

Basar Bilgic

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

135
papers

4,069
citations

23
h-index

63
g-index

148
ext. papers

4,986
ext. citations

3.7
avg, IF

4.6
L-index

#	Paper	IF	Citations
135	Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment.. <i>Acta Neurologica Belgica</i> , 2022 , 1	1.5	
134	Does transcranial direct current stimulation enhance cognitive performance in Parkinson's disease mild cognitive impairment? An event-related potentials and neuropsychological assessment study.. <i>Neurological Sciences</i> , 2022 , 1	3.5	
133	Genotype-Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review.. <i>BMC Neurology</i> , 2022 , 22, 122	3.1	
132	Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey. <i>Parkinsonism and Related Disorders</i> , 2021 , 93, 35-39	3.6	0
131	PHACTR1 genetic variability is not critical in small vessel ischemic disease patients and Pcoma recruitment in C57BL/6J mice. <i>Scientific Reports</i> , 2021 , 11, 6072	4.9	1
130	A novel PSEN2 p.Ser175Phe variant in a family with Alzheimer's disease. <i>Neurological Sciences</i> , 2021 , 42, 2497-2504	3.5	0
129	Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers. <i>Journal of Parkinson's Disease</i> , 2021 , 11, 559-568	5.3	0
128	Cerebellar ataxia, neuropathy and vestibular areflexia syndrome (canvas): an important cause of late-onset ataxia with unique clinical features. <i>Acta Neurologica Belgica</i> , 2021 , 1	1.5	
127	TREM2 variants as a possible cause of frontotemporal dementia with distinct neuroimaging features. <i>European Journal of Neurology</i> , 2021 , 28, 2603-2613	6	1
126	Intensive voice treatment (the Lee Silverman Voice Treatment [LSVTLOUD]) for individuals with Wilson's disease and adult cerebral palsy: two case reports. <i>Logopedics Phoniatrics Vocology</i> , 2021 , 1-9	1.3	
125	A novel PNPLA6 mutation in a Turkish family with intractable Holmes tremor and spastic ataxia. <i>Neurological Sciences</i> , 2021 , 42, 1535-1539	3.5	4
124	B-Tensor: Brain Connectome Tensor Factorization for Alzheimer's Disease. <i>IEEE Journal of Biomedical and Health Informatics</i> , 2021 , 25, 1591-1600	7.2	
123	Inflammation and regulatory T cell genes are differentially expressed in peripheral blood mononuclear cells of Parkinson's disease patients. <i>Scientific Reports</i> , 2021 , 11, 2316	4.9	4
122	A novel SACS p.Pro4154GlnfsTer20 mutation in a family with autosomal recessive spastic ataxia of Charlevoix-Saguenay. <i>Neurological Sciences</i> , 2021 , 42, 2969-2973	3.5	0
121	The Right Temporal Variant of Frontotemporal Dementia Is Not Genetically Sporadic: A Case Series. <i>Journal of Alzheimer's Disease</i> , 2021 , 79, 1195-1201	4.3	5
120	The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice. <i>Movement Disorders</i> , 2021 , 36, 1676-1688	7	3
119	A comprehensive analysis of copy number variation in a Turkish dementia cohort. <i>Human Genomics</i> , 2021 , 15, 48	6.8	

