Hakan I GÃ¼rvit

List of Publications by Year
in descending order

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163 papers	4,572 citations	136950 32 h-index	114465 63 g-index
176 all docs	176 docs citations	176 times ranked	7517 citing authors

#	Article	IF	Citations
1	TREM2 mutations implicated in neurodegeneration impair cell surface transport and phagocytosis. Science Translational Medicine, 2014, 6, 243ra86.	12.4	600
2	Using Exome Sequencing to Reveal Mutations in TREM2 Presenting as a Frontotemporal Dementiaâ€“like Syndrome Without Bone Involvement. JAMA Neurology, 2013, 70, 78.	9.0	311
3	<scp>EFNSâ€ENS</scp> Guidelines on the diagnosis and management of disorders associated with dementia. European Journal of Neurology, 2012, 19, 1159-1179.	3.3	239
4	The interleukin 1 alpha, interleukin 1 beta, interleukin 6 and alpha-2-macroglobulin serum levels in patients with early or late onset Alzheimer's disease, mild cognitive impairment or Parkinson's disease. Journal of Neuroimmunology, 2015, 283, 50-57.	2.3	197
5	Comparative analysis of event-related potentials during Go/NoGo and CPT: Decomposition of electrophysiological markers of response inhibition and sustained attention. Brain Research, 2006, 1104, 114-128.	2.2	182
6	BDNF, TNFÎ±, HSP90, CFH, and IL-10 Serum Levels in Patients with Early or Late Onset Alzheimer's Disease or Mild Cognitive Impairment. Journal of Alzheimer's Disease, 2013, 37, 185-195.	2.6	152
7	Association between Vitamin D Receptor Gene Polymorphism and Alzheimer's Disease. Tohoku Journal of Experimental Medicine, 2007, 212, 275-282.	1.2	134
8	Neuropsychological Patterns and Language Deficits in 20 Consecutive Cases of Autopsy-Confirmed Alzheimer's Disease. Archives of Neurology, 1993, 50, 931-937.	4.5	133
9	Cognitive impairment in amyotrophic lateral sclerosis: evidence from neuropsychological investigation and event-related potentials. Cognitive Brain Research, 2002, 14, 234-244.	3.0	116
10	A novel compound heterozygous mutation in TREM2 found in a Turkish frontotemporal dementia-like family. Neurobiology of Aging, 2013, 34, 2890.e1-2890.e5.	3.1	113
11	Seven-Year Follow-up of Neurologic Involvement in Behcet Syndrome. Archives of Neurology, 1996, 53, 691-694.	4.5	103
12	The effects of rasagiline on cognitive deficits in Parkinson's disease patients without dementia: A randomized, doubleâ€blind, placeboâ€controlled, multicenter study. Movement Disorders, 2011, 26, 1851-1858.	3.9	97
13	Neuropsychological follow-up of 12 patients with neuro-BehÃ§et disease. Journal of Neurology, 1999, 246, 113-119.	3.6	90
14	Exome sequencing reveals an unexpected genetic cause of disease: NOTCH3 mutation in a Turkish family with Alzheimer's disease. Neurobiology of Aging, 2012, 33, 1008.e17-1008.e23.	3.1	86
15	Pain is common in Parkinson's disease. Clinical Neurology and Neurosurgery, 2011, 113, 11-13.	1.4	78
16	The distinct genetic pattern of ALS in Turkey and novel mutations. Neurobiology of Aging, 2015, 36, 1764.e9-1764.e18.	3.1	78
17	Vitamin D Receptor Gene Haplotype Is Associated with Late-Onset Alzheimer's Disease. Tohoku Journal of Experimental Medicine, 2012, 228, 189-196.	1.2	77
18	Spontaneous dissection of the extracranial vertebral artery with spinal subarachnoid haemorrhage in a patient with Behï¿½et's disease. Neuroradiology, 1993, 35, 352-354.	2.2	76

