Basar Bilgic

List of Publications by Citations

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	TREM2 variants in Alzheimer's disease. *New England Journal of Medicine*, **2013**, 368, 117-27	59.2	1805
134	Using exome sequencing to reveal mutations in TREM2 presenting as a frontotemporal dementia-like syndrome without bone involvement. *JAMA Neurology*, **2013**, 70, 78-84	17.2	257
133	Loss of VPS13C Function in Autosomal-Recessive Parkinsonism Causes Mitochondrial Dysfunction and Increases PINK1/Parkin-Dependent Mitophagy. *American Journal of Human Genetics*, **2016**, 98, 500-513	11	225
132	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, 50-7	3.5	145
131	Progress toward standardized diagnosis of vascular cognitive impairment: Guidelines from the Vascular Impairment of Cognition Classification Consensus Study. *Alzheimer's and Dementia*, **2018**, 14, 280-292	1.2	136
130	BDNF, TNF□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, 185-95	4.3	124
129	The Vascular Impairment of Cognition Classification Consensus Study. *Alzheimer's and Dementia*, **2017**, 13, 624-633	1.2	106
128	Novel compound heterozygous mutation in TREM2 found in a Turkish frontotemporal dementia-like family. *Neurobiology of Aging*, **2013**, 34, 2890.e1-5	5.6	90
127	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.e17-23	5.6	72
126	The distinct genetic pattern of ALS in Turkey and novel mutations. *Neurobiology of Aging*, **2015**, 36, 1764.e9-1764.e18	5.6	69
125	Vitamin D receptor gene haplotype is associated with late-onset Alzheimer's disease. *Tohoku Journal of Experimental Medicine*, **2012**, 228, 189-96	2.4	64
124	Cognitive impairment and dementia in Parkinson's disease: practical issues and management. *Movement Disorders*, **2014**, 29, 663-72	7	56
123	The prevalence of dementia in an urban Turkish population. *American Journal of Alzheimer's Disease and Other Dementias*, **2008**, 23, 67-76	2.5	47
122	Are There Any Specific EEG Findings in Autoimmune Epilepsies?. *Clinical EEG and Neuroscience*, **2016**, 47, 224-34	2.3	42
121	Volumetric differences suggest involvement of cerebellum and brainstem in chronic migraine. *Cephalalgia*, **2016**, 36, 301-8	6.1	42
120	Risk factors for Alzheimer disease: a population-based case-control study in Istanbul, Turkey. *Alzheimer Disease and Associated Disorders*, **2003**, 17, 139-45	2.5	38
119	Simultaneous EEG/fMRI analysis of the resonance phenomena in steady-state visual evoked responses. *Clinical EEG and Neuroscience*, **2011**, 42, 98-106	2.3	36

118	Identification of PSEN1 and PSEN2 gene mutations and variants in Turkish dementia patients. *Neurobiology of Aging*, **2012**, 33, 1850.e17-27	5.6	31
117	GC and VDR SNPs and Vitamin D Levels in Parkinson's Disease: The Relevance to Clinical Features. *NeuroMolecular Medicine*, **2017**, 19, 24-40	4.6	29
116	Clinical and magnetic resonance imaging findings of HIV-negative patients with neurosyphilis. *Journal of Neurology*, **2007**, 254, 368-74	5.5	26
115	FBXO7-R498X mutation: phenotypic variability from chorea to early onset parkinsonism within a family. *Parkinsonism and Related Disorders*, **2014**, 20, 1253-6	3.6	24
114	Parkinson's Disease Dementia and Lewy Body Disease. *Seminars in Neurology*, **2019**, 39, 274-282	3.2	23
113	Vitamin D deficiency might pose a greater risk for ApoEe4 non-carrier Alzheimer's disease patients. *Neurological Sciences*, **2016**, 37, 1633-43	3.5	23
112	Unrecognized depression in community-dwelling elderly persons in Istanbul. *International Psychogeriatrics*, **2005**, 17, 303-12	3.4	22
111	The p.Thr11Met mutation in c19orf12 is frequent among adult Turkish patients with MPAN. *Parkinsonism and Related Disorders*, **2017**, 39, 64-70	3.6	21
110	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	6	21
109	Reduced orexin-A levels in frontotemporal dementia: possible association with sleep disturbance. *American Journal of Alzheimer's Disease and Other Dementias*, **2013**, 28, 606-11	2.5	21
108	The attitude of elderly lay people towards the symptoms of dementia. *International Psychogeriatrics*, **2006**, 18, 251-8	3.4	20
107	HPCA confirmed as a genetic cause of DYT2-like dystonia phenotype. *Movement Disorders*, **2018**, 33, 1354-1358	7	19
106	Evaluation of incidence and clinical features of antibody-associated autoimmune encephalitis mimicking dementia. *Behavioural Neurology*, **2014**, 2014, 935379	3	19
105	A novel homozygous DJ1 mutation causes parkinsonism and ALS in a Turkish family. *Parkinsonism and Related Disorders*, **2016**, 29, 117-20	3.6	16
104	Neuroimaging Findings in Sepsis-Induced Brain Dysfunction: Association with Clinical and Laboratory Findings. *Neurocritical Care*, **2019**, 30, 106-117	3.3	16
103	Non-convulsive status epilepticus associated with glutamic acid decarboxylase antibody. *Clinical EEG and Neuroscience*, **2013**, 44, 232-6	2.3	16
102	Mutation Frequency of the Major Frontotemporal Dementia Genes, MAPT, GRN and C9ORF72 in a Turkish Cohort of Dementia Patients. *PLoS ONE*, **2016**, 11, e0162592	3.7	16
101	The Association Between Clusterin and APOE Polymorphisms and Late-Onset Alzheimer Disease in a Turkish Cohort. *Journal of Geriatric Psychiatry and Neurology*, **2016**, 29, 221-6	3.8	16

