

Tatiana Foroud

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.

280
papers

20,247
citations

58
h-index

140
g-index

330
ext. papers

25,983
ext. citations

8.2
avg, IF

5.35
L-index

#	Paper	IF	Citations
280	Role of Lysosomal Gene Variants in Modulating GBA-Associated Parkinson's Disease Risk.. <i>Movement Disorders</i> , 2022 ,	7	3
279	New insights into the genetic etiology of Alzheimer's disease and related dementias.. <i>Nature Genetics</i> , 2022 ,	36.3	27
278	Comprehensive cross-sectional and longitudinal analyses of plasma neurofilament light across FTD spectrum disorders.. <i>Cell Reports Medicine</i> , 2022 , 3, 100607	18	0
277	A genome-wide association study of interhemispheric theta EEG coherence: implications for neural connectivity and alcohol use behavior. <i>Molecular Psychiatry</i> , 2021 , 26, 5040-5052	15.1	6
276	Genetic Testing for Parkinson Disease: Are We Ready?. <i>Neurology: Clinical Practice</i> , 2021 , 11, 69-77	1.7	5
275	Obesity, Diabetes, Coffee, Tea, and Cannabis Use Alter Risk for Alcohol-Related Cirrhosis in 2 Large Cohorts of High-Risk Drinkers. <i>American Journal of Gastroenterology</i> , 2021 , 116, 106-115	0.7	7
274	A genetic risk score and diabetes predict development of alcohol-related cirrhosis in drinkers. <i>Journal of Hepatology</i> , 2021 ,	13.4	4
273	Biomarkers of neurodegeneration and glial activation validated in Alzheimer's disease assessed in longitudinal cerebrospinal fluid samples of Parkinson's disease. <i>PLoS ONE</i> , 2021 , 16, e0257372	3.7	2
272	Can Salivary Innate Immune Molecules Provide Clue on Taste Dysfunction in COVID-19?. <i>Frontiers in Microbiology</i> , 2021 , 12, 727430	5.7	
271	Cross-Sectional Exploration of Plasma Biomarkers of Alzheimer's Disease in Down Syndrome: Early Data from the Longitudinal Investigation for Enhancing Down Syndrome Research (LIFE-DSR) Study. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	4
270	Longitudinal Analysis of Multiple Neurotransmitter Metabolites in Cerebrospinal Fluid in Early Parkinson's Disease. <i>Movement Disorders</i> , 2021 , 36, 1972-1978	7	1
269	Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. <i>Nature Genetics</i> , 2021 , 53, 817-829	36.3	83
268	The Longitudinal Early-onset Alzheimer's Disease Study (LEADS): Framework and methodology. <i>Alzheimer's and Dementia</i> , 2021 ,	1.2	5
267	Genomewide Association Studies of LRRK2 Modifiers of Parkinson's Disease. <i>Annals of Neurology</i> , 2021 , 90, 76-88	9.4	9
266	Genome-wide association study of stimulant dependence. <i>Translational Psychiatry</i> , 2021 , 11, 363	8.6	0
265	Genome-wide association identifies the first risk loci for psychosis in Alzheimer disease. <i>Molecular Psychiatry</i> , 2021 ,	15.1	5
264	Polygenic contributions to alcohol use and alcohol use disorders across population-based and clinically ascertained samples. <i>Psychological Medicine</i> , 2021 , 51, 1147-1156	6.9	5

263	Testing influences of APOE and BDNF genes and heart failure on cognitive function. <i>Heart and Lung: Journal of Acute and Critical Care</i> , 2021 , 50, 51-58	2.6	
262	Genome-wide admixture mapping of DSM-IV alcohol dependence, criterion count, and the self-rating of the effects of ethanol in African American populations. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2021 , 186, 151-161	3.5	0
261	Plasma Total-Tau and Neurofilament Light Chain as Diagnostic Biomarkers of Alzheimer's Disease Dementia and Mild Cognitive Impairment in Adults with Down Syndrome. <i>Journal of Alzheimer's Disease</i> , 2021 , 79, 671-681	4.3	12
260	Novel Alzheimer Disease Risk Loci and Pathways in African American Individuals Using the African Genome Resources Panel: A Meta-analysis. <i>JAMA Neurology</i> , 2021 , 78, 102-113	17.2	32
259	Genome-wide Association Study and Meta-analysis on Alcohol-Associated Liver Cirrhosis Identifies Genetic Risk Factors. <i>Hepatology</i> , 2021 , 73, 1920-1931	11.2	18
258	Allele-specific expression and high-throughput reporter assay reveal functional genetic variants associated with alcohol use disorders. <i>Molecular Psychiatry</i> , 2021 , 26, 1142-1151	15.1	15
257	The Contribution of Known Familial Cardiovascular Disease Genes to Sudden Cardiac Death in Patients Undergoing Hemodialysis. <i>CardioRenal Medicine</i> , 2021 , 11, 174-183	2.8	
256	The association of polygenic risk for schizophrenia, bipolar disorder, and depression with neural connectivity in adolescents and young adults: examining developmental and sex differences. <i>Translational Psychiatry</i> , 2021 , 11, 54	8.6	3
255	Assessment of Blood Biomarker Profile After Acute Concussion During Combative Training Among US Military Cadets: A Prospective Study From the NCAA and US Department of Defense CARE Consortium. <i>JAMA Network Open</i> , 2021 , 4, e2037731	10.4	7
254	Genome sequencing analysis identifies new loci associated with Lewy body dementia and provides insights into its genetic architecture. <i>Nature Genetics</i> , 2021 , 53, 294-303	36.3	31
253	Multi-omics integration analysis identifies novel genes for alcoholism with potential overlap with neurodegenerative diseases. <i>Nature Communications</i> , 2021 , 12, 5071	17.4	4
252	Dopamine transporter imaging predicts clinically-defined β -synucleinopathy in REM sleep behavior disorder. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 201-212	5.3	6
251	Outcomes of genetic test disclosure and genetic counseling in a large Parkinson's disease research study. <i>Journal of Genetic Counseling</i> , 2021 , 30, 755-765	2.5	3
250	Longitudinal Early-onset Alzheimer's Disease Study (LEADS) genetic screening: Initial results.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e056493	1.2	
249	ADSP follow-up study: NCRAD biospecimens.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e056242	1.2	
248	Association of a locus on chromosome 17 with earlier age at onset of cognitive impairment in a familial Amish dataset.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e056288	1.2	
247	NCRAD Family Study and NIA-LOAD brain tissue: A NCRAD resource.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e056284	1.2	
246	Establishing a centralized repository of human pluripotent stem cells for neurodegeneration research.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e053911	1.2	

