Hakan I GÃ¼rvit

List of Publications by Year
in descending order

Version: 2024-02-01

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	Elevated sTREM2 and NFL levels in patients with sepsis associated encephalopathy. International Journal of Neuroscience, 2023, 133, 327-333.	1.6	12
2	Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment. Acta Neurologica Belgica, 2023, 123, 823-829.	1.1	1
3	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
4	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
5	Default mode and dorsal attention network involvement in visually guided motor sequence learning. Cortex, 2022, 146, 89-105.	2.4	10
6	Volumetric changes within hippocampal subfields in Alzheimerâ€™s disease continuum. Neurological Sciences, 2022, , 1.	1.9	1
7	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
8	Genotypeâ€“Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review. BMC Neurology, 2022, 22, 122.	1.8	1
9	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
10	A novel PNPLA6 mutation in a Turkish family with intractable Holmes tremor and spastic ataxia. Neurological Sciences, 2021, 42, 1535-1539.	1.9	13
11	B-Tensor: Brain Connectome Tensor Factorization for Alzheimer's Disease. IEEE Journal of Biomedical and Health Informatics, 2021, 25, 1591-1600.	6.3	1
12	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
13	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
14	Normative data for the Turkish version of the pyramids and palm trees test. Applied Neuropsychology Adult, 2021, , 1-7.	1.2	3
15	A novel PSEN2 p.Ser175Phe variant in a family with Alzheimerâ€™s disease. Neurological Sciences, 2021, 42, 2497-2504.	1.9	1
16	Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers. Journal of Parkinson's Disease, 2021, 11, 559-568.	2.8	5
17	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
18	TREM2 variants as a possible cause of frontotemporal dementia with distinct neuroimaging features. European Journal of Neurology, 2021, 28, 2603-2613.	3.3	5

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19	A comprehensive analysis of copy number variation in a Turkish dementia cohort. Human Genomics, 2021, 15, 48.	2.9	0
20	Detection of visual and frontoparietal network perfusion deficits in Parkinsonâ€™s disease dementia. European Journal of Radiology, 2021, 144, 109985.	2.6	7
21	Frequency of frontotemporal dementia-related gene variants in Turkey. Neurobiology of Aging, 2021, 106, 332.e1-332.e11.	3.1	1
22	Neurological features and outcomes of Wilsonâ€™s disease: a single-center experience. Neurological Sciences, 2021, 42, 3829-3834.	1.9	9
23	Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey. Parkinsonism and Related Disorders, 2021, 93, 35-39.	2.2	1
24	Homozygosity analysis in a Turkish dementia cohort.. Alzheimer's and Dementia, 2021, 17 Suppl 3, e054052.	0.8	0
25	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
26	Peripheral TREM2 mRNA levels in early and late-onset Alzheimer diseaseâ€™s patients. Molecular Biology Reports, 2020, 47, 5903-5909.	2.3	8
27	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
28	Analysis of copy number variation in a Turkish dementia cohort. Alzheimer's and Dementia, 2020, 16, e044868.	0.8	1
29	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
30	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
31	Resting-state fMRI analysis in apathetic Alzheimer's disease. Diagnostic and Interventional Radiology, 2020, 26, 363-369.	1.5	12
32	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
33	Laboratory and clinical correlates of brain atrophy in Neuro-BehÃ§et's disease. Journal of the Neurological Sciences, 2020, 414, 116831.	0.6	2
34	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
35	Implicit contextual learning in spinocerebellar ataxia.. Neuropsychology, 2020, 34, 511-523.	1.3	2
36	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