100	Therapeutic effects of an acetylcholinesterase inhibitor (donepezil) on memory in Wernicke-Korsakoff's disease. *Clinical Neuropharmacology*, **2002**, 25, 16-20	1.4	15
99	Secondary paroxysmal kinesigenic dyskinesia associated with CLCN2 gene mutation. *Parkinsonism and Related Disorders*, **2015**, 21, 544-6	3.6	14
98	Expression changes of genes associated with apoptosis and survival processes in Parkinson's disease. *Neuroscience Letters*, **2016**, 615, 72-7	3.3	14
97	Genetic bases and phenotypes of autosomal recessive Parkinson disease in a Turkish population. *European Journal of Neurology*, **2012**, 19, 769-75	6	14
96	Silent neurological involvement in biopsy-defined coeliac patients. *Neurological Sciences*, **2013**, 34, 2199-204	3.3	14
95	Which cognitive dual-task walking causes most interference on the Timed Up and Go test in Parkinson's disease: a controlled study. *Neurological Sciences*, **2018**, 39, 2151-2157	3.5	14
94	Clozapine treatment in oromandibular dystonia. *Clinical Neuropharmacology*, **2004**, 27, 84-6	1.4	13
93	Perioral myoclonia with absence seizures: a rare epileptic syndrome. *Epileptic Disorders*, **2001**, 3, 23-7	1.9	12
92	Cognitive and anatomical correlates of anosognosia in amnestic mild cognitive impairment and early-stage Alzheimer's disease. *International Psychogeriatrics*, **2017**, 29, 293-302	3.4	11
91	Characterization of recessive Parkinson's disease in a large multicenter study. *Annals of Neurology*, **2020**, 88, 843	9.4	11
90	Interleukin-1alpha -889 C/T polymorphism in Turkish patients with late-onset Alzheimer's disease. *Dementia and Geriatric Cognitive Disorders*, **2009**, 27, 82-7	2.6	11
89	Obsessive-compulsive disorder secondary to bilateral frontal damage due to a closed head injury. *Cognitive and Behavioral Neurology*, **2004**, 17, 118-20	1.6	11
88	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	2.6	10
87	Differentiating symptomatic Parkin mutations carriers from patients with idiopathic Parkinson's disease: contribution of automated segmentation neuroimaging method. *Parkinsonism and Related Disorders*, **2012**, 18, 562-6	3.6	10
86	Brain Volume Changes in Patients with Acute Brain Dysfunction Due to Sepsis. *Neurocritical Care*, **2020**, 32, 459-468	3.3	9
85	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	1.5	9
84	Adult-onset phenylketonuria with rapidly progressive dementia and parkinsonism. *Neurocase*, **2016**, 22, 273-5	0.8	7
83	PLA2G6 Mutations Related to Distinct Phenotypes: A New Case with Early-onset Parkinsonism. *Tremor and Other Hyperkinetic Movements*, **2016**, 6, 363	2	7