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19	Neuropsychological function in obsessive-compulsive disorder. Comprehensive Psychiatry, 2012, 53, 167-175.	3.1	59
20	A new Fâ€box protein 7 gene mutation causing typical Parkinson's disease. Movement Disorders, 2015, 30, 1130-1133.	3.9	59
21	Paroxysmal dysarthria and ataxia in a patient with Behï¿½et's disease. Journal of Neurology, 1995, 242, 344-347.	3.6	54
22	Inflammatory/demyelinating central nervous system involvement in familial Mediterranean fever (FMF): coincidence or association?. Journal of Neurology, 2006, 253, 928-934.	3.6	53
23	The Arg194Trp polymorphism in DNA repair gene XRCC1 and the risk for sporadic late-onset Alzheimer's disease. Neurological Sciences, 2007, 28, 31-34.	1.9	48
24	Single nucleotide polymorphisms in base-excision repair genes hOGG1, APE1 and XRCC1 do not alter risk of Alzheimer's disease. Neuroscience Letters, 2008, 442, 287-291.	2.1	48
25	Risk Factors for Alzheimer Disease: A Population-Based Case-Control Study in Istanbul, Turkey. Alzheimer Disease and Associated Disorders, 2003, 17, 139-145.	1.3	47
26	Identification of PSEN1 and PSEN2 gene mutations and variants in Turkish dementia patients. Neurobiology of Aging, 2012, 33, 1850.e17-1850.e27.	3.1	44
27	The combinations of TNFÎ±-308 and IL-6 -174 or IL-10 -1082 genes polymorphisms suggest an association with susceptibility to sporadic late-onset Alzheimerâ€™s disease. Acta Neurologica Scandinavica, 2009, 120, 396-401.	2.1	42
28	A composite neural network model for perseveration and distractibility in the Wisconsin card sorting test. Neural Networks, 2006, 19, 375-387.	5.9	41
29	GC and VDR SNPs and Vitamin D Levels in Parkinsonâ€™s Disease: The Relevance to Clinical Features. NeuroMolecular Medicine, 2017, 19, 24-40.	3.4	38
30	The Central Biobank and Virtual Biobank of BIOMARKAPD: A Resource for Studies on Neurodegenerative Diseases. Frontiers in Neurology, 2015, 6, 216.	2.4	36
31	Vitamin D deficiency might pose a greater risk for ApoEÉ›4 non-carrier Alzheimerâ€™s disease patients. Neurological Sciences, 2016, 37, 1633-1643.	1.9	36
32	Clinical and magnetic resonance imaging findings of HIV-Negative patients with neurosyphilis. Journal of Neurology, 2007, 254, 368-374.	3.6	34
33	Salience network engagement with the detection of morally laden information. Social Cognitive and Affective Neuroscience, 2017, 12, 1118-1127.	3.0	30
34	Unrecognized depression in community-dwelling elderly persons in Istanbul. International Psychogeriatrics, 2005, 17, 303-312.	1.0	29
35	Paraneoplastic limbic encephalitis with immature ovarian teratoma--a case report. Journal of Neuro-Oncology, 1998, 37, 63-66.	2.9	28
36	An assessment of Movement Disorder Society Task Force diagnostic criteria for mild cognitive impairment in Parkinson's disease. European Journal of Neurology, 2018, 25, 148-153.	3.3	28

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37	Reduced Orexin-A Levels in Frontotemporal Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2013, 28, 606-611.	1.9	27
38	The attitude of elderly lay people towards the symptoms of dementia. International Psychogeriatrics, 2006, 18, 251-258.	1.0	26