118	Detection of visual and frontoparietal network perfusion deficits in Parkinson's disease dementia. <i>European Journal of Radiology</i> , 2021 , 144, 109985	4.7	1
117	Frequency of frontotemporal dementia-related gene variants in Turkey. <i>Neurobiology of Aging</i> , 2021 , 106, 332.e1-332.e11	5.6	
116	Neurological features and outcomes of Wilson's disease: a single-center experience. <i>Neurological Sciences</i> , 2021 , 42, 3829-3834	3.5	1
115	Homozygosity analysis in a Turkish dementia cohort.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e054052	5.2	
114	Analysis of copy number variation in a Turkish dementia cohort. <i>Alzheimer's and Dementia</i> , 2020 , 16, e044868	4.8	1
113	Do Alzheimer's Disease Patients Appear Younger than Their Real Age?. <i>Dementia and Geriatric Cognitive Disorders</i> , 2020 , 49, 483-488	2.6	
112	Functional neural substrates of football fanaticism: Different pattern of brain responses and connectivity in fanatics. <i>Psychiatry and Clinical Neurosciences</i> , 2020 , 74, 480-487	6.2	2
111	Characterization of recessive Parkinson's disease in a large multicenter study. <i>Annals of Neurology</i> , 2020 , 88, 843	9.4	11
110	Revisiting the complex architecture of ALS in Turkey: Expanding genotypes, shared phenotypes, molecular networks, and a public variant database. <i>Human Mutation</i> , 2020 , 41, e7-e45	4.7	4
109	An investigation of affective theory of mind ability and its relation to neuropsychological functions in Alzheimer's disease. <i>Journal of Neuropsychology</i> , 2020 , 14, 399-415	2.6	10
108	Small ubiquitin-related modifier (SUMO) 3 and SUMO4 gene polymorphisms in Parkinson's disease. <i>Neurological Research</i> , 2020 , 42, 451-457	2.7	1
107	Multicenter study of levodopa carbidopa intestinal gel in Parkinson's disease: the Turkish experience. <i>Turkish Journal of Medical Sciences</i> , 2020 , 50, 66-85	2.7	
106	White-matter changes in early and late stages of mild cognitive impairment. <i>Journal of Clinical Neuroscience</i> , 2020 , 78, 181-184	2.2	3
105	Laboratory and clinical correlates of brain atrophy in Neuro-Behçet's disease. <i>Journal of the Neurological Sciences</i> , 2020 , 414, 116831	3.2	1
104	The COVID-19 from Neurological Overview. <i>Turk Noroloji Dergisi = Turkish Journal of Neurology</i> , 2020 , 26, 58-108	1.1	2
103	Neuroinflammation Mediators are Reduced in Sera of Parkinson's Disease Patients with Mild Cognitive Impairment. <i>Noropsikiyatri Arsivi</i> , 2020 , 57, 15-17	0.6	2
102	The association of serum clusterin levels and Clusterin rs11136000 polymorphisms with Alzheimer disease in a Turkish cohort. <i>Neurological Sciences and Neurophysiology</i> , 2020 , 37, 134	0.4	
101	Peripheral TREM2 mRNA levels in early and late-onset Alzheimer disease's patients. <i>Molecular Biology Reports</i> , 2020 , 47, 5903-5909	2.8	3

100	The cerebral blood flow deficits in Parkinson's disease with mild cognitive impairment using arterial spin labeling MRI. <i>Journal of Neural Transmission</i> , 2020 , 127, 1285-1294	4.3	4
99	Genetic variants of vitamin D metabolism-related locus and gene are associated with clinical features of Parkinson's disease. <i>International Journal of Neuroscience</i> , 2020 , 1-11	2	0
98	Brain Volume Changes in Patients with Acute Brain Dysfunction Due to Sepsis. <i>Neurocritical Care</i> , 2020 , 32, 459-468	3.3	9
97	Hotspot Mutations in Pleuropulmonary Blastoma: A Case Series From a Tertiary Center. <i>Pediatric and Developmental Pathology</i> , 2020 , 23, 204-209	2.2	1
96	Caregiver Burden, Quality of Life and Related Factors in Family Caregivers of Dementia Patients in Turkey. <i>Issues in Mental Health Nursing</i> , 2020 , 41, 741-749	1.5	9
95	Late-onset phenotype associated with a homozygous GJC2 missense mutation in a Turkish family. <i>Parkinsonism and Related Disorders</i> , 2019 , 66, 228-231	3.6	2
94	Association between selected cholesterol-related gene polymorphisms and Alzheimer's disease in a Turkish cohort. <i>Molecular Biology Reports</i> , 2019 , 46, 1701-1707	2.8	4
93	Parkinson's Disease Dementia and Lewy Body Disease. <i>Seminars in Neurology</i> , 2019 , 39, 274-282	3.2	23
92	Neuroimaging Findings in Sepsis-Induced Brain Dysfunction: Association with Clinical and Laboratory Findings. <i>Neurocritical Care</i> , 2019 , 30, 106-117	3.3	16
91	A patient with early-onset Alzheimer's disease with a novel PSEN1 p.Leu424Pro mutation. <i>Neurobiology of Aging</i> , 2019 , 84, 238.e1-238.e4	5.6	3
90	Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis. <i>Neurodegenerative Diseases</i> , 2019 , 19, 218-224	2.3	4
89	Do Alzheimer's Patients Appear Younger than Their Age? A Study with Automatic Facial Age Estimation 2019 ,		1
88	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. <i>Journal of Alzheimer's Disease</i> , 2019 , 67, 159-167	4.3	7
87	Olfactory function and olfactory bulb volume in Wilson's disease. <i>European Archives of Oto-Rhino-Laryngology</i> , 2019 , 276, 139-142	3.5	4
86	The association between repeat number in C9orf72 and phenotypic variability in Turkish patients with frontotemporal lobar degeneration. <i>Neurobiology of Aging</i> , 2019 , 76, 216.e1-216.e7	5.6	3
85	Amyloid Beta Adsorption Problem with Transfer Plates in Amyloid Beta 1-42 IVD Kits. <i>Journal of Molecular Neuroscience</i> , 2019 , 67, 534-539	3.3	2
84	Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey. <i>Parkinsonism and Related Disorders</i> , 2018 , 48, 34-39	3.6	4
83	Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span. <i>Brain and Development</i> , 2018 , 40, 458-464	2.2	4