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37	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
38	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
39	The association of serum clusterin levels and Clusterin rs11136000 polymorphisms with Alzheimer disease in a Turkish cohort. Neurological Sciences and Neurophysiology, 2020, 37, 134.	0.3	0
40	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
41	Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis. Neurodegenerative Diseases, 2019, 19, 218-224.	1.4	11
42	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
43	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
44	Optimal population screening policies for Alzheimerâ€™s disease*. IISE Transactions on Healthcare Systems Engineering, 2019, 9, 14-25.	1.7	3
45	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
46	Development of somatic markers guiding decision-making along adolescence. International Journal of Psychophysiology, 2019, 137, 82-91.	1.0	8
47	Comparison of epidural analgesia combined to general anesthesia and general anesthesia for postoperative cognitive dysfunction in elderly patients. Noropsikiyatri Arsivi, 2019, , .	0.3	0
48	Neuroinflammation mediators are reduced in sera of Parkinson?s disease patients with mild cognitive impairment. Noropsikiyatri Arsivi, 2019, 57, 15-17.	0.3	4
49	Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey. Parkinsonism and Related Disorders, 2018, 48, 34-39.	2.2	8
50	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
51	Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span. Brain and Development, 2018, 40, 458-464.	1.1	13
52	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
53	P1â€317: TASKâ€RELATED AND TASKâ€FREE FUNCTIONAL NEUROIMAGING IN APATHETIC ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2018, 14, P413.	0.8	0
54	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

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55	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
56	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
57	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
58	P2â€302: CLINICAL FEATURES AND DIAGNOSIS OF EARLY ONSET DEGENERATIVE DEMENTIAS IN TURKEY. Alzheimer's and Dementia, 2018, 14, P798.	0.8	0
59	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
60	P4â€087: SPORADIC FATAL INSOMNIA IN A YOUNG TURKISH MAN. Alzheimer's and Dementia, 2018, 14, P1468.	0.8	0
61	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
62	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
63	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
64	Evaluation of cognitive performance in professional divers by means of event-related potentials and neuropsychology. Clinical Neurophysiology, 2017, 128, 579-588.	1.5	12
65	Salience network engagement with the detection of morally laden information. Social Cognitive and Affective Neuroscience, 2017, 12, 1118-1127.	3.0	30
66	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
67	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
68	SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey. Neurological Sciences, 2017, 38, 2203-2207.	1.9	8
69	[P2â€“405]: FUNCTIONAL NEUROIMAGING APPROACH TO APATHETIC ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2017, 13, P786.	0.8	0
70	[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
71	[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
72	[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

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73	[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
74	Determination of Diffusion Weighted Magnetic Resonance Imaging Based Biomarkers of Mild Cognitive Impairment in Parkinsonâ€™s Disease. , 2017, , .		0
75	NMDA Receptor Encephalitis with Cancer of Unknown Primary Origin. Tumori, 2016, 102, S3-S4.	1.1	8
76	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
77	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
78	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
79	P1â€233: Sporadic Fatal Insomnia In A Young Man. Alzheimer's and Dementia, 2016, 12, P496.	0.8	0
80	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
81	Expression changes of genes associated with apoptosis and survival processes in Parkinsonâ€™s disease. Neuroscience Letters, 2016, 615, 72-77.	2.1	22
82	Adult-onset phenylketonuria with rapidly progressive dementia and parkinsonism. Neurocase, 2016, 22, 273-275.	0.6	17
83	PLA2G6 Mutations Related to Distinct Phenotypes: A New Case with Early-onset Parkinsonism. Tremor and Other Hyperkinetic Movements, 2016, 6, 363.	2.0	11
84	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
85	P4-114: The applicability of new screening instruments for cognitive impairment in parkinson's disease in turkey. , 2015, 11, P819-P820.		0
86	PSEN1 mutation presenting as posterior cortical atrophy. Journal of the Neurological Sciences, 2015, 357, e127.	0.6	0
87	The Central Biobank and Virtual Biobank of BIOMARKAPD: A Resource for Studies on Neurodegenerative Diseases. Frontiers in Neurology, 2015, 6, 216.	2.4	36
88	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
89	P1-052: Association between clusterin polymorphisms and Alzheimer's disease. , 2015, 11, P358-P358.		0
90	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

