Basar Bilgic

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/493829/publications.pdf

Version: 2024-02-01

140 papers	5,673 citations	218662 26 h-index	82542 72 g-index
148 all docs	148 docs citations	148 times ranked	9731 citing authors

#	Article	IF	Citations
1	TREM2 Variants in Alzheimer's Disease. New England Journal of Medicine, 2013, 368, 117-127.	27.0	2,385
2	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.	6.2	333
3	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
4	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.	0.8	246
5	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
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	The Vascular Impairment of Cognition Classification Consensus Study. Alzheimer's and Dementia, 2017, 13, 624-633.	0.8	143
8	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
9	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
10	The distinct genetic pattern of ALS in Turkey and novel mutations. Neurobiology of Aging, 2015, 36, 1764.e9-1764.e18.	3.1	78
11	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
12	Simultaneous EEG/fMRI Analysis of the Resonance Phenomena in Steady-State Visual Evoked Responses. Clinical EEG and Neuroscience, 2011, 42, 98-106.	1.7	71
13	The Prevalence of Dementia in an Urban Turkish Population. American Journal of Alzheimer's Disease and Other Dementias, 2008, 23, 67-76.	1.9	70
14	Cognitive impairment and dementia in Parkinson's disease: Practical issues and management. Movement Disorders, 2014, 29, 663-672.	3.9	69
15	Volumetric differences suggest involvement of cerebellum and brainstem in chronic migraine. Cephalalgia, 2016, 36, 301-308.	3.9	57
16	Are There Any Specific EEG Findings in Autoimmune Epilepsies?. Clinical EEG and Neuroscience, 2016, 47, 224-234.	1.7	55
17	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
18	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

#	Article	IF	Citations
19	Parkinson's Disease Dementia and Lewy Body Disease. Seminars in Neurology, 2019, 39, 274-282.	1.4	40
20	Characterization of Recessive Parkinson Disease in a Large Multicenter Study. Annals of Neurology, 2020, 88, 843-850.	5.3	40
21	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
22	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
23	Clinical and magnetic resonance imaging findings of HIV-Negative patients with neurosyphilis. Journal of Neurology, 2007, 254, 368-374.	3.6	34
24	The p.Thr11Met mutation in c19orf12 is frequent among adult Turkish patients with MPAN. Parkinsonism and Related Disorders, 2017, 39, 64-70.	2.2	31
25	‹i›HPCA‹/i› confirmed as a genetic cause of DYT2â€like dystonia phenotype. Movement Disorders, 2018, 33, 1354-1358.	3.9	31
26	Unrecognized depression in community-dwelling elderly persons in Istanbul. International Psychogeriatrics, 2005, 17, 303-312.	1.0	29
27	FBXO7â€“R498X mutation: Phenotypic variability from chorea to early onset parkinsonism within a family. Parkinsonism and Related Disorders, 2014, 20, 1253-1256.	2.2	29
28	Which cognitive dual-task walking causes most interference on the Timed Up and Go test in Parkinsonâ€™s disease: a controlled study. Neurological Sciences, 2018, 39, 2151-2157.	1.9	29
29	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
30	Neuroimaging Findings in Sepsis-Induced Brain Dysfunction: Association with Clinical and Laboratory Findings. Neurocritical Care, 2019, 30, 106-117.	2.4	28
31	Reduced Orexin-A Levels in Frontotemporal Dementia. American Journal of Alzheimer's Disease and Other Dementias, 2013, 28, 606-611.	1.9	27
32	The attitude of elderly lay people towards the symptoms of dementia. International Psychogeriatrics, 2006, 18, 251-258.	1.0	26
33	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
34	Evaluation of Incidence and Clinical Features of Antibody-Associated Autoimmune Encephalitis Mimicking Dementia. Behavioural Neurology, 2014, 2014, 1-4.	2.1	25
35	Brain Volume Changes in Patients with Acute Brain Dysfunction Due to Sepsis. Neurocritical Care, 2020, 32, 459-468.	2.4	24
36	A novel homozygous DJ1 mutation causes parkinsonism and ALS in a Turkish family. Parkinsonism and Related Disorders, 2016, 29, 117-120.	2.2	23