82	Progress toward standardized diagnosis of vascular cognitive impairment: Guidelines from the Vascular Impairment of Cognition Classification Consensus Study. <i>Alzheimer's and Dementia</i> , 2018 , 14, 280-292	1.2	136
81	An assessment of Movement Disorder Society Task Force diagnostic criteria for mild cognitive impairment in Parkinson's disease. <i>European Journal of Neurology</i> , 2018 , 25, 148-153	6	21
80	HPCA confirmed as a genetic cause of DYT2-like dystonia phenotype. <i>Movement Disorders</i> , 2018 , 33, 1354-1358	7	19
79	Wernicke's Encephalopathy due to Non-Alcoholic Gastrointestinal Tract Disease. <i>Noropsikiyatri Arsivi</i> , 2018 , 55, 307-314	0.6	2
78	Neuroimaging and Dementia. <i>Noropsikiyatri Arsivi</i> , 2018 , 55, 1-2	0.6	1
77	Evaluation of OnabotulinumtoxinA Treatment in Patients with Concomitant Chronic Migraine and Temporomandibular Disorders. <i>Noropsikiyatri Arsivi</i> , 2018 , 55, 330-336	0.6	2
76	Which cognitive dual-task walking causes most interference on the Timed Up and Go test in Parkinson's disease: a controlled study. <i>Neurological Sciences</i> , 2018 , 39, 2151-2157	3.5	14
75	P3-109: ASSOCIATION BETWEEN NADSYN1/DHCR7 AND CYP2R1 GENOTYPES AND PARKINSON'S DISEASE AND ITS CLINICAL FEATURES 2018 , 14, P1109-P1109		
74	P4-085: CATATONIA AS CLINICAL PRESENTATION OF ANTI-N-METHYL-D-ASPARTATE (ANTI-NMDA) RECEPTOR ENCEPHALITIS 2018 , 14, P1467-P1468		
73	P2-229: THE EFFECT OF CSF AMYLOID BETA CONCENTRATIONS ON MEMORY PERFORMANCE OF THE INDIVIDUALS WITH SUBJECTIVE COGNITIVE IMPAIRMENT 2018 , 14, P756-P756		
72	P2-244: THE CORRELATION BETWEEN CSF AMYLOID BETA 1-42 LEVELS AND CSF VITAMIN D (25OHD) LEVELS IN PATIENTS WITH SPORADIC ALZHEIMER'S DISEASE 2018 , 14, P766-P766		0
71	P2-302: CLINICAL FEATURES AND DIAGNOSIS OF EARLY ONSET DEGENERATIVE DEMENTIAS IN TURKEY 2018 , 14, P798-P798		
70	P4-087: SPORADIC FATAL INSOMNIA IN A YOUNG TURKISH MAN 2018 , 14, P1468-P1469		
69	A Comparison Of The Relations Between Brain Atrophy, Cognition And Optic Coherence Tomography Between Multiple Sclerosis Patients And Healthy Controls. <i>Noropsikiyatri Arsivi</i> , 2018 , 55, 3-8	0.6	2
68	GC and VDR SNPs and Vitamin D Levels in Parkinson's Disease: The Relevance to Clinical Features. <i>NeuroMolecular Medicine</i> , 2017 , 19, 24-40	4.6	29
67	The p.Thr11Met mutation in c19orf12 is frequent among adult Turkish patients with MPAN. <i>Parkinsonism and Related Disorders</i> , 2017 , 39, 64-70	3.6	21
66	The Vascular Impairment of Cognition Classification Consensus Study. <i>Alzheimer's and Dementia</i> , 2017 , 13, 624-633	1.2	106
65	Cognitive and anatomical correlates of anosognosia in amnesic mild cognitive impairment and early-stage Alzheimer's disease. <i>International Psychogeriatrics</i> , 2017 , 29, 293-302	3.4	11