245	ADRC GWAS supplement: An NCRAD initiative.. <i>Alzheimer's and Dementia</i> , 2021 , 17 Suppl 3, e056176	1.2	
244	Web-based requisitioning, tracking and laboratory result reporting system for clinical trials using a central laboratory. <i>Alzheimer's and Dementia</i> , 2020 , 16, e038627	1.2	
243	Plasma biomarkers of inflammation reflect disease state in autosomal dominant forms of familial frontotemporal dementia. <i>Alzheimer's and Dementia</i> , 2020 , 16, e041533	1.2	
242	Amyloid and tau PET in sporadic early-onset Alzheimer's disease: Preliminary results from LEADS. <i>Alzheimer's and Dementia</i> , 2020 , 16, e041613	1.2	1
241	Associations of targeted genetic variants with Alzheimer's disease in African Americans and Nigerians. <i>Alzheimer's and Dementia</i> , 2020 , 16, e042323	1.2	
240	Studying the natural history of frontotemporal lobar degeneration (FTLD): The ARTFL LEFFTDS longitudinal FTLD (ALLFTD) protocol. <i>Alzheimer's and Dementia</i> , 2020 , 16, e045482	1.2	
239	Increased white matter MRI T1 hypointensity volume in young-onset Alzheimer's disease patients is not accounted for by age or cardiovascular risk factors. <i>Alzheimer's and Dementia</i> , 2020 , 16, e045577	1.2	
238	Plasma neurofilament light chain levels reflect caregiver burden and social cognition measures in familial frontotemporal lobar degeneration (FTLD). <i>Alzheimer's and Dementia</i> , 2020 , 16, e046247	1.2	
237	Neurodegeneration in the Longitudinal Evaluation of Early Onset Alzheimer's Disease Study (LEADS) sample: Results from the MRI core. <i>Alzheimer's and Dementia</i> , 2020 , 16, e046338	1.2	
236	The Alzheimer's disease sequencing project follow up study (ADSP-FUS): Increasing ethnic diversity in Alzheimer's genetics research with addition of potential new cohorts. <i>Alzheimer's and Dementia</i> , 2020 , 16, e046400	1.2	0
235	Sex-associated differences in pathology burden in early-onset Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2020 , 16, e046532	1.2	1
234	Evolution of Alzheimer's Disease Cerebrospinal Fluid Biomarkers in Early Parkinson's Disease. <i>Annals of Neurology</i> , 2020 , 88, 574-587	9.4	16
233	Harnessing peripheral DNA methylation differences in the Alzheimer's Disease Neuroimaging Initiative (ADNI) to reveal novel biomarkers of disease. <i>Clinical Epigenetics</i> , 2020 , 12, 84	7.7	15
232	Clinical and Dopamine Transporter Imaging Characteristics of Leucine Rich Repeat Kinase 2 (LRRK2) and Glucosylceramidase Beta (GBA) Parkinson's Disease Participants in the Parkinson's Progression Markers Initiative: A Cross-Sectional Study. <i>Movement Disorders</i> , 2020 , 35, 833-844	7	18
231	Leveraging genome-wide data to investigate differences between opioid use vs. opioid dependence in 41,176 individuals from the Psychiatric Genomics Consortium. <i>Molecular Psychiatry</i> , 2020 , 25, 1673-1687	15.1	30
230	Association of Blood Biomarkers With Acute Sport-Related Concussion in Collegiate Athletes: Findings From the NCAA and Department of Defense CARE Consortium. <i>JAMA Network Open</i> , 2020 , 3, e1919771	10.4	53
229	Exceptionally low likelihood of Alzheimer's dementia in APOE2 homozygotes from a 5,000-person neuropathological study. <i>Nature Communications</i> , 2020 , 11, 667	17.4	113
228	Clinical and dopamine transporter imaging characteristics of non-manifest LRRK2 and GBA mutation carriers in the Parkinson's Progression Markers Initiative (PPMI): a cross-sectional study. <i>Lancet Neurology</i> , 2020 , 19, 71-80	24.1	37

227	Exome-chip association analysis of intracranial aneurysms. <i>Neurology</i> , 2020 , 94, e481-e488	6.5	3
226	A large-scale genome-wide association study meta-analysis of cannabis use disorder. <i>Lancet Psychiatry</i> , 2020 , 7, 1032-1045	23.3	43
225	Genome-wide transcriptome analysis identifies novel dysregulated genes implicated in Alzheimer's pathology. <i>Alzheimer's and Dementia</i> , 2020 , 16, 1213-1223	1.2	8
224	A novel SNCA E83Q mutation in a case of dementia with Lewy bodies and atypical frontotemporal lobar degeneration. <i>Neuropathology</i> , 2020 , 40, 620-626	2	6
223	Longitudinal Measurements of Glucocerebrosidase activity in Parkinson's patients. <i>Annals of Clinical and Translational Neurology</i> , 2020 , 7, 1816-1830	5.3	10
222	Validation of Serum Neurofilament Light Chain as a Biomarker of Parkinson's Disease Progression. <i>Movement Disorders</i> , 2020 , 35, 1999-2008	7	32
221	Meta-analysis of up to 622,409 individuals identifies 40 novel smoking behaviour associated genetic loci. <i>Molecular Psychiatry</i> , 2020 , 25, 2392-2409	15.1	45
220	Genome-wide association studies of the self-rating of effects of ethanol (SRE). <i>Addiction Biology</i> , 2020 , 25, e12800	4.6	6
219	Fibroblast Growth Factor 23 Genotype and Cardiovascular Disease in Patients Undergoing Hemodialysis. <i>American Journal of Nephrology</i> , 2019 , 49, 125-132	4.6	5
218	Genome-wide association studies of alcohol dependence, DSM-IV criterion count and individual criteria. <i>Genes, Brain and Behavior</i> , 2019 , 18, e12579	3.6	22
217	Genome-wide association study identifies loci associated with liability to alcohol and drug dependence that is associated with variability in reward-related ventral striatum activity in African- and European-Americans. <i>Genes, Brain and Behavior</i> , 2019 , 18, e12580	3.6	8
216	Genome-wide association study identifies 30 loci associated with bipolar disorder. <i>Nature Genetics</i> , 2019 , 51, 793-803	36.3	662
215	Analysis of whole genome-transcriptomic organization in brain to identify genes associated with alcoholism. <i>Translational Psychiatry</i> , 2019 , 9, 89	8.6	33
214	Telomere Shortening in the Alzheimer's Disease Neuroimaging Initiative Cohort. <i>Journal of Alzheimer's Disease</i> , 2019 , 71, 33-43	4.3	9
213	Persistent Changes in Stress-Regulatory Genes in Pregnant Women or Children Exposed Prenatally to Alcohol. <i>Alcoholism: Clinical and Experimental Research</i> , 2019 , 43, 1887-1897	3.7	13
212	Cancer outcomes among Parkinson's disease patients with leucine rich repeat kinase 2 mutations, idiopathic Parkinson's disease patients, and nonaffected controls. <i>Movement Disorders</i> , 2019 , 34, 1392-1398	7	15
211	Psychosocial moderation of polygenic risk for cannabis involvement: the role of trauma exposure and frequency of religious service attendance. <i>Translational Psychiatry</i> , 2019 , 9, 269	8.6	3
210	Association of Polygenic Liability for Alcohol Dependence and EEG Connectivity in Adolescence and Young Adulthood. <i>Brain Sciences</i> , 2019 , 9,	3.4	7