#	Article	IF	Citations
37	Expression changes of genes associated with apoptosis and survival processes in Parkinsonâ€™s disease. Neuroscience Letters, 2016, 615, 72-77.	2.1	22
38	Therapeutic Effects of an Acetylcholinesterase Inhibitor (Donepezil) on Memory in Wernickeâ€“Korsakoff's Disease. Clinical Neuropharmacology, 2002, 25, 16-20.	0.7	21
39	Clozapine Treatment in Oromandibular Dystonia. Clinical Neuropharmacology, 2004, 27, 84-86.	0.7	21
40	Genetic bases and phenotypes of autosomal recessive Parkinson disease in a Turkish population. European Journal of Neurology, 2012, 19, 769-775.	3.3	20
41	Secondary paroxysmal kinesigenic dyskinesia associated with CLCN2 gene mutation. Parkinsonism and Related Disorders, 2015, 21, 544-546.	2.2	20
42	Inflammation and regulatory T cell genes are differentially expressed in peripheral blood mononuclear cells of Parkinsonâ€™s disease patients. Scientific Reports, 2021, 11, 2316.	3.3	20
43	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
44	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
45	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
46	Non-Convulsive Status Epilepticus Associated With Glutamic Acid Decarboxylase Antibody. Clinical EEG and Neuroscience, 2013, 44, 232-236.	1.7	17
47	Adult-onset phenylketonuria with rapidly progressive dementia and parkinsonism. Neurocase, 2016, 22, 273-275.	0.6	17
48	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
49	Silent neurological involvement in biopsy-defined coeliac patients. Neurological Sciences, 2013, 34, 2199-2204.	1.9	15
50	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
51	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
52	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
53	Clinical phenotype of hereditary spastic paraplegia due to KIF1C gene mutations across life span. Brain and Development, 2018, 40, 458-464.	1.1	13
54	A novel PNPLA6 mutation in a Turkish family with intractable Holmes tremor and spastic ataxia. Neurological Sciences, 2021, 42, 1535-1539.	1.9	13

#	Article	IF	Citations
55	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
56	Perioral myoclonia with absence seizures: a rare epileptic syndrome. Epileptic Disorders, 2001, 3, 23-7.	1.3	12
57	Patients with Lately Diagnosed Cerebrotendinous Xanthomatosis. Neurodegenerative Diseases, 2019, 19, 218-224.	1.4	11
58	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
59	PLA2G6 Mutations Related to Distinct Phenotypes: A New Case with Early-onset Parkinsonism. Tremor and Other Hyperkinetic Movements, 2016, 6, 363.	2.0	11
60	Achromatic temporal-frequency responses of human lateral geniculate nucleus and primary visual cortex. Vision Research, 2016, 127, 177-185.	1.4	10
61	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.	2.5	10
62	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
63	The Complex Genetic Landscape of Hereditary Ataxias in Turkey and Implications in Clinical Practice. Movement Disorders, 2021, 36, 1676-1688.	3.9	9
64	Neurological features and outcomes of Wilsonâ€™s disease: a single-center experience. Neurological Sciences, 2021, 42, 3829-3834.	1.9	9
65	Neuroimaging, Biomarkers, and Management of Dementia with Lewy Bodies. Frontiers in Neurology, 2013, 4, 151.	2.4	8
66	NMDA Receptor Encephalitis with Cancer of Unknown Primary Origin. Tumori, 2016, 102, S3-S4.	1.1	8
67	SYNE1 related cerebellar ataxia presents with variable phenotypes in a consanguineous family from Turkey. Neurological Sciences, 2017, 38, 2203-2207.	1.9	8
68	Role of LRRK2 and SNCA in autosomal dominant Parkinson's disease in Turkey. Parkinsonism and Related Disorders, 2018, 48, 34-39.	2.2	8
69	Peripheral TREM2 mRNA levels in early and late-onset Alzheimer diseaseâ€™s patients. Molecular Biology Reports, 2020, 47, 5903-5909.	2.3	8
70	Small ubiquitin-related modifier (SUMO) 3 and SUMO4 gene polymorphisms in Parkinsonâ€™s disease. Neurological Research, 2020, 42, 451-457.	1.3	7
71	Detection of visual and frontoparietal network perfusion deficits in Parkinsonâ€™s disease dementia. European Journal of Radiology, 2021, 144, 109985.	2.6	7
72	The COVID-19 from Neurological Overview. Turk Noroloji Dergisi = Turkish Journal of Neurology, 2020, 26, 58-108.	0.3	7