82	NMDA receptor encephalitis with cancer of unknown primary origin. *Tumori*, **2016**, 102,	1.7	7
81	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	4.3	7
80	SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey. *Neurological Sciences*, **2017**, 38, 2203-2207	3.5	6
79	Neuroimaging, biomarkers, and management of dementia with lewy bodies. *Frontiers in Neurology*, **2013**, 4, 151	4.1	6
78	Achromatic temporal-frequency responses of human lateral geniculate nucleus and primary visual cortex. *Vision Research*, **2016**, 127, 177-185	2.1	5
77	Mutations in TYROBP are not a common cause of dementia in a Turkish cohort. *Neurobiology of Aging*, **2017**, 58, 240.e1-240.e3	5.6	5
76	The Right Temporal Variant of Frontotemporal Dementia Is Not Genetically Sporadic: A Case Series. *Journal of Alzheimer's Disease*, **2021**, 79, 1195-1201	4.3	5
75	Association between selected cholesterol-related gene polymorphisms and Alzheimer's disease in a Turkish cohort. *Molecular Biology Reports*, **2019**, 46, 1701-1707	2.8	4
74	Revisiting the complex architecture of ALS in Turkey: Expanding genotypes, shared phenotypes, molecular networks, and a public variant database. *Human Mutation*, **2020**, 41, e7-e45	4.7	4
73	Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey. *Parkinsonism and Related Disorders*, **2018**, 48, 34-39	3.6	4
72	Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span. *Brain and Development*, **2018**, 40, 458-464	2.2	4
71	Analysis of the tremor in juvenile myoclonic epilepsy. *Epilepsy Research*, **2016**, 128, 140-148	3	4
70	The cerebral blood flow deficits in Parkinson's disease with mild cognitive impairment using arterial spin labeling MRI. *Journal of Neural Transmission*, **2020**, 127, 1285-1294	4.3	4
69	Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis. *Neurodegenerative Diseases*, **2019**, 19, 218-224	2.3	4
68	Olfactory function and olfactory bulb volume in Wilson's disease. *European Archives of Oto-Rhino-Laryngology*, **2019**, 276, 139-142	3.5	4
67	A novel PNPLA6 mutation in a Turkish family with intractable Holmes tremor and spastic ataxia. *Neurological Sciences*, **2021**, 42, 1535-1539	3.5	4
66	Inflammation and regulatory T cell genes are differentially expressed in peripheral blood mononuclear cells of Parkinson's disease patients. *Scientific Reports*, **2021**, 11, 2316	4.9	4
65	First-episode psychotic disorder improving after immunotherapy. *Acta Neurologica Belgica*, **2016**, 116, 113-4	1.5	3

64	White-matter changes in early and late stages of mild cognitive impairment. *Journal of Clinical Neuroscience*, **2020**, 78, 181-184	2.2	3
63	A patient with early-onset Alzheimer's disease with a novel PSEN1 p.Leu424Pro mutation. *Neurobiology of Aging*, **2019**, 84, 238.e1-238.e4	5.6	3
62	Comparison of Cognitive Parameters Between Bilateral and Unilateral Hippocampal Sclerosis. *Noropsikiyatri Arsivi*, **2016**, 53, 199-204	0.4	3
61	Peripheral TREM2 mRNA levels in early and late-onset Alzheimer disease's patients. *Molecular Biology Reports*, **2020**, 47, 5903-5909	2.8	3
60	The association between repeat number in C9orf72 and phenotypic variability in Turkish patients with frontotemporal lobar degeneration. *Neurobiology of Aging*, **2019**, 76, 216.e1-216.e7	5.6	3
59	The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice. *Movement Disorders*, **2021**, 36, 1676-1688	7	3
58	Late-onset phenotype associated with a homozygous GJC2 missense mutation in a Turkish family. *Parkinsonism and Related Disorders*, **2019**, 66, 228-231	3.6	2
57	Functional neural substrates of football fanaticism: Different pattern of brain responses and connectivity in fanatics. *Psychiatry and Clinical Neurosciences*, **2020**, 74, 480-487	6.2	2
56	Wernicke's Encephalopathy due to Non-Alcoholic Gastrointestinal Tract Disease. *Noropsikiyatri Arsivi*, **2018**, 55, 307-314	0.6	2
55	The Relationship of White Matter Hyperintensities with Depressive Symptoms and Daily Living Activities in Early-Stage Alzheimer's Disease Patients. *Noropsikiyatri Arsivi*, **2013**, 50, 360-363	0.4	2
54	The COVID-19 from Neurological Overview. *Turk Noroloji Dergisi = Turkish Journal of Neurology*, **2020**, 26, 58-108	1.1	2
53	Neuroinflammation Mediators are Reduced in Sera of Parkinson's Disease Patients with Mild Cognitive Impairment. *Noropsikiyatri Arsivi*, **2020**, 57, 15-17	0.6	2
52	Evaluation of OnabotulinumtoxinA Treatment in Patients with Concomitant Chronic Migraine and Temporomandibular Disorders. *Noropsikiyatri Arsivi*, **2018**, 55, 330-336	0.6	2
51	Amyloid Beta Adsorption Problem with Transfer Plates in Amyloid Beta 1-42 IVD Kits. *Journal of Molecular Neuroscience*, **2019**, 67, 534-539	3.3	2
50	A Comparison Of The Relations Between Brain Atrophy, Cognition And Optic Coherance Tomography Between Multiple Sclerosis Patients And Healthy Controls. *Noropsikiyatri Arsivi*, **2018**, 55, 3-8	0.6	2
49	Analysis of copy number variation in a Turkish dementia cohort. *Alzheimer's and Dementia*, **2020**, 16, e044868	12.8	1
48	Small ubiquitin-related modifier (SUMO) 3 and SUMO4 gene polymorphisms in Parkinson's disease. *Neurological Research*, **2020**, 42, 451-457	2.7	1
47	Laboratory and clinical correlates of brain atrophy in Neuro-Behçet's disease. *Journal of the Neurological Sciences*, **2020**, 414, 116831	3.2	1

