	Autosomal recessive cerebellar ataxia with spasticity due to a rare mutation in GBA2 gene in a large consanguineous Saudi family. Genes and Diseases, 2021, 8, 110-114.	1.5	2
11	Contactin-associated protein-like 2 (CASPR2) autoantibody-related pancerebellar syndrome. Acta Neurologica Belgica, 2021, , 1.	0.5	1
12	Mills' syndrome: Reporting the disease course with a monthly intravenous immunoglobulin program. Journal of Neuroimmunology, 2021, 355, 577562.	1.1	1
13	Multiple Sclerosis in Saudi Arabia: Clinical, Social, and Psychological Aspects of the Disease. Cureus, 2021, 13, e16484.	0.2	1
14	Vaccine-Induced Immune Thrombotic Thrombocytopenia with Disseminated Intravascular Coagulation and Death following the ChAdOx1 nCoV-19 Vaccine. Journal of Stroke and Cerebrovascular Diseases, 2021, 30, 105938.	0.7	32
15	Novel Heterozygous Variants in the HLA-DRB1 Gene in a Saudi Family with Early-Onset Familial Multiple Sclerosis: Therapeutic Failure and Success. International Journal of MS Care, 2021, , .	0.4	0
16	Prevalence of Pediatric Onset Multiple Sclerosis in Saudi Arabia. Multiple Sclerosis International, 2021, 2021, 1-6.	0.4	3
17	Cocaine-induced massive ischemic stroke treated by decompressive craniectomy with favorable outcome. Journal of Innovative Optical Health Sciences, 2021, 16, 830.	0.5	1
18	Clinical Presentation, Management, and Outcome in Patients With Myasthenia Gravis: A Retrospective Study From Two Tertiary Care Centers in Saudi Arabia. Cureus, 2021, 13, e20765.	0.2	1

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19	Chronic progressive external ophthalmoplegia in a Saudi patient with a mutation in the POLG gene successfully managed with bilateral frontalis sling. Journal of Genetic Medicine, 2021, 18, 121-126.	0.1	0
20	Perception and attitude of the general population towards Alzheimerâ€™s disease in Jeddah, Saudi Arabia. Acta Neurologica Belgica, 2020, 120, 313-320.	0.5	19
21	Familial aggregation of multiple sclerosis: Results from the national registry of the disease in Saudi Arabia. Multiple Sclerosis Journal - Experimental, Translational and Clinical, 2020, 6, 205521732096049.	0.5	5
22	Amyotrophic lateral sclerosis care in Saudi Arabia: A survey of providersâ€™ perceptions. Brain and Behavior, 2020, 10, e01795.	1.0	1
23	The spectrum of muscle pathologies: Three decades of experience from a reference laboratory in Saudi Arabia. Annals of Diagnostic Pathology, 2020, 47, 151532.	0.6	2
24	A Novel Heterozygous Variant in Exon 19 of NOTCH3 in a Saudi Family with Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy. Journal of Stroke and Cerebrovascular Diseases, 2020, 29, 104832.	0.7	2
25	Concurrence of multiple sclerosis, oligodendroglioma, and autosomal recessive cerebellar ataxia with spasticity in the same patient: A challenging diagnosis. Multiple Sclerosis and Related Disorders, 2020, 40, 101945.	0.9	3
26	Are Repeated Cycles of Intravenous Immunoglobulin Justified in Patients With Poorly Responsive Guillain-BarrÃ© Syndrome?. Neurohospitalist, The, 2020, 10, 224-228.	0.3	1
27	Assessment of the Burden of Multiple Sclerosis Patients' Caregivers in Saudi Arabia. Cureus, 2020, 12, e6658.	0.2	5
28	A Novel Intronic Variant in SLC2A1 Gene in a Saudi Patient with Myoclonic Epilepsy. Journal of Epilepsy Research, 2020, 10, 40-43.	0.1	0
29	A novel variant c.3706C>T p.(Avg 1236Cys) in the <i>ABCA7</i> gene in a Saudi patient with susceptibility to Alzheimer's disease 9. Intractable and Rare Diseases Research, 2020, 9, 151-155.	0.3	1