64	Mutations in TYROBP are not a common cause of dementia in a Turkish cohort. <i>Neurobiology of Aging</i> , 2017 , 58, 240.e1-240.e3	5.6	5
63	SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey. <i>Neurological Sciences</i> , 2017 , 38, 2203-2207	3.5	6
62	[P2047]: A COMPARISON OF AFFECTIVE THEORY OF MIND ABILITIES BETWEEN PARTICIPANTS WITH AMNESTIC MILD COGNITIVE IMPAIRMENT AND HEALTHY CONTROLS 2017 , 13, P808-P809		
61	[P1090]: THE RELATIONSHIP BETWEEN CSF AMYLOID BETA CONCENTRATIONS AND FREE AND CUED RECALL PERFORMANCE AMONG PARTICIPANTS WITH AMNESTIC MILD COGNITIVE IMPAIRMENT 2017 , 13, P362-P362		
60	[P1043]: INTERCORRELATIONS BETWEEN CSF AMYLOID BETA LEVELS AND NEUROPSYCHOLOGICAL PROFILES AMONG PARTICIPANTS WITH SUBJECTIVE COGNITIVE IMPAIRMENT AND MCI OF THE AMNESTIC TYPE 2017 , 13, P387-P388		
59	[P2028]: 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) 2017 , 13, P744-P745		
58	INCIDENTAL PAGET'S DISEASE DISGUISED AS BONE METASTASIS IN A PATIENT WITH ENDOMETRIUM CARCINOMA. <i>Acta Endocrinologica</i> , 2017 , 13, 111-114	0.9	
57	First-episode psychotic disorder improving after immunotherapy. <i>Acta Neurologica Belgica</i> , 2016 , 116, 113-4	1.5	3
56	Are There Any Specific EEG Findings in Autoimmune Epilepsies?. <i>Clinical EEG and Neuroscience</i> , 2016 , 47, 224-34	2.3	42
55	Achromatic temporal-frequency responses of human lateral geniculate nucleus and primary visual cortex. <i>Vision Research</i> , 2016 , 127, 177-185	2.1	5
54	Analysis of the tremor in juvenile myoclonic epilepsy. <i>Epilepsy Research</i> , 2016 , 128, 140-148	3	4
53	Volumetric differences suggest involvement of cerebellum and brainstem in chronic migraine. <i>Cephalalgia</i> , 2016 , 36, 301-8	6.1	42
52	Expression changes of genes associated with apoptosis and survival processes in Parkinson's disease. <i>Neuroscience Letters</i> , 2016 , 615, 72-7	3.3	14
51	A novel homozygous DJ1 mutation causes parkinsonism and ALS in a Turkish family. <i>Parkinsonism and Related Disorders</i> , 2016 , 29, 117-20	3.6	16
50	Adult-onset phenylketonuria with rapidly progressive dementia and parkinsonism. <i>Neurocase</i> , 2016 , 22, 273-5	0.8	7
49	Loss of VPS13C Function in Autosomal-Recessive Parkinsonism Causes Mitochondrial Dysfunction and Increases PINK1/Parkin-Dependent Mitophagy. <i>American Journal of Human Genetics</i> , 2016 , 98, 500-513	11.3	225
48	Comparison of Cognitive Parameters Between Bilateral and Unilateral Hippocampal Sclerosis. <i>Noropsikiyatri Arsivi</i> , 2016 , 53, 199-204	0.4	3
47	PLA2G6 Mutations Related to Distinct Phenotypes: A New Case with Early-onset Parkinsonism. <i>Tremor and Other Hyperkinetic Movements</i> , 2016 , 6, 363	2	7