46	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
45	Dementia and behavioral neurology: recent advances. *Journal of Neurology*, **2012**, 259, 1006-10	5.5	1
44	Ilen Bulgular ile Seyreden bir Marchiafava-Bignami Hastasen Tansda Diffzyon Aklk Manyetik Rezonans Gtlemenin Katks *Noropsikiyatri Arsivi*, **2011**, 48, 277-280	0.4	1
43	OC1Bilateral central giant cell granuloma of the mandible: a case report. *Oral Diseases*, **2006**, 12, 10-10	3.5	1
42	Neuroimaging and Dementia. *Noropsikiyatri Arsivi*, **2018**, 55, 1-2	0.6	1
41	PHACTR1 genetic variability is not critical in small vessel ischemic disease patients and PcomA recruitment in C57BL/6J mice. *Scientific Reports*, **2021**, 11, 6072	4.9	1
40	TREM2 variants as a possible cause of frontotemporal dementia with distinct neuroimaging features. *European Journal of Neurology*, **2021**, 28, 2603-2613	6	1
39	Do Alzheimers Patients Appear Younger than Their AgeA Study with Automatic Facial Age Estimation **2019**,		1
38	Hotspot Mutations in Pleuropulmonary Blastoma: A Case Series From a Tertiary Center. *Pediatric and Developmental Pathology*, **2020**, 23, 204-209	2.2	1
37	Detection of visual and frontoparietal network perfusion deficits in Parkinson's disease dementia. *European Journal of Radiology*, **2021**, 144, 109985	4.7	1
36	Neurological features and outcomes of Wilson's disease: a single-center experience. *Neurological Sciences*, **2021**, 42, 3829-3834	3.5	1
35	Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey. *Parkinsonism and Related Disorders*, **2021**, 93, 35-39	3.6	0
34	Genetic variants of vitamin D metabolism-related locus and gene are associated with clinical features of Parkinson's disease. *International Journal of Neuroscience*, **2020**, 1-11	2	0
33	A novel PSEN2 p.Ser175Phe variant in a family with Alzheimer's disease. *Neurological Sciences*, **2021**, 42, 2497-2504	3.5	0
32	Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers. *Journal of Parkinsons Disease*, **2021**, 11, 559-568	5.3	0
31	A novel SACS p.Pro4154GlnfsTer20 mutation in a family with autosomal recessive spastic ataxia of Charlevoix-Saguenay. *Neurological Sciences*, **2021**, 42, 2969-2973	3.5	0
30	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
29	Do Alzheimer's Disease Patients Appear Younger than Their Real Age?. *Dementia and Geriatric Cognitive Disorders*, **2020**, 49, 483-488	2.6	