30	Hashimotoâ€™s encephalopathy presenting with progressive cerebellar ataxia. Neurosciences, 2020, 25, 327-327.	0.1	0
31	Unusual presentation of basilar artery thrombosis. Journal of Cerebrovascular and Endovascular Neurosurgery, 2020, 22, 282-286.	0.2	1
32	Irreversible Cerebellar Atrophy as a Complication of Short-Term Phenytoin Exposure: Clinical Improvement Following Discontinuation of the Culprit. Journal of Epilepsy Research, 2020, 10, 96-99.	0.1	4
33	Effectiveness and Needs Assessment of Faculty Development Programme for Medical Education: Experience from Saudi Arabia. Sultan Qaboos University Medical Journal, 2020, 20, 83.	0.3	14
34	Factors to be considered in designing a faculty development program for medical education: local experience from the Western region of Saudi Arabia. Yeungnam University Journal of Medicine, 2020, 37, 210-216.	0.7	3
35	Carotid Artery Angioplasty and Stenting for Carotid Stenosis: A Single-Center Experience from Saudi Arabia. Neurointervention, 2020, 15, 133-139.	0.5	1
36	Epilepsy and driving: Local experience from Saudi Arabia. Epilepsy and Behavior, 2019, 99, 106401.	0.9	5

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37	A very rare form of autosomal dominant progressive myoclonus epilepsy caused by a novel variant in the PRICKLE1 gene. Seizure: the Journal of the British Epilepsy Association, 2019, 69, 133-139.	0.9	10
38	A novel mutation in CACNA1A gene in a Saudi female with episodic ataxia type 2 with no response to acetazolamide or 4-aminopyridine. Intractable and Rare Diseases Research, 2019, 8, 67-71.	0.3	6
39	Erythema nodosum in a patient with multiple sclerosis on dimethyl fumarate. Multiple Sclerosis and Related Disorders, 2019, 28, 155-158.	0.9	4
40	Perception and Attitude of the General Population towards Epilepsy in Jeddah, Saudi Arabia. Journal of Epilepsy Research, 2019, 9, 42-50.	0.1	18
41	Antiepileptic Drugs Usage in Pregnant Women with Epilepsy in Saudi Arabia. Journal of Epilepsy Research, 2019, 9, 134-138.	0.1	2
42	Unilateral headache and convulsive-like movements as a manifestation of pontine warning syndrome. Journal of King Abdulaziz University, Islamic Economics, 2019, 24, 332-334.	0.5	1
43	Hashimotoâ€™s Encephalopathy Presenting with Progressive Cerebellar Ataxia. Journal of King Abdulaziz University, Islamic Economics, 2019, 24, 315-319.	0.5	6
44	Unethical human research in the field of neuroscience: a historical review. Neurological Sciences, 2018, 39, 829-834.	0.9	12
45	Transverse myelitis-like presentation of methanol intoxication: A case report and review of the literature. Journal of Spinal Cord Medicine, 2018, 41, 72-76.	0.7	5
46	Urea cycle disorder misdiagnosed as multiple sclerosis: a case report and review of the literature. Neuroradiology Journal, 2018, 31, 213-217.	0.6	8
47	Susac syndrome misdiagnosed as multiple sclerosis with exacerbation by interferon beta therapy. Neuroradiology Journal, 2018, 31, 207-212.	0.6	21
48	Teratoma of the nervous system: A case series. Neurocirugia, 2018, 29, 143-149.	0.2	3
49	Fatal serotonin syndrome in a patient with Marchiafavaâ€“Bignami disease: Combined neurological and psychiatric emergency. Neurologia I Neurochirurgia Polska, 2018, 52, 277-280.	0.6	4
50	Neurological recovery from multiple cardiac arrests due to acute massive pulmonary embolism managed by cardiopulmonary resuscitation and extracorporeal membrane oxygenation. Cardiovascular Revascularization Medicine, 2018, 19, 120-122.	0.3	2