46	NMDA receptor encephalitis with cancer of unknown primary origin. <i>Tumori</i> , 2016 , 102,	1.7	7
45	Mutation Frequency of the Major Frontotemporal Dementia Genes, MAPT, GRN and C9ORF72 in a Turkish Cohort of Dementia Patients. <i>PLoS ONE</i> , 2016 , 11, e0162592	3.7	16
44	Vitamin D deficiency might pose a greater risk for ApoEε4 non-carrier Alzheimer's disease patients. <i>Neurological Sciences</i> , 2016 , 37, 1633-43	3.5	23
43	The Association Between Clusterin and APOE Polymorphisms and Late-Onset Alzheimer Disease in a Turkish Cohort. <i>Journal of Geriatric Psychiatry and Neurology</i> , 2016 , 29, 221-6	3.8	16
42	Secondary paroxysmal kinesigenic dyskinesia associated with CLCN2 gene mutation. <i>Parkinsonism and Related Disorders</i> , 2015 , 21, 544-6	3.6	14
41	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		1
40	Parkinson's Disease Dementia. <i>Neuropsychiatric Symptoms of Neurological Disease</i> , 2015 , 53-77		
39	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. <i>Journal of Neuroimmunology</i> , 2015 , 283, 50-7	3.5	145
38	P1-052: Association between clusterin polymorphisms and Alzheimer's disease 2015 , 11, P358-P358		
37	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		
36	The distinct genetic pattern of ALS in Turkey and novel mutations. <i>Neurobiology of Aging</i> , 2015 , 36, 1764.e9-1764.e18	3.5	18
35	Cognitive impairment and dementia in Parkinson's disease: practical issues and management. <i>Movement Disorders</i> , 2014 , 29, 663-72	7	56
34	FBXO7-R498X mutation: phenotypic variability from chorea to early onset parkinsonism within a family. <i>Parkinsonism and Related Disorders</i> , 2014 , 20, 1253-6	3.6	24
33	P1-287: NEUROIMAGING FINDINGS OF NASU-HAKOLA DISEASE 2014 , 10, P415-P415		
32	P1-165: EVALUATION OF INCIDENCE AND CLINICAL FEATURES OF ANTIBODY-ASSOCIATED AUTOIMMUNE ENCEPHALITIS MIMICKING DEMENTIA 2014 , 10, P360-P360		
31	Evaluation of incidence and clinical features of antibody-associated autoimmune encephalitis mimicking dementia. <i>Behavioural Neurology</i> , 2014 , 2014, 935379	3	19
30	Silent neurological involvement in biopsy-defined coeliac patients. <i>Neurological Sciences</i> , 2013 , 34, 2199-204	3.9	14
29	Non-convulsive status epilepticus associated with glutamic acid decarboxylase antibody. <i>Clinical EEG and Neuroscience</i> , 2013 , 44, 232-6	2.3	16