28	Multicenter study of levodopa carbidopa intestinal gel in Parkinson disease: the Turkish experience. *Turkish Journal of Medical Sciences*, **2020**, 50, 66-85	2.7
27	P1-287: NEUROIMAGING FINDINGS OF NASU-HAKOLA DISEASE **2014**, 10, P415-P415	
26	P1-165: EVALUATION OF INCIDENCE AND CLINICAL FEATURES OF ANTIBODY-ASSOCIATED AUTOIMMUNE ENCEPHALITIS MIMICKING DEMENTIA **2014**, 10, P360-P360	
25	[P2047]: A COMPARISON OF AFFECTIVE THEORY OF MIND ABILITIES BETWEEN PARTICIPANTS WITH AMNESTIC MILD COGNITIVE IMPAIRMENT AND HEALTHY CONTROLS **2017**, 13, P808-P809	
24	[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	
23	[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	
22	[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	
21	Parkinson Disease Dementia. *Neuropsychiatric Symptoms of Neurological Disease*, **2015**, 53-77	
20	P1-052: Association between clusterin polymorphisms and Alzheimer's disease **2015**, 11, P358-P358	
19	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	
18	Parkinson disease dementia177-191	
17	Impact of Earthquake on Multiple Sclerosis Attacks. *Noropsikiyatri Arsivi*, **2011**, 48, 1-1	0.4
16	The association of serum clusterin levels and Clusterin rs11136000 polymorphisms with Alzheimer disease in a Turkish cohort. *Neurological Sciences and Neurophysiology*, **2020**, 37, 134	0.4
15	INCIDENTAL PAGET'S DISEASE DISGUISED AS BONE METASTASIS IN A PATIENT WITH ENDOMETRIUM CARCINOMA. *Acta Endocrinologica*, **2017**, 13, 111-114	0.9
14	Cerebellar ataxia, neuropathy and vestibular areflexia syndrome (canvas): an important cause of late-onset ataxia with unique clinical features. *Acta Neurologica Belgica*, **2021**, 1	1.5
13	Intensive voice treatment (the Lee Silverman Voice Treatment [LSVTLOUD]) for individuals with Wilson's disease and adult cerebral palsy: two case reports. *Logopedics Phoniatrics Vocology*, **2021**, 1-9	1.3
12	B-Tensor: Brain Connectome Tensor Factorization for Alzheimer's Disease. *IEEE Journal of Biomedical and Health Informatics*, **2021**, 25, 1591-1600	7.2
11	P3-109: ASSOCIATION BETWEEN NADSYN1/DHCR7 AND CYP2R1 GENOTYPES AND PARKINSON'S DISEASE AND ITS CLINICAL FEATURES **2018**, 14, P1109-P1109	

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10	P4-085: CATATONIA AS CLINICAL PRESENTATION OF ANTI-N-METHYL-D-ASPARTATE (ANTI-NMDA) RECEPTOR ENCEPHALITIS **2018**, 14, P1467-P1468	
9	P2-229: THE EFFECT OF CSF AMYLOID BETA CONCENTRATIONS ON MEMORY PERFORMANCE OF THE INDIVIDUALS WITH SUBJECTIVE COGNITIVE IMPAIRMENT **2018**, 14, P756-P756	
8	P2-302: CLINICAL FEATURES AND DIAGNOSIS OF EARLY ONSET DEGENERATIVE DEMENTIAS IN TURKEY **2018**, 14, P798-P798	
7	P4-087: SPORADIC FATAL INSOMNIA IN A YOUNG TURKISH MAN **2018**, 14, P1468-P1469	
6	A comprehensive analysis of copy number variation in a Turkish dementia cohort. *Human Genomics*, **2021**, 15, 48	6.8
5	Frequency of frontotemporal dementia-related gene variants in Turkey. *Neurobiology of Aging*, **2021**, 106, 332.e1-332.e11	5.6
4	Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment.. *Acta Neurologica Belgica*, **2022**, 1	1.5
3	Does transcranial direct current stimulation enhance cognitive performance in Parkinson's disease mild cognitive impairment? An event-related potentials and neuropsychological assessment study.. *Neurological Sciences*, **2022**, 1	3.5
2	Genotype-Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review.. *BMC Neurology*, **2022**, 22, 122	3.1
1	Homozygosity analysis in a Turkish dementia cohort.. *Alzheimer's and Dementia*, **2021**, 17 Suppl 3, e054052	12