#	Article	IF	Citations
73	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
74	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
75	White-matter changes in early and late stages of mild cognitive impairment. Journal of Clinical Neuroscience, 2020, 78, 181-184.	1.5	6
76	Late-onset phenotype associated with a homozygous GJC2 missense mutation in a Turkish family. Parkinsonism and Related Disorders, 2019, 66, 228-231.	2.2	5
77	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
78	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
79	Functional Connectivity Analysis in Heterozygous Glucocerebrosidase Mutation Carriers. Journal of Parkinson's Disease, 2021, 11, 559-568.	2.8	5
80	TREM2 variants as a possible cause of frontotemporal dementia with distinct neuroimaging features. European Journal of Neurology, 2021, 28, 2603-2613.	3.3	5
81	Comparison of Cognitive Parameters between Bilateral and Unilateral Hippocampal Sclerosis. Noropsikiyatri Arsivi, 2016, 53, 199-204.	0.7	5
82	Analysis of the tremor in juvenile myoclonic epilepsy. Epilepsy Research, 2016, 128, 140-148.	1.6	4
83	First-episode psychotic disorder improving after immunotherapy. Acta Neurologica Belgica, 2016, 116, 113-114.	1.1	4
84	A patient with early-onset Alzheimer's disease with a novel PSEN1 p.Leu424Pro mutation. Neurobiology of Aging, 2019, 84, 238.e1-238.e4.	3.1	4
85	Olfactory function and olfactory bulb volume in Wilsonâ€™s disease. European Archives of Oto-Rhino-Laryngology, 2019, 276, 139-142.	1.6	4
86	Neuroinflammation mediators are reduced in sera of Parkinson?s disease patients with mild cognitive impairment. Noropsikiyatri Arsivi, 2019, 57, 15-17.	0.3	4
87	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.	3.1	3
88	DICER1 Hotspot Mutations in Pleuropulmonary Blastoma: A Case Series From a Tertiary Center. Pediatric and Developmental Pathology, 2020, 23, 204-209.	1.0	3
89	Does transcranial direct current stimulation enhance cognitive performance in Parkinsonâ€™s disease mild cognitive impairment? An event-related potentials and neuropsychological assessment study. Neurological Sciences, 2022, 43, 4029-4044.	1.9	3
90	Dementia and behavioral neurology: recent advances. Journal of Neurology, 2012, 259, 1006-1010.	3.6	2

#	Article	IF	Citations
91	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
92	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
93	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
94	Multicenter study of levodopa carbidopa intestinal gel in Parkinson's disease: the Turkish experince. Turkish Journal of Medical Sciences, 2020, 50, 66-85.	0.9	2
95	Laboratory and clinical correlates of brain atrophy in Neuro-BehÃ§et's disease. Journal of the Neurological Sciences, 2020, 414, 116831.	0.6	2
96	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
97	PHACTR1 genetic variability is not critical in small vessel ischemic disease patients and PcomA recruitment in C57BL/6J mice. Scientific Reports, 2021, 11, 6072.	3.3	2
98	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
99	Wernickeâ€™s Encephalopathy due to Non-alcoholic Gastrointestinal Tract Disease. Noropsikiyatri Arsivi, 2018, 55, 307-314.	0.3	2
100	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
101	Evaluation of OnabotulinumtoxinA Treatment in Patients with Concomitant Chronic Migraine and Temporomandibular Disorders. Noropsikiyatri Arsivi, 2018, 55, 330-336.	0.3	2
102	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.	0.3	2
103	OC1 Bilateral central giant cell granuloma of the mandible: a case report. Oral Diseases, 2006, 12, 10-10.	3.0	1
104	IlÄ±mlÄ± Bulgular ile Seyreden bir Marchiafava-Bignami HastasÄ±nÄ±n TanÄ±sÄ±nda DiffÃ¼zyon AÄŸÄ±rlÄ±klÄ± Manyetik Rezonans GÃ¶rÃ¼ntÃ¼lemenin KatkÄ±sÄ±. Noropsikiyatri Arsivi, 2011, 48, 277-280.	0.7	1
105	Do Alzheimerâ€™s Patients Appear Younger than Their AgeÆ’ A Study with Automatic Facial Age Estimation. , 2019, , .		1
106	Analysis of copy number variation in a Turkish dementia cohort. Alzheimer's and Dementia, 2020, 16, e044868.	0.8	1
107	Do Alzheimerâ€™s Disease Patients Appear Younger than Their Real Age?. Dementia and Geriatric Cognitive Disorders, 2020, 49, 483-488.	1.5	1
108	B-Tensor: Brain Connectome Tensor Factorization for Alzheimer's Disease. IEEE Journal of Biomedical and Health Informatics, 2021, 25, 1591-1600.	6.3	1