28	BDNF, TNF α , HSP90, CFH, and IL-10 serum levels in patients with early or late onset Alzheimer's disease or mild cognitive impairment. <i>Journal of Alzheimer's Disease</i> , 2013 , 37, 185-95	4.3	124
27	TREM2 variants in Alzheimer's disease. <i>New England Journal of Medicine</i> , 2013 , 368, 117-27	59.2	1805
26	Novel compound heterozygous mutation in TREM2 found in a Turkish frontotemporal dementia-like family. <i>Neurobiology of Aging</i> , 2013 , 34, 2890.e1-5	5.6	90
25	Using exome sequencing to reveal mutations in TREM2 presenting as a frontotemporal dementia-like syndrome without bone involvement. <i>JAMA Neurology</i> , 2013 , 70, 78-84	17.2	257
24	Reduced orexin-A levels in frontotemporal dementia: possible association with sleep disturbance. <i>American Journal of Alzheimer's Disease and Other Dementias</i> , 2013 , 28, 606-11	2.5	21
23	Neuroimaging, biomarkers, and management of dementia with lewy bodies. <i>Frontiers in Neurology</i> , 2013 , 4, 151	4.1	6
22	The Relationship of White Matter Hyperintensities with Depressive Symptoms and Daily Living Activities in Early-Stage Alzheimer's Disease Patients. <i>Noropsikiyatri Arsivi</i> , 2013 , 50, 360-363	0.4	2
21	Genetic bases and phenotypes of autosomal recessive Parkinson disease in a Turkish population. <i>European Journal of Neurology</i> , 2012 , 19, 769-75	6	14
20	Exome sequencing reveals an unexpected genetic cause of disease: NOTCH3 mutation in a Turkish family with Alzheimer's disease. <i>Neurobiology of Aging</i> , 2012 , 33, 1008.e17-23	5.6	72
19	Identification of PSEN1 and PSEN2 gene mutations and variants in Turkish dementia patients. <i>Neurobiology of Aging</i> , 2012 , 33, 1850.e17-27	5.6	31
18	Differentiating symptomatic Parkin mutations carriers from patients with idiopathic Parkinson's disease: contribution of automated segmentation neuroimaging method. <i>Parkinsonism and Related Disorders</i> , 2012 , 18, 562-6	3.6	10
17	Dementia and behavioral neurology: recent advances. <i>Journal of Neurology</i> , 2012 , 259, 1006-10	5.5	1
16	Vitamin D receptor gene haplotype is associated with late-onset Alzheimer's disease. <i>Tohoku Journal of Experimental Medicine</i> , 2012 , 228, 189-96	2.4	64
15	Simultaneous EEG/fMRI analysis of the resonance phenomena in steady-state visual evoked responses. <i>Clinical EEG and Neuroscience</i> , 2011 , 42, 98-106	2.3	36
14	İlhan Bulgular ile Seyreden bir Marchiafava-Bignami Hastası Tanısında Diffzyon Ağıllı Manyetik Rezonans Gütlemenin Katkısı <i>Noropsikiyatri Arsivi</i> , 2011 , 48, 277-280	0.4	1
13	Impact of Earthquake on Multiple Sclerosis Attacks. <i>Noropsikiyatri Arsivi</i> , 2011 , 48, 1-1	0.4	
12	Interleukin-1alpha -889 C/T polymorphism in Turkish patients with late-onset Alzheimer's disease. <i>Dementia and Geriatric Cognitive Disorders</i> , 2009 , 27, 82-7	2.6	11
11	The prevalence of dementia in an urban Turkish population. <i>American Journal of Alzheimer's Disease and Other Dementias</i> , 2008 , 23, 67-76	2.5	47

10	Clinical and magnetic resonance imaging findings of HIV-negative patients with neurosyphilis. <i>Journal of Neurology</i> , 2007 , 254, 368-74	5.5	26
9	The attitude of elderly lay people towards the symptoms of dementia. <i>International Psychogeriatrics</i> , 2006 , 18, 251-8	3.4	20
8	OC1Bilateral central giant cell granuloma of the mandible: a case report. <i>Oral Diseases</i> , 2006 , 12, 10-10	3.5	1
7	Unrecognized depression in community-dwelling elderly persons in Istanbul. <i>International Psychogeriatrics</i> , 2005 , 17, 303-12	3.4	22
6	Obsessive-compulsive disorder secondary to bilateral frontal damage due to a closed head injury. <i>Cognitive and Behavioral Neurology</i> , 2004 , 17, 118-20	1.6	11
5	Clozapine treatment in oromandibular dystonia. <i>Clinical Neuropharmacology</i> , 2004 , 27, 84-6	1.4	13
4	Risk factors for Alzheimer disease: a population-based case-control study in Istanbul, Turkey. <i>Alzheimer Disease and Associated Disorders</i> , 2003 , 17, 139-45	2.5	38
3	Therapeutic effects of an acetylcholinesterase inhibitor (donepezil) on memory in Wernicke-Korsakoff's disease. <i>Clinical Neuropharmacology</i> , 2002 , 25, 16-20	1.4	15
2	Perioral myoclonia with absence seizures: a rare epileptic syndrome. <i>Epileptic Disorders</i> , 2001 , 3, 23-7	1.9	12
1	Parkinson disease dementia 177-191		