209	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates Aβ tau, immunity and lipid processing. <i>Nature Genetics</i> , 2019 , 51, 414-430	36.3	917
208	Genetic architecture of subcortical brain structures in 38,851 individuals. <i>Nature Genetics</i> , 2019 , 51, 1624-1636	36.3	81
207	Exome Chip Meta-analysis Fine Maps Causal Variants and Elucidates the Genetic Architecture of Rare Coding Variants in Smoking and Alcohol Use. <i>Biological Psychiatry</i> , 2019 , 85, 946-955	7.9	35
206	Efficient region-based test strategy uncovers genetic risk factors for functional outcome in bipolar disorder. <i>European Neuropsychopharmacology</i> , 2019 , 29, 156-170	1.2	5
205	Angiotensin-related genetic determinants of cardiovascular disease in patients undergoing hemodialysis. <i>Nephrology Dialysis Transplantation</i> , 2019 , 34, 1924-1931	4.3	3
204	The Promise and Pitfalls of Facebook Advertising: a Genetic Counselor's Perspective. <i>Journal of Genetic Counseling</i> , 2018 , 27, 326-328	2.5	4
203	Functional variants in the gene confer shared effects on risk for Crohn's disease and Parkinson's disease. <i>Science Translational Medicine</i> , 2018 , 10,	17.5	165
202	Genome-wide association analysis identifies new candidate risk loci for familial intracranial aneurysm in the French-Canadian population. <i>Scientific Reports</i> , 2018 , 8, 4356	4.9	10
201	Genetic risk for schizophrenia and psychosis in Alzheimer disease. <i>Molecular Psychiatry</i> , 2018 , 23, 963-972	25.1	35
200	Ondansetron blocks wild-type and p.F503L variant small-conductance Ca-activated K channels. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2018 , 315, H375-H388	5.2	15
199	Immunohistochemical Method and Histopathology Judging for the Systemic Synuclein Sampling Study (S4). <i>Journal of Neuropathology and Experimental Neurology</i> , 2018 , 77, 793-802	3.1	22
198	Finding useful biomarkers for Parkinson's disease. <i>Science Translational Medicine</i> , 2018 , 10,	17.5	69
197	L1 coupling to ankyrin and the spectrin-actin cytoskeleton modulates ethanol inhibition of L1 adhesion and ethanol teratogenesis. <i>FASEB Journal</i> , 2018 , 32, 1364-1374	0.9	8
196	Meta-Analyses of Externalizing Disorders: Genetics or Prenatal Alcohol Exposure?. <i>Alcoholism: Clinical and Experimental Research</i> , 2018 , 42, 162-172	3.7	1
195	Genome-wide association study identifies a novel locus for cannabis dependence. <i>Molecular Psychiatry</i> , 2018 , 23, 1293-1302	15.1	28
194	P1-433: GRAY MATTER DEFICITS IN SYMPTOMATIC AND PRESYMPTOMATIC MAPT MUTATION CARRIERS 2018 , 14, P475-P476		
193	P1-597: AMYLOID NEUROIMAGING AND GENETICS INITIATIVE: IMPLEMENTING DNA COLLECTION USING NOVEL CONSENTING APPROACHES FOR AN IDEAS ADD-ON STUDY 2018 , 14, P566-P567		
192	P1-149: THE ALZHEIMER'S DISEASE SEQUENCING PROJECT (ADSP) DATA UPDATE 2018 2018 , 14, P333-P334		

191	O2-14-01: CHARACTERISTICS AND PROGRESS OF 320 SUBJECTS IN THE LONGITUDINAL EVALUATION OF FAMILIAL FRONTOTEMPORAL DEMENTIA SUBJECTS (LEFFTDS) PROTOCOL 2018 , 14, P656-P656		
190	Detecting significant genotype-phenotype association rules in bipolar disorder: market research meets complex genetics. <i>International Journal of Bipolar Disorders</i> , 2018 , 6, 24	5.4	4
189	Transancestral GWAS of alcohol dependence reveals common genetic underpinnings with psychiatric disorders. <i>Nature Neuroscience</i> , 2018 , 21, 1656-1669	25.5	257
188	P1-419: USING A BRAIN NETWORK APPROACH TO PREDICT GENETIC MUTATION IN INDIVIDUAL PATIENTS WITH FAMILIAL FRONTOTEMPORAL DEMENTIA 2018 , 14, P465-P466		
187	The Parkinson's progression markers initiative (PPMI) - establishing a PD biomarker cohort. <i>Annals of Clinical and Translational Neurology</i> , 2018 , 5, 1460-1477	5.3	142
186	Collagen COL22A1 maintains vascular stability and mutations in are potentially associated with intracranial aneurysms. <i>DMM Disease Models and Mechanisms</i> , 2018 , 11,	4.1	5
185	Feasibility and Safety of Multicenter Tissue and Biofluid Sampling for β Synuclein in Parkinson's Disease: The Systemic Synuclein Sampling Study (S4). <i>Journal of Parkinsons Disease</i> , 2018 , 8, 517-527	5.3	11
184	Meta-Analysis of Genetic Influences on Initial Alcohol Sensitivity. <i>Alcoholism: Clinical and Experimental Research</i> , 2018 , 42, 2349-2359	3.7	12
183	Combined Face-Brain Morphology and Associated Neurocognitive Correlates in Fetal Alcohol Spectrum Disorders. <i>Alcoholism: Clinical and Experimental Research</i> , 2018 , 42, 1769-1782	3.7	24
182	Novel genetic loci associated with hippocampal volume. <i>Nature Communications</i> , 2017 , 8, 13624	17.4	173
181	A genome wide association study of fast beta EEG in families of European ancestry. <i>International Journal of Psychophysiology</i> , 2017 , 115, 74-85	2.9	5
180	Comparison of Parent, Peer, Psychiatric, and Cannabis Use Influences Across Stages of Offspring Alcohol Involvement: Evidence from the COGA Prospective Study. <i>Alcoholism: Clinical and Experimental Research</i> , 2017 , 41, 359-368	3.7	37
179	An endophenotype approach to the genetics of alcohol dependence: a genome wide association study of fast beta EEG in families of African ancestry. <i>Molecular Psychiatry</i> , 2017 , 22, 1767-1775	15.1	15
178	Inflammatory profile discriminates clinical subtypes in LRRK2-associated Parkinson's disease. <i>European Journal of Neurology</i> , 2017 , 24, 427-e6	6	30
177	Neurology Individualized Medicine: When to Use Next-Generation Sequencing Panels. <i>Mayo Clinic Proceedings</i> , 2017 , 92, 292-305	6.4	36
176	Transethnic genome-wide scan identifies novel Alzheimer's disease loci. <i>Alzheimer's and Dementia</i> , 2017 , 13, 727-738	1.2	106
175	The Systemic Synuclein Sampling Study: toward a biomarker for Parkinson's disease. <i>Biomarkers in Medicine</i> , 2017 , 11, 359-368	2.3	33
174	Calcium-Sensing Receptor Genotype and Response to Cinacalcet in Patients Undergoing Hemodialysis. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2017 , 12, 1128-1138	6.9	12