39	Cognitive Impairment in Neuro-Behcet's Disease and Multiple Sclerosis: A Comparative Study. International Journal of Neuroscience, 2012, 122, 650-656.	1.6	26
40	Caregiver Burden, Quality of Life and Related Factors in Family Caregivers of Dementia Patients in Turkey. Issues in Mental Health Nursing, 2020, 41, 741-749.	1.2	26
41	Evaluation of Incidence and Clinical Features of Antibody-Associated Autoimmune Encephalitis Mimicking Dementia. Behavioural Neurology, 2014, 2014, 1-4.	2.1	25
42	Brainâ€derived neurotrophic factor gene Val66Met polymorphism and cognitive function in obsessiveâ€“compulsive disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 850-858.	1.7	24
43	Intracranial hypertension in BehÃ§et's Disease*. European Journal of Neurology, 1996, 3, 66-70.	3.3	23
44	Expression changes of genes associated with apoptosis and survival processes in Parkinsonâ€™s disease. Neuroscience Letters, 2016, 615, 72-77.	2.1	22
45	Therapeutic Effects of an Acetylcholinesterase Inhibitor (Donepezil) on Memory in Wernickeâ€“Korsakoff's Disease. Clinical Neuropharmacology, 2002, 25, 16-20.	0.7	21
46	Clozapine Treatment in Oromandibular Dystonia. Clinical Neuropharmacology, 2004, 27, 84-86.	0.7	21
47	Episodic memory and metamemory in Parkinsonâ€™s disease patients.. Neuropsychology, 2009, 23, 736-745.	1.3	21
48	Genetic bases and phenotypes of autosomal recessive Parkinson disease in a Turkish population. European Journal of Neurology, 2012, 19, 769-775.	3.3	20
49	Secondary paroxysmal kinesigenic dyskinesia associated with CLCN2 gene mutation. Parkinsonism and Related Disorders, 2015, 21, 544-546.	2.2	20
50	Mutation Frequency of the Major Frontotemporal Dementia Genes, MAPT, GRN and C9ORF72 in a Turkish Cohort of Dementia Patients. PLoS ONE, 2016, 11, e0162592.	2.5	19
51	The Association Between Clusterin and APOE Polymorphisms and Late-Onset Alzheimer Disease in a Turkish Cohort. Journal of Geriatric Psychiatry and Neurology, 2016, 29, 221-226.	2.3	19
52	An investigation of affective theory of mind ability and its relation to neuropsychological functions in Alzheimer's disease. Journal of Neuropsychology, 2020, 14, 399-415.	1.4	18
53	Magnetic nerve root stimulation in two types of brachial plexus injury: Segmental demyelination and axonal degeneration. , 1997, 20, 823-832.		17
54	Non-Convulsive Status Epilepticus Associated With Glutamic Acid Decarboxylase Antibody. Clinical EEG and Neuroscience, 2013, 44, 232-236.	1.7	17

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55	Adult-onset phenylketonuria with rapidly progressive dementia and parkinsonism. Neurocase, 2016, 22, 273-275.	0.6	17
56	Association between inflammatory markers and cognitive outcome in patients with acute brain dysfunction due to sepsis. Noropsikiyatri Arsivi, 2018, 56, 63-70.	0.3	17
57	The cerebral blood flow deficits in Parkinsonâ€™s disease with mild cognitive impairment using arterial spin labeling MRI. Journal of Neural Transmission, 2020, 127, 1285-1294.	2.8	16
58	Polymorphisms in the DNA repair genes XPD (ERCC2) and XPF (ERCC4) are not associated with sporadic late-onset Alzheimer's disease. Neuroscience Letters, 2006, 404, 258-261.	2.1	15
59	Crossed Aphasia in a Dextral Patient With Logopenic/Phonological Variant of Primary Progressive Aphasia. Alzheimer Disease and Associated Disorders, 2012, 26, 282-284.	1.3	15