51	Ataxia with ocular apraxia type 2 not responding to 4-aminopyridine: A rare mutation in the <i>SETX</i> gene in a Saudi patient. Intractable and Rare Diseases Research, 2018, 7, 275-279.	0.3	4
52	Adult-onset hemophagocytic lymphohistiocytosis type 2 presenting as a demyelinating disease. Multiple Sclerosis and Related Disorders, 2018, 25, 77-82.	0.9	13
53	Limb-girdle muscular dystrophy type 2B: An unusual cause of proximal muscular weakness in Saudi Arabia. Journal of Back and Musculoskeletal Rehabilitation, 2018, 31, 999-1004.	0.4	9
54	Vertebral Artery Occlusion Causing Facial Colliculus and Opalski Stroke Syndromes Simultaneously. Neurologist, 2018, 23, 100-103.	0.4	1

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55	Post-partum headache caused by dual pathology: A message to the anesthetist. Journal of King Abdulaziz University, Islamic Economics, 2018, 23, 262-264.	0.5	1
56	Chromosomal Micro-aberration in a Saudi Family with Juvenile Myoclonic Epilepsy. CNS and Neurological Disorders - Drug Targets, 2018, 16, 1010-1017.	0.8	7
57	De novo Craniopharyngioma of the fourth ventricle: Case report and review of literature. Journal of Innovative Optical Health Sciences, 2018, 13, 62-65.	0.5	6
58	Vogt Koyanagi Harada Syndrome mimicking multiple sclerosis: A case report and review of the literature. Multiple Sclerosis and Related Disorders, 2017, 12, 44-48.	0.9	12
59	Tumefactive demyelinating lesions: A comprehensive review. Multiple Sclerosis and Related Disorders, 2017, 14, 72-79.	0.9	87
60	Biotinâ€“thiamineâ€“responsive basal ganglia disease: catastrophic consequences of delay in diagnosis and treatment. Neurological Research, 2017, 39, 117-125.	0.6	21
61	Posterior Reversible Encephalopathy Syndrome. Neurohospitalist, The, 2017, 7, 24-29.	0.3	5
62	Cough syncope induced by post nasal drip successfully managed by Gabapentin. Respiratory Medicine Case Reports, 2017, 22, 47-50.	0.2	0
63	Autosomal Recessive Cerebellar Ataxia type 1 mimicking multiple sclerosis: A report of two siblings with a novel mutation in SYNE1 gene in a Saudi family. Journal of the Neurological Sciences, 2017, 372, 97-100.	0.3	20
64	Nosocomial herpes simplex encephalitis: A challenging diagnosis. Journal of Infection and Public Health, 2017, 10, 343-347.	1.9	3
65	Quality of life among multiple sclerosis patients in Saudi Arabia. Journal of King Abdulaziz University, Islamic Economics, 2017, 22, 261-266.	0.5	22
66	Occupational Neurobrucellosis Mimicking a Brain Tumor: A Case Report and Review of the Literature. Case Reports in Infectious Diseases, 2017, 2017, 1-5.	0.2	9
67	Idiopathic Harlequin Syndrome Manifesting during Exercise: A Case Report and Review of the Literature. Case Reports in Medicine, 2017, 2017, 1-3.	0.3	12
68	More Than a Decade of Misdiagnosis of Alternating Hemiplegia of Childhood with Catastrophic Outcome. Case Reports in Medicine, 2017, 2017, 1-5.	0.3	8
69	Neurology Research in Saudi Arabia: Urgent call for action. Sultan Qaboos University Medical Journal, 2017, 17, e324-328.	0.3	4
70	Perception of hospital accreditation among health professionals in Saudi Arabia. Annals of Saudi Medicine, 2017, 37, 326-332.	0.5	31
71	Levetiracetam-Induced Skin Hyperpigmentation: An Extremely Rare Undesirable Side Effect. Journal of Epilepsy Research, 2017, 7, 106-108.	0.1	11
72	Desmoplastic infantile astrocytoma and ganglioglioma: case report and review of the literature. , 2017, 36, 31-40.		11