173	Exome Sequencing Identifies Candidate Genetic Modifiers of Syndromic and Familial Thoracic Aortic Aneurysm Severity. <i>Journal of Cardiovascular Translational Research</i> , 2017 , 10, 423-432	3.3	17
172	Genome-Wide Association Study for Anthracycline-Induced Congestive Heart Failure. <i>Clinical Cancer Research</i> , 2017 , 23, 43-51	12.9	44
171	Targeted neurogenesis pathway-based gene analysis identifies ADORA2A associated with hippocampal volume in mild cognitive impairment and Alzheimer's disease. <i>Neurobiology of Aging</i> , 2017 , 60, 92-103	5.6	42
170	Alzheimer's Disease Sequencing Project discovery and replication criteria for cases and controls: Data from a community-based prospective cohort study with autopsy follow-up. <i>Alzheimer's and Dementia</i> , 2017 , 13, 1410-1413	1.2	6
169	Impact of Genetic Ancestry on Outcomes in ECOG-ACRIN-E5103. <i>JCO Precision Oncology</i> , 2017 , 2017,	3.6	12
168	A GABRA2 polymorphism improves a model for prediction of drinking initiation. <i>Alcohol</i> , 2017 , 63, 1-8	2.7	4
167	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. <i>Nature Genetics</i> , 2017 , 49, 1373-1384	36.3	508
166	Polygenic Scores for Major Depressive Disorder and Risk of Alcohol Dependence. <i>JAMA Psychiatry</i> , 2017 , 74, 1153-1160	14.5	56
165	[P3102]: GENETIC RISK FOR SCHIZOPHRENIA AND PSYCHOSIS IN ALZHEIMER DISEASE 2017 , 13, P973-P974		
164	Penetrance estimate of LRRK2 p.G2019S mutation in individuals of non-Ashkenazi Jewish ancestry. <i>Movement Disorders</i> , 2017 , 32, 1432-1438	7	74
163	Facial Curvature Detects and Explicates Ethnic Differences in Effects of Prenatal Alcohol Exposure. <i>Alcoholism: Clinical and Experimental Research</i> , 2017 , 41, 1471-1483	3.7	15
162	Parkinson's disease biomarkers: perspective from the NINDS Parkinson's Disease Biomarkers Program. <i>Biomarkers in Medicine</i> , 2017 , 11, 451-473	2.3	33
161	Association analysis of rare variants near the APOE region with CSF and neuroimaging biomarkers of Alzheimer's disease. <i>BMC Medical Genomics</i> , 2017 , 10, 29	3.7	17
160	Two novel loci, COBL and SLC10A2, for Alzheimer's disease in African Americans. <i>Alzheimer's and Dementia</i> , 2017 , 13, 119-129	1.2	48
159	[P3098]: ALZHEIMER'S DISEASE SEQUENCING PROJECT DISCOVERY AND REPLICATION CRITERIA FOR CASES AND CONTROLS: DATA FROM A COMMUNITY-BASED PROSPECTIVE COHORT STUDY WITH AUTOPSY FOLLOW-UP 2017 , 13, P971-P972		
158	[P1142]: DNA METHYLATION DYNAMICS IN ALZHEIMER'S DISEASE DIAGNOSIS AND PROGRESSION 2017 , 13, P297-P297		3
157	[P1054]: CHARACTERISTICS AND PROGRESS ON THE INITIAL 209 SUBJECTS IN THE LONGITUDINAL EVALUATION OF FAMILIAL FRONTOTEMPORAL DEMENTIA SUBJECTS (LEFFTDS) PROTOCOL 2017 , 13, P345-P345		
156	[P2B03]: ADVANCING RESEARCH AND TREATMENT IN FRONTOTEMPORAL LOBAR DEGENERATION (ARTFL) NORTH AMERICAN RARE DISEASE CLINICAL RESEARCH CONSORTIUM: PROGRESS AND CHARACTERIZATION OF INITIAL PARTICIPANTS 2017 , 13, P733-P734		