60	Affective theory of mind in human aging: is there any relation with executive functioning?. Aging, Neuropsychology, and Cognition, 2020, 27, 207-219.	1.3	15
61	Obsessiveâ€“Compulsive Disorder Secondary to Bilateral Frontal Damage Due to a Closed Head Injury. Cognitive and Behavioral Neurology, 2004, 17, 118-120.	0.9	14
62	Interleukin-1Î± â€“889 C/T Polymorphism in Turkish Patients with Late-Onset Alzheimerâ€™s Disease. Dementia and Geriatric Cognitive Disorders, 2009, 27, 82-87.	1.5	14
63	Dysexecutive Syndrome. Cognitive and Behavioral Neurology, 2012, 25, 57-62.	0.9	14
64	COMT Val158Met Polymorphism and Executive Functions in Obsessive-Compulsive Disorder. Journal of Neuropsychiatry and Clinical Neurosciences, 2013, 25, 214-221.	1.8	13
65	Cognitive and anatomical correlates of anosognosia in amnestic mild cognitive impairment and early-stage Alzheimer's disease. International Psychogeriatrics, 2017, 29, 293-302.	1.0	13
66	Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span. Brain and Development, 2018, 40, 458-464.	1.1	13
67	A novel PNPLA6 mutation in a Turkish family with intractable Holmes tremor and spastic ataxia. Neurological Sciences, 2021, 42, 1535-1539.	1.9	13
68	Modelling the Stroop effect: A connectionist approach. Neurocomputing, 2007, 70, 1414-1423.	5.9	12
69	Differentiating symptomatic Parkin mutations carriers from patients with idiopathic Parkinsonâ€™s disease: Contribution of automated segmentation neuroimaging method. Parkinsonism and Related Disorders, 2012, 18, 562-566.	2.2	12
70	Evaluation of cognitive performance in professional divers by means of event-related potentials and neuropsychology. Clinical Neurophysiology, 2017, 128, 579-588.	1.5	12
71	Resting-state fMRI analysis in apathetic Alzheimer's disease. Diagnostic and Interventional Radiology, 2020, 26, 363-369.	1.5	12
72	Elevated sTREM2 and NFL levels in patients with sepsis associated encephalopathy. International Journal of Neuroscience, 2023, 133, 327-333.	1.6	12

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73	P3 response during short-term memory retrieval revisited by a spatio-temporal analysis. International Journal of Psychophysiology, 2012, 84, 205-210.	1.0	11
74	Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis. Neurodegenerative Diseases, 2019, 19, 218-224.	1.4	11
75	Peripheral GRN mRNA and Serum Progranulin Levels as a Potential Indicator for Both the Presence of Splice Site Mutations and Individuals at Risk for Frontotemporal Dementia. Journal of Alzheimer's Disease, 2019, 67, 159-167.	2.6	11
76	PLA2G6 Mutations Related to Distinct Phenotypes: A New Case with Early-onset Parkinsonism. Tremor and Other Hyperkinetic Movements, 2016, 6, 363.	2.0	11
77	Comparison of epidural analgesia combined to general anesthesia and general anesthesia for postoperative cognitive dysfunction in elderly patients. Ulusal Travma Ve Acil Cerrahi Dergisi, 2020, 26, 30-36.	0.3	11
78	The Right Temporal Variant of Frontotemporal Dementia Is Not Genetically Sporadic: A Case Series. Journal of Alzheimer's Disease, 2021, 79, 1195-1201.	2.6	10
79	Default mode and dorsal attention network involvement in visually guided motor sequence learning. Cortex, 2022, 146, 89-105.	2.4	10
