Carmel Armon

List of Publications by Year
in descending order

33 papers	1,204 citations	759233 12 h-index	552781 26 g-index
34 all docs	34 docs citations	34 times ranked	1419 citing authors

#	Article	IF	Citations
1	Reasons for delayed treatment initiation in Guillain-Barre syndrome. Journal of the Neurological Sciences, 2022, 434, 120179.	0.6	3
2	Shortâ€•and longâ€term outcome and predictors in an international cohort of patients with neuroâ€COVIDâ€19. European Journal of Neurology, 2022, 29, 1663-1684.	3.3	18
3	Oral and Topical Treatment of Painful Diabetic Polyneuropathy: Practice Guideline Update Summary. Neurology, 2022, 98, 31-43.	1.1	64
4	Estimating the X chromosome-mediated risk for developing Alzheimerâ€™s disease. Journal of Neurology, 2021, , 1.	3.6	0
5	Validation of MRI biomarker of white matter degeneration for ALS clinical trials. Neurology, 2020, 95, 327-328.	1.1	0
6	High BMI is associated with low ALS risk. Neurology, 2019, 93, 189-191.	1.1	2
7	Theme 1 Epidemiology and informatics. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 101-113.	1.7	0
8	Smoking is a cause of ALS. High LDLâ€cholesterol levels? Unsure. Annals of Neurology, 2019, 85, 465.	5.3	4
9	CT-guided thrombolytic treatment of patients with wake-up strokes. ENeurologicalSci, 2019, 14, 91-97.	1.3	9
10	Smoking is a cause of amyotrophic lateral sclerosis. High lowâ€density lipoprotein cholesterol levels? Unsure. Annals of Neurology, 2019, 85, 465-469.	5.3	8
11	Intrinsic race differences in ALS survival in a US clinic population independent of ventilation. Neurology, 2019, 92, 781-783.	1.1	1
12	Ethics of clinical research in patients with ALS: is there a risk of exploitation?. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 161-166.	1.7	2
13	From Snow to Hill to ALS: An epidemiological odyssey in search of ALS causation. Journal of the Neurological Sciences, 2018, 391, 134-140.	0.6	16
14	The beginning of precision medicine in ALS?. Neurology, 2017, 89, 1850-1851.	1.1	5
15	Accrued somatic mutations (nucleic acid changes) trigger ALS: 2005-2015 update. Muscle and Nerve, 2016, 53, 842-849.	2.2	12
16	Effect of the 2013 AHA/ASA guidelines on TPA use in acute ischemic stroke at Assaf Harofeh Medical Center in Israel. Journal of the Neurological Sciences, 2016, 369, 306-309.	0.6	3
17	A blow to the head traumaâ€“ALS hypothesis. Neurology, 2015, 84, 1728-1729.	1.1	0
18	The underestimation of familial ALS and counseling patients with sporadic ALS. Neurology, 2014, 82, 13-14.	1.1	2

#	Article	IF	Citations
19	Is head trauma a risk factor for amyotrophic lateral sclerosis? An evidence based review. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 351-356.	2.1	42
20	Smoking may be considered an established risk factor for sporadic ALS. Neurology, 2009, 73, 1693-1698.	1.1	136
21	Three drawers. Neurology, 2008, 70, 2347-2347.	1.1	0
22	From clues to mechanisms. Neurology, 2008, 71, 872-873.	1.1	16
23	Assessment: Use of epidural steroid injections to treat radicular lumbosacral pain: Report of the Therapeutics and Technology Assessment Subcommittee of the American Academy of Neurology. Neurology, 2007, 68, 723-729.	1.1	198
24	Sports and trauma in amyotrophic lateral sclerosis revisited. Journal of the Neurological Sciences, 2007, 262, 45-53.	0.6	80
25	Acquired nucleic acid changes may trigger sporadic amyotrophic lateral sclerosis. Muscle and Nerve, 2005, 32, 373-377.	2.2	19
26	Addendum to assessment: Prevention of postâ€“lumbar puncture headaches [RETIRED]. Neurology, 2005, 65, 510-512.	1.1	149
27	Chapter 7 Epidemiology of Amyotrophic Lateral Sclerosis/Motor Neuron Disease. Blue Books of Practical Neurology, 2003, 28, 167-205.	0.1	12
28	An Evidence-Based Medicine Approach to the Evaluation of the Role of Exogenous Risk Factors in Sporadic Amyotrophic Lateral Sclerosis. Neuroepidemiology, 2003, 22, 217-228.	2.3	186
29	Limitations of inferences from observational databases in amyotrophic lateral sclerosis: all that glitters is not gold. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases, 2002, 3, 109-112.	1.2	9
30	Linear estimates of disease progression predict survival in patients with amyotrophic lateral sclerosis. Muscle and Nerve, 2000, 23, 874-882.	2.2	59
31	Motor unit number estimate-based rates of progression of ALS predict patient survival. , 1999, 22, 1571-1575.		79
32	Motor unit number estimates and quantitative muscle strength measurements of distal muscles in patients with amyotrophic lateral sclerosis. , 1997, 20, 499-501.		21
33	Mechanical trauma as a risk factor in classic amyotrophic lateral sclerosis: Lack of epidemiologic evidence. Journal of the Neurological Sciences, 1992, 113, 133-143.	0.6	49