155	[P3090]: THE ALZHEIMER'S DISEASE SEQUENCING PROJECT (ADSP) DATA UPDATE 2017 2017 , 13, P968-P968		
154	Association of the OPRM1 Variant rs1799971 (A118G) with Non-Specific Liability to Substance Dependence in a Collaborative de novo Meta-Analysis of European-Ancestry Cohorts. <i>Behavior Genetics</i> , 2016 , 46, 151-69	3.2	77
153	Global and local ancestry in African-Americans: Implications for Alzheimer's disease risk. <i>Alzheimer's and Dementia</i> , 2016 , 12, 233-43	1.2	27
152	The Role of Cardiovascular Risk Factors and Stroke in Familial Alzheimer Disease. <i>JAMA Neurology</i> , 2016 , 73, 1231-1237	17.2	35
151	Shared Genetic Risk Factors of Intracranial, Abdominal, and Thoracic Aneurysms. <i>Journal of the American Heart Association</i> , 2016 , 5,	6	34
150	Integration of bioinformatics and imaging informatics for identifying rare PSEN1 variants in Alzheimer's disease. <i>BMC Medical Genomics</i> , 2016 , 9 Suppl 1, 30	3.7	16
149	Identification of TMEM230 mutations in familial Parkinson's disease. <i>Nature Genetics</i> , 2016 , 48, 733-9	36.3	122
148	Genome-wide association study of 40,000 individuals identifies two novel loci associated with bipolar disorder. <i>Human Molecular Genetics</i> , 2016 , 25, 3383-3394	5.6	125
147	Genome-wide association study of serum iron phenotypes in premenopausal women of European descent. <i>Blood Cells, Molecules, and Diseases</i> , 2016 , 57, 50-3	2.1	2
146	A MULTIVARIATE FINITE MIXTURE LATENT TRAJECTORY MODEL WITH APPLICATION TO DEMENTIA STUDIES. <i>Journal of Applied Statistics</i> , 2016 , 43, 2503-2523	1	11
145	Clinical-Genetic Associations in the Prospective Huntington at Risk Observational Study (PHAROS): Implications for Clinical Trials. <i>JAMA Neurology</i> , 2016 , 73, 102-10	17.2	29
144	Whole-Exome Sequencing in Familial Parkinson Disease. <i>JAMA Neurology</i> , 2016 , 73, 68-75	17.2	52
143	Steroid Pathway Genes and Neonatal Respiratory Distress After Betamethasone Use in Anticipated Preterm Birth. <i>Reproductive Sciences</i> , 2016 , 23, 680-6	3	8
142	Genetic Influences on Plasma Homocysteine Levels in African Americans and Yoruba Nigerians. <i>Journal of Alzheimer's Disease</i> , 2016 , 49, 991-1003	4.3	10
141	Genome wide association study for anthracycline-induced congestive heart failure.. <i>Journal of Clinical Oncology</i> , 2016 , 34, 1017-1017	2.2	1
140	Charcot-Marie-Tooth gene, SBF2, associated with taxane-induced peripheral neuropathy in African Americans. <i>Oncotarget</i> , 2016 , 7, 82244-82253	3.3	26
139	Association of Charcot-Marie-Tooth gene, SBF2, with taxane-induced peripheral neuropathy in African Americans.. <i>Journal of Clinical Oncology</i> , 2016 , 34, 1026-1026	2.2	
138	Inflammatory profile in LRRK2-associated prodromal and clinical PD. <i>Journal of Neuroinflammation</i> , 2016 , 13, 122	10.1	33

137	Genome-wide polygenic scores for age at onset of alcohol dependence and association with alcohol-related measures. <i>Translational Psychiatry</i> , 2016 , 6, e761	8.6	14
136	P2-097: The Alzheimer's Disease Sequencing Project (ADSP): Data Production, Management, and Availability 2016 , 12, P648-P648		
135	IC-P-072: Gene Expression Of ABCA7 Dysregulated in Peripheral Blood is Associated With Decreased Metabolic Activity in Hippocampus 2016 , 12, P56-P58		
134	IC-P-074: Genome-Wide Meta-Analysis of Transcriptome Profiling Identifies Novel Dysregulated Genes Implicated in Alzheimer's Disease 2016 , 12, P58-P60		
133	IC-P-075: The Growth and Impact of ADNI Genetics Publications as Measured by Science Mapping 2016 , 12, P60-P61		
132	P1-348: Neuropathology of Familial Alzheimer's Disease Associated with a Presenilin 1 A396T Mutation Reveals The Coexistence of A β TAU, and A-Synuclein Proteinopathies 2016 , 12, P562-P563		
131	P2-258: The Growth and Impact of ADNI Genetics Publications as Measured by Science Mapping 2016 , 12, P725-P726		
130	P3-087: Gene Expression of ABCA7 Dysregulated in Peripheral Blood is Associated With Decreased Metabolic Activity in Hippocampus 2016 , 12, P851-P853		
129	O2-06-02: Genome-Wide Meta-Analysis of Transcriptome Profiling Identifies Novel Dysregulated Genes Implicated in Alzheimer's Disease 2016 , 12, P238-P239		
128	NIPT and Informed Consent: an Assessment of Patient Understanding of a Negative NIPT Result. <i>Journal of Genetic Counseling</i> , 2016 , 25, 1127-37	2.5	27
127	Genome-wide linkage analyses of non-Hispanic white families identify novel loci for familial late-onset Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2016 , 12, 2-10	1.2	18
126	DNM3 and genetic modifiers of age of onset in LRRK2 Gly2019Ser parkinsonism: a genome-wide linkage and association study. <i>Lancet Neurology</i> , 2016 , 15, 1248-1256	24.1	50
125	Novel genetic loci underlying human intracranial volume identified through genome-wide association. <i>Nature Neuroscience</i> , 2016 , 19, 1569-1582	25.5	147
124	Knowledge gaps and research recommendations for essential tremor. <i>Parkinsonism and Related Disorders</i> , 2016 , 33, 27-35	3.6	33
123	Common genetic variants influence human subcortical brain structures. <i>Nature</i> , 2015 , 520, 224-9	50.4	601
122	A multi-ancestral genome-wide exome array study of Alzheimer disease, frontotemporal dementia, and progressive supranuclear palsy. <i>JAMA Neurology</i> , 2015 , 72, 414-22	17.2	33
121	Genome-Wide Association Studies for Taxane-Induced Peripheral Neuropathy in ECOG-5103 and ECOG-1199. <i>Clinical Cancer Research</i> , 2015 , 21, 5082-5091	12.9	79
120	Genetic studies of quantitative MCI and AD phenotypes in ADNI: Progress, opportunities, and plans. <i>Alzheimer's and Dementia</i> , 2015 , 11, 792-814	1.2	167