#	Article	IF	Citations
109	A novel PSEN2 p.Ser175Phe variant in a family with Alzheimerâ€™s disease. Neurological Sciences, 2021, 42, 2497-2504.	1.9	1
110	Cerebellar ataxia, neuropathy and vestibular areflexia syndrome (canvas): an important cause of late-onset ataxia with unique clinical features. Acta Neurologica Belgica, 2022, 122, 939-945.	1.1	1
111	Frequency of frontotemporal dementia-related gene variants in Turkey. Neurobiology of Aging, 2021, 106, 332.e1-332.e11.	3.1	1
112	Neuroimaging and Dementia. Noropsikiyatri Arsivi, 2018, 55, 1-2.	0.3	1
113	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
114	Clinical and molecular genetic findings of hereditary Parkinson's patients from Turkey. Parkinsonism and Related Disorders, 2021, 93, 35-39.	2.2	1
115	Medication management and treatment adherence in Parkinson's disease patients with mild cognitive impairment. Acta Neurologica Belgica, 2023, 123, 823-829.	1.1	1
116	Genotypeâ€“Phenotype correlations of SCARB2 associated clinical presentation: a case report and in-depth literature review. BMC Neurology, 2022, 22, 122.	1.8	1
117	Impact of Earthquake on Multiple Sclerosis Attacks. Noropsikiyatri Arsivi, 2011, 48, 1-1.	0.7	0
118	Parkinsonâ€™s disease dementia. , 0, , 177-191.		0
119	P1-287: NEUROIMAGING FINDINGS OF NASU-HAKOLA DISEASE. , 2014, 10, P415-P415.		0
120	P1-165: EVALUATION OF INCIDENCE AND CLINICAL FEATURES OF ANTIBODY-ASSOCIATED AUTOIMMUNE ENCEPHALITIS MIMICKING DEMENTIA. , 2014, 10, P360-P360.		0
121	PSEN1 mutation presenting as posterior cortical atrophy. Journal of the Neurological Sciences, 2015, 357, e127.	0.6	0
122	Parkinsonâ€™s Disease Dementia. Neuropsychiatric Symptoms of Neurological Disease, 2015, , 53-77.	0.3	0
123	Group independent component analysis of alzheimer's disease and mild cognitive impairment patients. , 2015, , .		0
124	P1-052: Association between clusterin polymorphisms and Alzheimer's disease. , 2015, 11, P358-P358.		0
125	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
126	Task related modulation of group independent components of Alzheimer's disease and mild cognitive impairment patients. , 2015, , .		0

#	Article	IF	Citations
127	[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
128	[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
129	[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
130	[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
131	Determination of Diffusion Weighted Magnetic Resonance Imaging Based Biomarkers of Mild Cognitive Impairment in Parkinsonâ€™s Disease. , 2017, , .		0
132	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
133	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
134	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
135	P2â€“302: CLINICAL FEATURES AND DIAGNOSIS OF EARLY ONSET DEGENERATIVE DEMENTIAS IN TURKEY. Alzheimer's and Dementia, 2018, 14, P798.	0.8	0
136	P4â€“087: SPORADIC FATAL INSOMNIA IN A YOUNG TURKISH MAN. Alzheimer's and Dementia, 2018, 14, P1468.	0.8	0
137	A comprehensive analysis of copy number variation in a Turkish dementia cohort. Human Genomics, 2021, 15, 48.	2.9	0
138	Incidental Pagetâ€™s Disease Disguised as Bone Metastasis in a Patient with Endometrium Carcinoma. Acta Endocrinologica, 2017, 13, 111-114.	0.3	0
139	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
140	Homozygosity analysis in a Turkish dementia cohort.. Alzheimer's and Dementia, 2021, 17 Suppl 3, e054052.	0.8	0