80	Neuromuscular consequences of prolonged hunger strike: an electrophysiological study. Clinical Neurophysiology, 2000, 111, 2064-2070.	1.5	9
81	Discrimination ability of the Short Test of Mental Status (STMS) compared to the Mini Mental State Examination (MMSE) in the spectrum of normal cognition, mild cognitive impairment, and probable Alzheimerâ€™s disease dementia: The Turkish standardization study. Journal of Clinical and Experimental Neuropsychology, 2020, 42, 450-458.	1.3	9
82	Neurological features and outcomes of Wilsonâ€™s disease: a single-center experience. Neurological Sciences, 2021, 42, 3829-3834.	1.9	9
83	Cortical excitability in Duchenne muscular dystrophy. Clinical Neurophysiology, 2008, 119, 459-465.	1.5	8
84	Presenilin-1 Gene Intronic Polymorphism and Late-onset Alzheimer's Disease. Journal of Geriatric Psychiatry and Neurology, 2008, 21, 268-273.	2.3	8
85	NMDA Receptor Encephalitis with Cancer of Unknown Primary Origin. Tumori, 2016, 102, S3-S4.	1.1	8
86	SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey. Neurological Sciences, 2017, 38, 2203-2207.	1.9	8
87	Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey. Parkinsonism and Related Disorders, 2018, 48, 34-39.	2.2	8
88	G82S polymorphism of receptor for advanced glycation end products gene and serum soluble RAGE levels in mild cognitive impairment and dementia of Alzheimer's type patients in Turkish population. Journal of Clinical Neuroscience, 2019, 59, 197-201.	1.5	8
89	Development of somatic markers guiding decision-making along adolescence. International Journal of Psychophysiology, 2019, 137, 82-91.	1.0	8
90	Peripheral TREM2 mRNA levels in early and late-onset Alzheimer diseaseâ€™s patients. Molecular Biology Reports, 2020, 47, 5903-5909.	2.3	8

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91	Detection of visual and frontoparietal network perfusion deficits in Parkinsonâ€™s disease dementia. European Journal of Radiology, 2021, 144, 109985.	2.6	7
92	LRRK2 mutations are uncommon in Turkey. European Journal of Neurology, 2011, 18, e137.	3.3	6
93	Akinetic Mutism Without a Structural Prefrontal Lesion. Cognitive and Behavioral Neurology, 2013, 26, 59-62.	0.9	6
94	Serotonin transporter promoter polymorphism is associated with executive function impairments in patients with obsessive compulsive disorder. Clinical Neuropsychologist, 2016, 30, 536-546.	2.3	6
95	Mutations in TYROBP are not a common cause of dementia in a Turkish cohort. Neurobiology of Aging, 2017, 58, 240.e1-240.e3.	3.1	6
96	Effects of cerebral oxygen changes during coronary bypass surgery on postoperative cognitive dysfunction in elderly patients: a pilot study. Brazilian Journal of Anesthesiology (Elsevier), 2018, 68, 142-148.	0.4	6
97	Association between selected cholesterol-related gene polymorphisms and Alzheimerâ€™s disease in a Turkish cohort. Molecular Biology Reports, 2019, 46, 1701-1707.	2.3	6
98	Genetic variants of vitamin D metabolism-related DHCR7/NADSYN1 locus and CYP2R1 gene are associated with clinical features of Parkinsonâ€™s disease. International Journal of Neuroscience, 2022, 132, 439-449.	1.6	5
99	Functional neural substrates of football fanaticism: Different pattern of brain responses and connectivity in fanatics. Psychiatry and Clinical Neurosciences, 2020, 74, 480-487.	1.8	5