119	Genome-wide association data suggest ABCB1 and immune-related gene sets may be involved in adult antisocial behavior. <i>Translational Psychiatry</i> , 2015 , 5, e558	8.6	32
118	A description of the methods of the Nulliparous Pregnancy Outcomes Study: monitoring mothers-to-be (nuMoM2b). <i>American Journal of Obstetrics and Gynecology</i> , 2015 , 212, 539.e1-539.e24	6.4	79
117	APOE effect on Alzheimer's disease biomarkers in older adults with significant memory concern. <i>Alzheimer's and Dementia</i> , 2015 , 11, 1417-1429	1.2	116
116	TREM2 is associated with increased risk for Alzheimer's disease in African Americans. <i>Molecular Neurodegeneration</i> , 2015 , 10, 19	19	108
115	Polygenic risk for externalizing disorders: Gene-by-development and gene-by-environment effects in adolescents and young adults. <i>Clinical Psychological Science</i> , 2015 , 3, 189-201	6	70
114	Characteristics of Bipolar I patients grouped by externalizing disorders. <i>Journal of Affective Disorders</i> , 2015 , 178, 206-14	6.6	7
113	Adaptation of Subjective Responses to Alcohol is Affected by an Interaction of GABRA2 Genotype and Recent Drinking. <i>Alcoholism: Clinical and Experimental Research</i> , 2015 , 39, 1148-57	3.7	16
112	Fine mapping of bone structure and strength QTLs in heterogeneous stock rat. <i>Bone</i> , 2015 , 81, 417-426	4.7	9
111	GWAS of longitudinal amyloid accumulation on 18F-florbetapir PET in Alzheimer's disease implicates microglial activation gene IL1RAP. <i>Brain</i> , 2015 , 138, 3076-88	11.2	88
110	Convergent genetic and expression data implicate immunity in Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2015 , 11, 658-71	1.2	146
109	P3-011: Genome-wide association of plasma homocysteine in the indianapolis-ibadan dementia study cohort 2015 , 11, P623-P624		
108	IC-P-042: Influence of rare reelin variants on quantitative PET imaging and CSF phenotypes in late-onset Alzheimer's disease 2015 , 11, P36-P36		
107	P1-201: Genetic findings using ADNI multimodal quantitative phenotypes: A 2014 update 2015 , 11, P426-P426		
106	P3-014: Influence of rare RELN variants on quantitative PET imaging and CSF phenotypes in late-onset Alzheimer's disease 2015 , 11, P624-P625		
105	Assessment of first and second degree relatives of individuals with bipolar disorder shows increased genetic risk scores in both affected relatives and young At-Risk Individuals. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2015 , 168, 617-29	3.5	34
104	Novel recruitment strategy to enrich for LRRK2 mutation carriers. <i>Molecular Genetics & Genomic Medicine</i> , 2015 , 3, 404-12	2.3	14
103	Brief report: genetics of alcoholic cirrhosis-GenomALC multinational study. <i>Alcoholism: Clinical and Experimental Research</i> , 2015 , 39, 836-42	3.7	22
102	Genes Associated With Alcohol Outcomes Show Enrichment of Effects With Broad Externalizing and Impulsivity Phenotypes in an Independent Sample. <i>Journal of Studies on Alcohol and Drugs</i> , 2015 , 76, 38-46	1.9	9

101	P4-197: Gene expression profiling identifies altered networks in late-onset Alzheimer's disease: Immune response and mitochondrial process 2015 , 11, P855-P856		
100	Genetic variants associated with susceptibility to psychosis in late-onset Alzheimer's disease families. <i>Neurobiology of Aging</i> , 2015 , 36, 3116.e9-3116.e16	5.6	9
99	O4-05-01: Gwas of longitudinal amyloid PET identifies IL1RAP as a new potential Alzheimer's disease target 2015 , 11, P277-P278		
98	Comprehensive gene- and pathway-based analysis of depressive symptoms in older adults. <i>Journal of Alzheimer's Disease</i> , 2015 , 45, 1197-206	4.3	27
97	Lessons learned from whole exome sequencing in multiplex families affected by a complex genetic disorder, intracranial aneurysm. <i>PLoS ONE</i> , 2015 , 10, e0121104	3.7	27
96	Rarity of the Alzheimer disease-protective APP A673T variant in the United States. <i>JAMA Neurology</i> , 2015 , 72, 209-16	17.2	31
95	Guidelines for the standardization of preanalytic variables for blood-based biomarker studies in Alzheimer's disease research. <i>Alzheimer's and Dementia</i> , 2015 , 11, 549-60	1.2	159
94	AluY-mediated germline deletion, duplication and somatic stem cell reversion in UBE2T defines a new subtype of Fanconi anemia. <i>Human Molecular Genetics</i> , 2015 , 24, 5093-108	5.6	56
93	Linkage analyses in Caribbean Hispanic families identify novel loci associated with familial late-onset Alzheimer's disease. <i>Alzheimer's and Dementia</i> , 2015 , 11, 1397-1406	1.2	18
92	O3-13-04: Genome-wide rare variant analysis identifies candidate genes significantly associated with composite scores for memory 2015 , 11, P251-P252		0
91	Genes associated with alcohol outcomes show enrichment of effects with broad externalizing and impulsivity phenotypes in an independent sample. <i>Journal of Studies on Alcohol and Drugs</i> , 2015 , 76, 38-46	1.9	8
90	Identification of pathways for bipolar disorder: a meta-analysis. <i>JAMA Psychiatry</i> , 2014 , 71, 657-64	14.5	172
89	DSM-5 cannabis use disorder: a phenotypic and genomic perspective. <i>Drug and Alcohol Dependence</i> , 2014 , 134, 362-369	4.9	32
88	Ethanol treatment of lymphoblastoid cell lines from alcoholics and non-alcoholics causes many subtle changes in gene expression. <i>Alcohol</i> , 2014 , 48, 603-10	2.7	14
87	Large-scale meta-analysis of genome-wide association data identifies six new risk loci for Parkinson's disease. <i>Nature Genetics</i> , 2014 , 46, 989-93	36.3	1261
86	Parent-of-origin-specific allelic associations among 106 genomic loci for age at menarche. <i>Nature</i> , 2014 , 514, 92-97	50.4	401
85	Genome-wide survival analysis of age at onset of alcohol dependence in extended high-risk COGA families. <i>Drug and Alcohol Dependence</i> , 2014 , 142, 56-62	4.9	21
84	Genetics of alcoholism. <i>Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn</i> , 2014 , 125, 561-71	3	28