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91	The distinct genetic pattern of ALS in Turkey and novel mutations. Neurobiology of Aging, 2015, 36, 1764.e9-1764.e18.	3.1	78
92	Secondary paroxysmal kinesigenic dyskinesia associated with CLCN2 gene mutation. Parkinsonism and Related Disorders, 2015, 21, 544-546.	2.2	20
93	A new Fâ€box protein 7 gene mutation causing typical Parkinson's disease. Movement Disorders, 2015, 30, 1130-1133.	3.9	59
94	Neuro-BehÃ§etâ€™s Disease with Chorea. Noropsikiyatri Arsivi, 2015, 52, 200-201.	0.7	5
95	Evaluation of Incidence and Clinical Features of Antibody-Associated Autoimmune Encephalitis Mimicking Dementia. Behavioural Neurology, 2014, 2014, 1-4.	2.1	25
96	TREM2 mutations implicated in neurodegeneration impair cell surface transport and phagocytosis. Science Translational Medicine, 2014, 6, 243ra86.	12.4	600
97	Sporadik Creutzfeldt-Jacob HastalÄ±ÄŸÄ±nda SupranÃ¼kleer BakÄ±ÅŸ Parezisi. Noropsikiyatri Arsivi, 2014, 51, 91-92.	0.7	0
98	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
99	P1-287: NEUROIMAGING FINDINGS OF NASU-HAKOLA DISEASE. , 2014, 10, P415-P415.		0
100	P4-266: MAPT MUTATION CAUSING RIGHT TEMPORAL VARIANT SEMANTIC DEMENTIA. , 2014, 10, P882-P882.		0
101	P1-165: EVALUATION OF INCIDENCE AND CLINICAL FEATURES OF ANTIBODY-ASSOCIATED AUTOIMMUNE ENCEPHALITIS MIMICKING DEMENTIA. , 2014, 10, P360-P360.		0
102	Social Cognition or Towards a Mental Psychology/Neurology as a Dialectical Synthesis of Faculty Psychology. Noropsikiyatri Arsivi, 2014, 51, 298-302.	0.7	1
103	Non-Convulsive Status Epilepticus Associated With Glutamic Acid Decarboxylase Antibody. Clinical EEG and Neuroscience, 2013, 44, 232-236.	1.7	17
104	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
105	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
106	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
107	Reduced Orexin-A Levels in Frontotemporal Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2013, 28, 606-611.	1.9	27
108	Akinetic Mutism Without a Structural Prefrontal Lesion. Cognitive and Behavioral Neurology, 2013, 26, 59-62.	0.9	6

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109	COMT Val158Met Polymorphism and Executive Functions in Obsessive-Compulsive Disorder. Journal of Neuropsychiatry and Clinical Neurosciences, 2013, 25, 214-221.	1.8	13
110	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
111	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
112	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
113	A Rare Dementing Disease: Adult Neuronal Ceroid Lipofuscinoses. Journal of Neuropsychiatry and Clinical Neurosciences, 2012, 24, 493-498.	1.8	4
114	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
115	<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
116	Cognitive Impairment in Neuro-Behcet's Disease and Multiple Sclerosis: A Comparative Study. International Journal of Neuroscience, 2012, 122, 650-656.	1.6	26
117	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
118	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
119	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
120	P3 response during short-term memory retrieval revisited by a spatio-temporal analysis. International Journal of Psychophysiology, 2012, 84, 205-210.	1.0	11
121	Neuropsychological function in obsessive-compulsive disorder. Comprehensive Psychiatry, 2012, 53, 167-175.	3.1	59
122	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
123	Dysexecutive Syndrome. Cognitive and Behavioral Neurology, 2012, 25, 57-62.	0.9	14
124	Genetic bases and phenotypes of autosomal recessive Parkinson disease in a Turkish population. European Journal of Neurology, 2012, 19, 769-775.	3.3	20
125	Pain is common in Parkinson's disease. Clinical Neurology and Neurosurgery, 2011, 113, 11-13.	1.4	78
126	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