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73	A novel mutation in the DNMT1 gene in a patient presenting with pure cerebellar ataxia. Journal of Genetic Medicine, 2017, 14, 71-74.	0.1	3
74	Neurological Manifestations of Acute Posterior Multifocal Placoid Pigment Epitheliopathy. Journal		

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91	Dialysis Dementia:. Saudi Endodontic Journal, 2014, 4, 29-35.	0.0	0
92	Cerebral venous sinus thrombosis. Neurosciences, 2014, 19, 11-6.	0.1	9
93	Acute paraplegia caused by Schistosoma mansoni. Neurosciences, 2014, 19, 47-51.	0.1	3
94	Crossed cerebro-cerebellar atrophy with Dyke Davidoff Masson syndrome. Neurosciences, 2014, 19, 52-5.	0.1	5
95	Memory boosting effect of Citrus limon, Pomegranate and their combinations. Pakistan Journal of Pharmaceutical Sciences, 2014, 27, 1837-40.	0.2	5
96	Suprasellar Meningioma Causing Papilledema and Hallucinations. Saudi Endodontic Journal, 2013, 3, 43-45.	0.0	0
97	Luminaries in Medicine - Ibn al-Nafis. Saudi Endodontic Journal, 2013, 3, 7.	0.0	0
98	Suprasellar Meningioma Causing Papilledema and Hallucinations. Saudi Endodontic Journal, 2013, 3, 43-45.	0.0	0
99	Luminaries in Medicine - Ibn al-Nafis. Saudi Endodontic Journal, 2013, 3, 7.	0.0	0
100	Posterior fossa teratoma. Neurosciences, 2013, 18, 371-4.	0.1	5
101	Reversible Parkinsonism caused by deep cerebral venous sinus thrombosis. Neurosciences, 2013, 18, 378-81.	0.1	3
102	Intractable Seizures Caused by a Rare Treatable Condition. Saudi Endodontic Journal, 2012, 2, 33-37.	0.0	0
103	A Young Patient with Leukemia and New-Onset Headache. Saudi Endodontic Journal, 2012, 2, 39-44.	0.0	0
104	Sir William Osler:. Saudi Endodontic Journal, 2012, 2, 7.	0.0	0
105	Sir William Osler:. Saudi Endodontic Journal, 2012, 2, 7.	0.0	0
106	Intractable Seizures Caused by a Rare Treatable Condition. Saudi Endodontic Journal, 2012, 2, 33-37.	0.0	0
107	A Young Patient with Leukemia and New-Onset Headache. Saudi Endodontic Journal, 2012, 2, 39-44.	0.0	0
108	Delirium. A comprehensive review. Neurosciences, 2012, 17, 205-12.	0.1	2

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109	A middle aged man with respiratory failure and weakness. Neurosciences, 2012, 17, 274-5.	0.1	0
110	The Status of Acute Stroke Care in Saudi Arabia: An Urgent Call for Action!. International Journal of Stroke, 2011, 6, 75-76.	2.9	25
111	A young women who couldn't see or move. Neurosciences, 2011, 16, 87-8.	0.1	0
112	A woman with seizures and papilledema. Neurosciences, 2011, 16, 295-7.	0.1	0
113	Cerebral venous sinus thrombosis in Saudi Arabia. Neurosciences, 2011, 16, 329-34.	0.1	19
114	Inability to walk due to scurvy: a forgotten disease. Annals of Saudi Medicine, 2010, 30, 325-328.	0.5	24
115	Guillain-Barre syndrome following cardiac surgery. Difficult diagnosis in the intensive care unit. Neurosciences, 2009, 14, 374-8.	0.1	15
116	Central pontine myelinolysis due to rapid correction of hyponatremia induced by excessive water intake. Neurosciences, 2008, 13, 296-8.	0.1	1
117	Sporadic creutzfeldt jacob disease. Neurosciences, 2008, 13, 430-2.	0.1	0
118	Idiopathic intracranial hypertension. Atypical presentation. Journal of King Abdulaziz University, Islamic Economics, 2007, 28, 762-5.	0.5	4
119	Bilateral Femoral Neuropathy Complicating Rhabdomyolysis and Acute Renal Failure. Journal of Clinical Neuromuscular Disease, 2005, 6, 153-156.	0.3	3