83	SIBLING family genes and bone mineral density: association and allele-specific expression in humans. <i>Bone</i> , 2014 , 64, 166-72	4.7	7
82	Association of plasma and cortical amyloid beta is modulated by APOE ϵ status. <i>Alzheimer's and Dementia</i> , 2014 , 10, e9-e18	1.2	35
81	Two rare AKAP9 variants are associated with Alzheimer's disease in African Americans. <i>Alzheimer's and Dementia</i> , 2014 , 10, 609-618.e11	1.2	83
80	Translational research on aging: clinical epidemiology as a bridge between the sciences. <i>Translational Research</i> , 2014 , 163, 439-45	11	6
79	IC-P-177: GENETIC FINDINGS USING ADNI MULTIMODAL QUANTITATIVE PHENOTYPES: A REVIEW OF PAPERS PUBLISHED IN 2013 2014 , 10, P99-P100		
78	IC-P-174: RARE VARIANT IN PLD3 IS ASSOCIATED WITH ALZHEIMER'S PATTERN OF NEURODEGENERATIVE CHANGES 2014 , 10, P97-P97		
77	P3-024: NEXT-GENERATION SEQUENCING OF THE BCHE LOCUS IDENTIFIES A FUNCTIONAL SNP ASSOCIATED WITH ALZHEIMER'S DISEASE BIOMARKERS AND AGE OF ONSET 2014 , 10, P636-P636		
76	P3-017: ASSOCIATION ANALYSIS OF RARE VARIANTS NEAR THE APOE REGION WITH CEREBROSPINAL FLUID (CSF) BIOMARKERS OF ALZHEIMER'S DISEASE 2014 , 10, P632-P633		
75	P1-141: GENETIC FINDINGS USING ADNI MULTIMODAL QUANTITATIVE PHENOTYPES: A REVIEW OF PAPERS PUBLISHED IN 2013 2014 , 10, P351-P351		
74	P3-019: RARE VARIANT IN PLD3 IS ASSOCIATED WITH ALZHEIMER'S PATTERN OF NEURODEGENERATIVE CHANGES 2014 , 10, P634-P634		
73	Genetic variant predicts bevacizumab-induced hypertension in ECOG-5103 and ECOG-2100. <i>British Journal of Cancer</i> , 2014 , 111, 1241-8	8.7	31
72	Phenotypic dissection of bone mineral density reveals skeletal site specificity and facilitates the identification of novel loci in the genetic regulation of bone mass attainment. <i>PLoS Genetics</i> , 2014 , 10, e1004423	6	107
71	P3-018: INFLUENCE OF RARE PSEN1 VARIANTS ON QUANTITATIVE STRUCTURAL IMAGING AND CSF PHENOTYPES IN LATE ONSET ALZHEIMER'S DISEASE 2014 , 10, P633-P633		
70	Michael J. Fox Foundation LRRK2 Consortium: geographical differences in returning genetic research data to study participants. <i>Genetics in Medicine</i> , 2014 , 16, 644-5	8.1	7
69	Age-specific incidence rates for dementia and Alzheimer disease in NIA-LOAD/NCRAD and EFIGA families: National Institute on Aging Genetics Initiative for Late-Onset Alzheimer Disease/National Cell Repository for Alzheimer Disease (NIA-LOAD/NCRAD) and Estudio Familiar de Influenza Genetica en Alzheimer (EFIGA). <i>JAMA Neurology</i> , 2014 , 71, 315-23	17.2	36
68	APOE ϵ and the risk for Alzheimer disease and cognitive decline in African Americans and Yoruba. <i>International Psychogeriatrics</i> , 2014 , 26, 977-85	3.4	54
67	Effects of multiple genetic loci on age at onset in late-onset Alzheimer disease: a genome-wide association study. <i>JAMA Neurology</i> , 2014 , 71, 1394-404	17.2	129
66	Gene-wide analysis detects two new susceptibility genes for Alzheimer's disease. <i>PLoS ONE</i> , 2014 , 9, e94661	3.7	90

65	Meta-analysis of 74,046 individuals identifies 11 new susceptibility loci for Alzheimer's disease. <i>Nature Genetics</i> , 2013 , 45, 1452-8	36.3	2714
64	Whole exome sequencing of intracranial aneurysm. <i>Stroke</i> , 2013 , 44, S26-8	6.7	9
63	Genetic influences on craving for alcohol. <i>Addictive Behaviors</i> , 2013 , 38, 1501-1508	4.2	38
62	Stress-response pathways are altered in the hippocampus of chronic alcoholics. <i>Alcohol</i> , 2013 , 47, 505-15	5.7	67
61	O40602: Genetic variation in PLXNA4 associated with susceptibility of Alzheimer's disease through tau phosphorylation 2013 , 9, P692-P692		1
60	A multicenter study of glucocerebrosidase mutations in dementia with Lewy bodies. <i>JAMA Neurology</i> , 2013 , 70, 727-35	17.2	285
59	Mutations disrupting PI3K signaling act as dominant enhancers of ethanol teratogenicity. <i>FASEB Journal</i> , 2013 , 27, 962.5	0.9	
58	Relation over time between facial measurements and cognitive outcomes in fetal alcohol-exposed children. <i>Alcoholism: Clinical and Experimental Research</i> , 2012 , 36, 1634-46	3.7	18
57	Genome-wide association study of intracranial aneurysms confirms role of Anril and SOX17 in disease risk. <i>Stroke</i> , 2012 , 43, 2846-52	6.7	90
56	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. <i>Nature Genetics</i> , 2011 , 43, 436-41	36.3	1367
55	Meta-analysis confirms CR1, CLU, and PICALM as alzheimer disease risk loci and reveals interactions with APOE genotypes. <i>Archives of Neurology</i> , 2010 , 67, 1473-84		330
54	GABRR1 and GABRR2, encoding the GABA-A receptor subunits rho1 and rho2, are associated with alcohol dependence. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2010 , 153B, 418-427	3.5	33
53	Alzheimer's Disease Neuroimaging Initiative biomarkers as quantitative phenotypes: Genetics core aims, progress, and plans. <i>Alzheimer's and Dementia</i> , 2010 , 6, 265-73	1.2	279
52	O3-03-01: Genome-wide association study of CSF biomarkers amyloid beta 1-42, tau and tau phosphorylated at threonine 181 in the ADNI cohort 2010 , 6, S129-S129		1
51	Genetic research: who is at risk for alcoholism. <i>Alcohol Research</i> , 2010 , 33, 64-75		14
50	Genomewide association study for susceptibility genes contributing to familial Parkinson disease. <i>Human Genetics</i> , 2009 , 124, 593-605	6.3	363
49	Genome screen in familial intracranial aneurysm. <i>BMC Medical Genetics</i> , 2009 , 10, 3	2.1	23
48	The tachykinin receptor 3 is associated with alcohol and cocaine dependence. <i>Alcoholism: Clinical and Experimental Research</i> , 2008 , 32, 1023-30	3.7	39