100	Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers. Journal of Parkinson's Disease, 2021, 11, 559-568.	2.8	5
101	TREM2 variants as a possible cause of frontotemporal dementia with distinct neuroimaging features. European Journal of Neurology, 2021, 28, 2603-2613.	3.3	5
102	Neuro-BehÃ§etâ€™s Disease with Chorea. Noropsikiyatri Arsivi, 2015, 52, 200-201.	0.7	5
103	A Rare Dementing Disease: Adult Neuronal Ceroid Lipofuscinoses. Journal of Neuropsychiatry and Clinical Neurosciences, 2012, 24, 493-498.	1.8	4
104	Prevalence of HIVâ€associated neurocognitive disorder (HAND) in Turkey and assessment of Addenbrookeâ€™s Cognitive Examination Revised (ACEâ€R) test as a screening tool. HIV Medicine, 2021, 22, 60-66.	2.2	4
105	Neuroinflammation mediators are reduced in sera of Parkinson?s disease patients with mild cognitive impairment. Noropsikiyatri Arsivi, 2019, 57, 15-17.	0.3	4
106	P 043 Neuropsychological follow-up of 12 patients with neuro-BehÃ§et's disease. Revue De Medecine Interne, 1993, 14, 74s.	1.0	3
107	Erdheim Chester disease presenting as slowly progressive cerebellar syndrome and asymptomatic widespread skeletal involvement. European Journal of Neurology, 2011, 18, e93-e93.	3.3	3
108	The effect of two different glycemic management protocols on postoperative cognitive dysfunction in coronary artery bypass surgery. Brazilian Journal of Anesthesiology (Elsevier), 2017, 67, 258-265.	0.4	3

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109	Optimal population screening policies for Alzheimerâ€™s disease*. IISE Transactions on Healthcare Systems Engineering, 2019, 9, 14-25.	1.7	3
110	Normative data for the Turkish version of the pyramids and palm trees test. Applied Neuropsychology Adult, 2021, , 1-7.	1.2	3
111	P3-097: Serum interleukin 1 alpha and alpha 2 macroglobulin levels in patients with early- or late-onset Alzheimer's disease or mild cognitive impairment. , 2015, 11, P657-P657.		2
112	P2â€244: THE CORRELATION BETWEEN CSF AMYLOID BETA 1â€42 LEVELS AND CSF VITAMIN D (25OHD) LEVELS IN PATIENTS WITH SPORADIC ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2018, 14, P766.	0.8	2
113	Amyloid Beta Adsorption Problem with Transfer Plates in Amyloid Beta 1â€“42 IVD Kits. Journal of Molecular Neuroscience, 2019, 67, 534-539.	2.3	2
114	Laboratory and clinical correlates of brain atrophy in Neuro-BehÃ§et's disease. Journal of the Neurological Sciences, 2020, 414, 116831.	0.6	2
115	A novel SACS p.Pro4154GlnfsTer20 mutation in a family with autosomal recessive spastic ataxia of Charlevoix-Saguenay. Neurological Sciences, 2021, 42, 2969-2973.	1.9	2
116	Intensive voice treatment (the Lee Silverman Voice Treatment [LSVT^{Â®}LOUD]) for individuals with Wilsonâ€™s disease and adult cerebral palsy: two case reports. Logopedics Phoniatrics Vocology, 2022, 47, 262-270.	1.0	2
117	Implicit contextual learning in spinocerebellar ataxia.. Neuropsychology, 2020, 34, 511-523.	1.3	2
118	Erken Evre Alzheimer HastalÄ±ÄŸÄ±nda Ä°zlenen Ak Madde Hiperintensitelerinin Depresif Semptomlar ve GÃ¼nlÃ¼k YaÅŸam Aktiviteleri ile Ä°liÅŸkisi. Noropsikiyatri Arsivi, 2013, 50, 360-363.	0.7	2
119	P 036 Clinical experience with neuro-BehÃ§et's disease. Revue De Medecine Interne, 1993, 14, 71s.	1.0	1
120	Analysis of copy number variation in a Turkish dementia cohort. Alzheimer's and Dementia, 2020, 16, e044868.	0.8	1