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127	LRRK2 mutations are uncommon in Turkey. European Journal of Neurology, 2011, 18, e137.	3.3	6
128	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
129	Parkinson HastalÄ±ÄŸÄ± GeliÅŸtirenEsansiyel Tremor ile Esansiyel Tremor ArasÄ±ndaki FarklÄ±lÄ±klar. Noropsikiyatri Arsivi, 2011, 48, 1-1.	0.7	0
130	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
131	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
132	Episodic memory and metamemory in Parkinsonâ€™s disease patients.. Neuropsychology, 2009, 23, 736-745.	1.3	21
133	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
134	Cortical excitability in Duchenne muscular dystrophy. Clinical Neurophysiology, 2008, 119, 459-465.	1.5	8
135	Presenilin-1 Gene Intronic Polymorphism and Late-onset Alzheimer's Disease. Journal of Geriatric Psychiatry and Neurology, 2008, 21, 268-273.	2.3	8
136	Association between Vitamin D Receptor Gene Polymorphism and Alzheimer's Disease. Tohoku Journal of Experimental Medicine, 2007, 212, 275-282.	1.2	134
137	Modelling the Stroop effect: A connectionist approach. Neurocomputing, 2007, 70, 1414-1423.	5.9	12
138	Clinical and magnetic resonance imaging findings of HIV-Negative patients with neurosyphilis. Journal of Neurology, 2007, 254, 368-374.	3.6	34
139	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
140	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
141	The attitude of elderly lay people towards the symptoms of dementia. International Psychogeriatrics, 2006, 18, 251-258.	1.0	26
142	A composite neural network model for perseveration and distractibility in the Wisconsin card sorting test. Neural Networks, 2006, 19, 375-387.	5.9	41
143	Inflammatory/demyelinating central nervous system involvement in familial Mediterranean fever (FMF): coincidence or association?. Journal of Neurology, 2006, 253, 928-934.	3.6	53
144	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

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145	Unrecognized depression in community-dwelling elderly persons in Istanbul. International Psychogeriatrics, 2005, 17, 303-312.	1.0	29
146	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
147	Clozapine Treatment in Oromandibular Dystonia. Clinical Neuropharmacology, 2004, 27, 84-86.	0.7	21
148	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
149	Therapeutic Effects of an Acetylcholinesterase Inhibitor (Donepezil) on Memory in Wernickeâ€“Korsakoff's Disease. Clinical Neuropharmacology, 2002, 25, 16-20.	0.7	21
150	Cognitive impairment in amyotrophic lateral sclerosis: evidence from neuropsychological investigation and event-related potentials. Cognitive Brain Research, 2002, 14, 234-244.	3.0	116
151	Neuromuscular consequences of prolonged hunger strike: an electrophysiological study. Clinical Neurophysiology, 2000, 111, 2064-2070.	1.5	9
152	Neuropsychological follow-up of 12 patients with neuro-BehÃ§et disease. Journal of Neurology, 1999, 246, 113-119.	3.6	90
153	Paraneoplastic limbic encephalitis with immature ovarian teratoma–a case report. Journal of Neuro-Oncology, 1998, 37, 63-66.	2.9	28
154	Magnetic nerve root stimulation in two types of brachial plexus injury: Segmental demyelination and axonal degeneration. , 1997, 20, 823-832.		17
155	Intracranial hypertension in BehÃ§et's Disease*. European Journal of Neurology, 1996, 3, 66-70.	3.3	23
156	Seven-Year Follow-up of Neurologic Involvement in Behcet Syndrome. Archives of Neurology, 1996, 53, 691-694.	4.5	103
157	Paroxysmal dysarthria and ataxia in a patient with Behï¿½et's disease. Journal of Neurology, 1995, 242, 344-347.	3.6	54
158	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
159	P 036 Clinical experience with neuro-BehÃ§et's disease. Revue De Medecine Interne, 1993, 14, 71s.	1.0	1
160	P 043 Neuropsychological follow-up of 12 patients with neuro-BehÃ§et's disease. Revue De Medecine Interne, 1993, 14, 74s.	1.0	3
161	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
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