47	Genome screen to detect linkage to intracranial aneurysm susceptibility genes: the Familial Intracranial Aneurysm (FIA) study. <i>Stroke</i> , 2008 , 39, 1434-40	6.7	45
46	Association studies of ALOX5 and bone mineral density in healthy adults. <i>Osteoporosis International</i> , 2008 , 19, 637-43	5.3	7
45	Association of alcohol craving with alpha-synuclein (SNCA). <i>Alcoholism: Clinical and Experimental Research</i> , 2007 , 31, 537-45	3.7	48
44	Lack of association of alcohol dependence and habitual smoking with catechol-O-methyltransferase. <i>Alcoholism: Clinical and Experimental Research</i> , 2007 , 31, 1773-9	3.7	36
43	Meta-analysis of genome-wide scans provides evidence for sex- and site-specific regulation of bone mass. <i>Journal of Bone and Mineral Research</i> , 2007 , 22, 173-183	6.3	128
42	Chromosome 5 and Parkinson disease. <i>European Journal of Human Genetics</i> , 2006 , 14, 1106-10	5.3	4
41	Polymorphisms in the bone morphogenetic protein 2 (BMP2) gene do not affect bone mineral density in white men or women. <i>Osteoporosis International</i> , 2006 , 17, 587-92	5.3	12
40	A mutation in myotilin causes spheroid body myopathy. <i>Neurology</i> , 2005 , 65, 1936-40	6.5	70
39	Genetics of Parkinson disease. <i>Neurotherapeutics</i> , 2004 , 1, 235-242	6.4	
38	Confirmation of alcohol preference quantitative trait loci in the replicate high alcohol drinking and low alcohol drinking rat lines. <i>Psychiatric Genetics</i> , 2003 , 13, 155-61	2.9	17
37	Reliability of reported age at onset for Parkinson's disease. <i>Movement Disorders</i> , 2003 , 18, 275-279	7	27
36	Prenatal Alcohol Exposure: Advancing Knowledge Through International Collaborations. <i>Alcoholism: Clinical and Experimental Research</i> , 2003 , 27, 118-135	3.7	36
35	Prenatal Alcohol Exposure: Advancing Knowledge Through International Collaborations. <i>Alcoholism: Clinical and Experimental Research</i> , 2003 , 27, 118-135	3.7	
34	Saccadic Eye Movements Are Associated With a Family History of Alcoholism at Baseline and After Exposure to Alcohol. <i>Alcoholism: Clinical and Experimental Research</i> , 2002 , 26, 1568-1573	3.7	29
33	Mapping of QTL influencing saccharin consumption in the selectively bred alcohol-preferring and -nonpreferring rat lines. <i>Behavior Genetics</i> , 2002 , 32, 57-67	3.2	19
32	Parametric linkage analysis and disequilibrium methods to identify loci for complex disease. <i>Genetic Epidemiology</i> , 2001 , 21 Suppl 1, S528-33	2.6	1
31	Genome screen for quantitative trait loci underlying normal variation in femoral structure. <i>Journal of Bone and Mineral Research</i> , 2001 , 16, 985-91	6.3	99
30	Variability in skeletal mass, structure, and biomechanical properties among inbred strains of rats. <i>Journal of Bone and Mineral Research</i> , 2001 , 16, 1532-9	6.3	58

29	Mutations in a member of the ADAMTS gene family cause thrombotic thrombocytopenic purpura. <i>Nature</i> , 2001 , 413, 488-94	50.4	1394
28	Stoppage: An issue for segregation analysis 2001 , 20, 328		1
27	Suggestive evidence of a locus on chromosome 10p using the NIMH genetics initiative bipolar affective disorder pedigrees 2000 , 96, 18-23		58
26	A genome screen of maximum number of drinks as an alcoholism phenotype. <i>American Journal of Medical Genetics Part A</i> , 2000 , 96, 632-7		181
25	Alcoholism Susceptibility Loci: Confirmation Studies in a Replicate Sample and Further Mapping. <i>Alcoholism: Clinical and Experimental Research</i> , 2000 , 24, 933-945	3.7	205
24	Alcoholism Susceptibility Loci: Confirmation Studies in a Replicate Sample and Further Mapping 2000 , 24, 933		17
23	Suggestive evidence of a locus on chromosome 10p using the NIMH genetics initiative bipolar affective disorder pedigrees. <i>American Journal of Medical Genetics Part A</i> , 2000 , 96, 18-23		11
22	Alcoholism susceptibility loci: confirmation studies in a replicate sample and further mapping. <i>Alcoholism: Clinical and Experimental Research</i> , 2000 , 24, 933-45	3.7	103
21	Differences in duration of Huntington's disease based on age at onset. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 1999 , 66, 52-6	5.5	144
20	Genetics of alcoholism: a review of recent studies in human and animal models. <i>American Journal on Addictions</i> , 1999 , 8, 261-78	3.7	31
19	Chromosome workshop: Chromosomes 11, 14, and 15. <i>American Journal of Medical Genetics Part A</i> , 1999 , 88, 244-254		48
18	Linkage of type II and type III cystinuria to 19q13.1: Codominant inheritance of two cystinuric alleles at 19q13.1 produces an extreme stone-forming phenotype 1999 , 86, 134-139		16
17	Genome screen for platelet monoamine oxidase (MAO) activity. <i>American Journal of Medical Genetics Part A</i> , 1999 , 88, 517-21		12
16	Nonparametric linkage and family-based association studies of a simulated complex disorder. <i>Genetic Epidemiology</i> , 1999 , 17 Suppl 1, S627-32	2.6	
15	A Quantitative Trait Locus for Alcohol Consumption in Selectively Bred Rat Lines. <i>Alcoholism: Clinical and Experimental Research</i> , 1998 , 22, 884-887	3.7	171
14	Linkage of an Alcoholism-Related Severity Phenotype to Chromosome 16. <i>Alcoholism: Clinical and Experimental Research</i> , 1998 , 22, 2035-2042	3.7	59
13	Heterogeneity in hereditary pancreatitis 1998 , 77, 47-53		29
12	Genome-wide search for genes affecting the risk for alcohol dependence 1998 , 81, 207-215		496

11	Genomic screen for QTLs underlying alcohol consumption in the P and NP rat lines. <i>Mammalian Genome</i> , 1998 , 9, 949-55	3.2	87
10	Genetics of alcoholism. <i>Science</i> , 1998 , 282, 1269	33.3	6
9	A Quantitative Trait Locus for Alcohol Consumption in Selectively Bred Rat Lines 1998 , 22, 884		4
8	Genome-wide search for genes affecting the risk for alcohol dependence 1998 , 81, 207		44
7	Linkage of an alcoholism-related severity phenotype to chromosome 16. <i>Alcoholism: Clinical and Experimental Research</i> , 1998 , 22, 2035-42	3.7	20
6	Localization of the gene for familial primary pulmonary hypertension to chromosome 2q31-32. <i>Nature Genetics</i> , 1997 , 15, 277-80	36.3	218
5	Initial genomic scan of the NIMH genetics initiative bipolar pedigrees: Chromosomes 3, 5, 15, 16, 17, and 22 1997 , 74, 238-246		133
4	Spheroid body myopathy revisited. <i>Muscle and Nerve</i> , 1997 , 20, 1127-36	3.4	23
3	Cognitive scores in carriers of Huntington's disease gene compared to noncarriers. <i>Annals of Neurology</i> , 1995 , 37, 657-64	9.4	110
2	Linkage of the Indiana kindred of Gerstmann-Strüssler-Scheinker disease to the prion protein gene. <i>Nature Genetics</i> , 1992 , 1, 64-7	36.3	177
1	Trans-ancestral GWAS of alcohol dependence reveals common genetic underpinnings with psychiatric disorders		7