121	B-Tensor: Brain Connectome Tensor Factorization for Alzheimer's Disease. IEEE Journal of Biomedical and Health Informatics, 2021, 25, 1591-1600.	6.3	1
122	A novel PSEN2 p.Ser175Phe variant in a family with Alzheimerâ€™s disease. Neurological Sciences, 2021, 42, 2497-2504.	1.9	1
123	Frequency of frontotemporal dementia-related gene variants in Turkey. Neurobiology of Aging, 2021, 106, 332.e1-332.e11.	3.1	1
124	Social Cognition or Towards a Mental Psychology/Neurology as a Dialectical Synthesis of Faculty Psychology. Noropsikiyatri Arsivi, 2014, 51, 298-302.	0.7	1
125	Factors Predicting Falls in Parkinsonâ€™s Disease: Investigation of Motor, Non-motor Findings and Different Dual Task Activities. Turk Noroloji Dergisi = Turkish Journal of Neurology, 2020, 26, 126-132.	0.3	1
126	Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey. Parkinsonism and Related Disorders, 2021, 93, 35-39.	2.2	1

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127	Volumetric changes within hippocampal subfields in Alzheimerâ€™s disease continuum. Neurological Sciences, 2022, , 1.	1.9	1
128	Verbal and Nonverbal Memory in Neurodegenerative and Stroke Aphasia: Evidence From the Turkish Version of the Three Words Three Shapes Test. Cognitive and Behavioral Neurology, 2022, 35, 49-65.	0.9	1
129	Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment. Acta Neurologica Belgica, 2023, 123, 823-829.	1.1	1
130	Genotypeâ€“Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review. BMC Neurology, 2022, 22, 122.	1.8	1
131	Parkinson HastalÄ±ÄŸÄ± GeliÅŸtirenEsansiyel Tremor ile Esansiyel Tremor ArasÄ±ndaki FarklÄ±lÄ±klar. Noropsikiyatri Arsivi, 2011, 48, 1-1.	0.7	0
132	The Impact of Familial Structure on Parkinson's Disease in Istanbul Medical School, Turkey. International Journal of Neuroscience, 2012, 122, 102-105.	1.6	0
133	Sporadik Creutzfeldt-Jacob HastalÄ±ÄŸÄ±nda SupranÃ¼kleer BakÄ±ÅŸ Parezisi. Noropsikiyatri Arsivi, 2014, 51, 91-92.	0.7	0
134	P1-343: MOCA, MMSE, AND ACE-R FOR THE ASSESSMENT OF MILD COGNITIVE IMPAIRMENT IN PATIENTS WITH PARKINSON's DISEASE. , 2014, 10, P438-P438.		0
135	P1-287: NEUROIMAGING FINDINGS OF NASU-HAKOLA DISEASE. , 2014, 10, P415-P415.		0
136	P4-266: MAPT MUTATION CAUSING RIGHT TEMPORAL VARIANT SEMANTIC DEMENTIA. , 2014, 10, P882-P882.		0
137	P1-165: EVALUATION OF INCIDENCE AND CLINICAL FEATURES OF ANTIBODY-ASSOCIATED AUTOIMMUNE ENCEPHALITIS MIMICKING DEMENTIA. , 2014, 10, P360-P360.		0
138	P4-114: The applicability of new screening instruments for cognitive impairment in parkinson's disease in turkey. , 2015, 11, P819-P820.		0
139	PSEN1 mutation presenting as posterior cortical atrophy. Journal of the Neurological Sciences, 2015, 357, e127.	0.6	0
140	P1-052: Association between clusterin polymorphisms and Alzheimer's disease. , 2015, 11, P358-P358.		0
141	P3-082: Compromised regulation of serum cytokine levels and BDNF due to low levels of vitamin d in patients with early- or late-onset Alzheimer's disease or parkinson's disease. , 2015, 11, P649-P650.		0
142	P1â€233: Sporadic Fatal Insomnia In A Young Man. Alzheimer's and Dementia, 2016, 12, P496.	0.8	0
143	[P2â€“405]: FUNCTIONAL NEUROIMAGING APPROACH TO APATHETIC ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2017, 13, P786.	0.8	0
144	[P2â€“447]: A COMPARISON OF AFFECTIVE THEORY OF MIND ABILITIES BETWEEN PARTICIPANTS WITH AMNESTIC MILD COGNITIVE IMPAIRMENT AND HEALTHY CONTROLS. Alzheimer's and Dementia, 2017, 13, P808.	0.8	0

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145	[P1â€“290]: THE RELATIONSHIP BETWEEN CSF AMYLOID BETA CONCENTRATIONS AND FREE AND CUED RECALL PERFORMANCE AMONG PARTICIPANTS WITH AMNESTIC MILD COGNITIVE IMPAIRMENT. Alzheimer's and Dementia, 2017, 13, P362.	0.8	0
146	[P1â€“343]: INTERCORRELATIONS BETWEEN CSF AMYLOID BETA LEVELS AND NEUROPSYCHOLOGICAL PROFILES AMONG PARTICIPANTS WITH SUBJECTIVE COGNITIVE IMPAIRMENT AND MCI OF THE AMNESTIC TYPE. Alzheimer's and Dementia, 2017, 13, P387.	0.8	0
147	[P2â€“328]: DIFFERENTIAL RESTING STATE CONNECTIVITY AND ITS RELATIONSHIP WITH COGNITIVE PERFORMANCE AMONG PARTICIPANTS WITH SUBJECTIVE COGNITIVE IMPAIRMENT (SCI), MILD COGNITIVE IMPAIRMENT (MCI), AND EARLY STAGE ALZHEIMER's DISEASE (AD). Alzheimer's and Dementia, 2017, 13, P744.	0.8	0
148	Determination of Diffusion Weighted Magnetic Resonance Imaging Based Biomarkers of Mild Cognitive Impairment in Parkinsonâ€™s Disease. , 2017, , .		0
149	P1â€317: TASKâ€RELATED AND TASKâ€FREE FUNCTIONAL NEUROIMAGING IN APATHETIC ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2018, 14, P413.	0.8	0
150	P3â€109: ASSOCIATION BETWEEN NADSYN1/DHCR7 AND CYP2R1 GENOTYPES AND PARKINSON'S DISEASE AND ITS CLINICAL FEATURES. Alzheimer's and Dementia, 2018, 14, P1109.	0.8	0
151	P4â€085: CATATONIA AS CLINICAL PRESENTATIONÂOF ANTIâ€Nâ€METHYLâ€Dâ€ASPARTATE (ANTIâ€NMDA) RECEPTOR ENCEPHALITIS. Alzheimer's and Dementia, 2018, 14, P1467.	0.8	0
152	P2â€229: THE EFFECT OF CSF AMYLOID BETA CONCENTRATIONS ON MEMORY PERFORMANCE OF THE INDIVIDUALS WITH SUBJECTIVE COGNITIVE IMPAIRMENT. Alzheimer's and Dementia, 2018, 14, P756.	0.8	0
153	P2â€302: CLINICAL FEATURES AND DIAGNOSIS OF EARLY ONSET DEGENERATIVE DEMENTIAS IN TURKEY. Alzheimer's and Dementia, 2018, 14, P798.	0.8	0
154	ADDIA, A MULTICENTER CLINICAL STUDY OF PROOF-OF-PERFORMANCE TO VALIDATE BLOOD-BASED BIOMARKERS FOR THE DIAGNOSIS OF ALZHEIMERâ€™S DISEASE (AD): THE STUDY PROTOCOL. , 2018, 14, P781.		0
155	P4â€087: SPORADIC FATAL INSOMNIA IN A YOUNG TURKISH MAN. Alzheimer's and Dementia, 2018, 14, P1468.	0.8	0
156	Association between PSEN1 p.E318G Variant and APOE Polymorphism and Alzheimer Disease in Turkish Patients. Turk Noroloji Dergisi = Turkish Journal of Neurology, 2021, 27, 117-122.	0.3	0
157	A comprehensive analysis of copy number variation in a Turkish dementia cohort. Human Genomics, 2021, 15, 48.	2.9	0
158	NÃ–RO-BEHÃ‡ET HASTALIÄžINDA KORE: OLGU SUNUMU. Noropsikiyatri Arsivi, 0, , .	0.7	0
159	Comparison of epidural analgesia combined to general anesthesia and general anesthesia for postoperative cognitive dysfunction in elderly patients. Noropsikiyatri Arsivi, 2019, , .	0.3	0
160	Factors Predicting Falls in Parkinsonâ€™s Disease: Investigation of Motor, Non-motor Findings and Different Dual Task Activities. Turk Noroloji Dergisi = Turkish Journal of Neurology, 0, , .	0.3	0
161	lncRNAs as a novel source of diagnostic applications for early Alzheimerâ€™s disease and other dementia types. Alzheimer's and Dementia, 2020, 16, e039788.	0.8	0
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