# Lindsay A Farrer

# 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.

30,800 80 169 379 h-index g-index citations papers 6.52 8.3 412 37,510 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
379	Blood and brain transcriptome analysis reveals APOE genotype-mediated and immune-related pathways involved in Alzheimer disease <i>Alzheimerm Research and Therapy</i> , <b>2022</b> , 14, 30	9	O
378	An association test of the spatial distribution of rare missense variants within protein structures identify Alzheimer's disease-related patterns <i>Genome Research</i> , <b>2022</b> ,	9.7	1
377	Genome-wide association study of brain arteriolosclerosis <i>Journal of Cerebral Blood Flow and Metabolism</i> , <b>2022</b> , 271678X211066299	7.3	O
376	New insights into the genetic etiology of Alzheimer's disease and related dementias <i>Nature Genetics</i> , <b>2022</b> ,	36.3	27
375	Alzheimer's disease associated AKAP9 I2558M mutation alters posttranslational modification and interactome of tau and cellular functions in CRISPR-edited human neuronal cells <i>Aging Cell</i> , <b>2022</b> , e136	5 <del>77</del>	O
374	Molecular Quantitative Trait Locus Mapping in Human Complex Diseases <i>Current Protocols</i> , <b>2022</b> , 2, e426		0
373	A missense variant in SHARPIN mediates Alzheimer's disease-specific brain damages. <i>Translational Psychiatry</i> , <b>2021</b> , 11, 590	8.6	2
372	Exploration of alcohol use disorder-associated brain miRNA-mRNA regulatory networks. <i>Translational Psychiatry</i> , <b>2021</b> , 11, 504	8.6	5
371	Analysis of telomere length variation and Shelterin complex subunit gene expression changes in ethanol-exposed human embryonic stem cells. <i>Journal of Psychiatric Research</i> , <b>2021</b> , 143, 543-549	5.2	1
370	Set-Based Rare Variant Expression Quantitative Trait Loci in Blood and Brain from Alzheimer Disease Study Participants. <i>Genes</i> , <b>2021</b> , 12,	4.2	3
369	Cell-type-specific expression quantitative trait loci associated with Alzheimer disease in blood and brain tissue. <i>Translational Psychiatry</i> , <b>2021</b> , 11, 250	8.6	4
368	Common variants in Alzheimer's disease and risk stratification by polygenic risk scores. <i>Nature Communications</i> , <b>2021</b> , 12, 3417	17.4	23
367	Genome-wide association study of stimulant dependence. <i>Translational Psychiatry</i> , <b>2021</b> , 11, 363	8.6	O
366	Causal Associations Between Modifiable Risk Factors and the Alzheimer's Phenome. <i>Annals of Neurology</i> , <b>2021</b> , 89, 54-65	9.4	21
365	Novel Alzheimer Disease Risk Loci and Pathways in African American Individuals Using the African Genome Resources Panel: A Meta-analysis. <i>JAMA Neurology</i> , <b>2021</b> , 78, 102-113	17.2	32
364	Genome-wide association study of phenotypes measuring progression from first cocaine or opioid use to dependence reveals novel risk genes. <i>Exploration of Medicine</i> , <b>2021</b> , 2, 60-73	1.1	1
363	Integrative brain transcriptome analysis links complement component 4 and HSPA2 to the APOE Depote to the APOE Depote time and HSPA2 to the AP	15.1	1

362	Enhancing Discovery of Genetic Variants for Posttraumatic Stress Disorder Through Integration of Quantitative Phenotypes and Trauma Exposure Information. <i>Biological Psychiatry</i> , <b>2021</b> ,	7.9	3
361	Alzheimer's disease associated AKAP9 I2558M mutation alters posttranslational modification and interactome of tau and cellular functions in CRISPR-edited human neuronal cells <i>Alzheimermand Dementia</i> , <b>2021</b> , 17 Suppl 2, e058592	1.2	
360	APOE-stratified genome-wide association analysis identifies novel Alzheimer disease candidate risk loci for African Americans <i>Alzheimerm and Dementia</i> , <b>2021</b> , 17 Suppl 3, e056383	1.2	O
359	Multivariate analysis of blood and brain transcriptome in Alzheimer's reveals unique APOE 4-related immune pathways <i>Alzheimermand Dementia</i> , <b>2021</b> , 17 Suppl 3, e054237	1.2	
358	Domain specific cognitive functions predict neuropathological traits in the Framingham Heart Study <i>Alzheimermand Dementia</i> , <b>2021</b> , 17 Suppl 3, e054249	1.2	
357	Genome-wide association and multi-omics studies identify MGMT as a novel risk gene for Alzheimer disease among women <i>Alzheimermand Dementia</i> , <b>2021</b> , 17 Suppl 3, e054483	1.2	
356	Multiple viruses detected in human DNA are associated with Alzheimer disease risk <i>Alzheimern</i> a and Dementia, <b>2021</b> , 17 Suppl 3, e054585	1.2	
355	Sex differences in the genetic architecture underlying resilience in AD <i>Alzheimermand Dementia</i> , <b>2021</b> , 17 Suppl 3, e055010	1.2	
354	Sex-specific genetic predictors of memory performance <i>Alzheimerm and Dementia</i> , <b>2021</b> , 17 Suppl 3, e056083	1.2	
353	Admixture mapping identifies novel regions influencing Alzheimer disease in African Americans <i>Alzheimermand Dementia</i> , <b>2021</b> , 17 Suppl 3, e056443	1.2	
352	Novel mechanism underlying the APOE 2 protective effect for Alzheimer disease implicated by integrative genome and transcriptome analysis. <i>Alzheimerm and Dementia</i> , <b>2020</b> , 16, e040065	1.2	
351	Cell-type specific eQTLs (ct-eQTLs) associated with Alzheimer disease in blood and brain tissue. <i>Alzheimermand Dementia</i> , <b>2020</b> , 16, e044149	1.2	
350	Genome-wide meta-analysis of late-onset Alzheimer disease using rare variant imputation in 65,602 subjects identifies risk loci with roles in memory, neurodevelopment, and cardiometabolic traits: The international genomics of Alzheimer project (IGAP). Alzheimer and Dementia, 2020,	1.2	O
349	16, e044193 Differential effects of apolipoprotein E on the molecular and cellular phenotypes associated with Alzheimer disease in isogenic human iPSC-derived neurons. <i>Alzheimer and Dementia</i> , 2020, 16, e04457	<del>1</del> 92	
348	Mechanism for the protective effect of APOE I against Alzheimer disease is linked to tau and the classical complement pathway. <i>Alzheimermand Dementia</i> , <b>2020</b> , 16, e044881	1.2	
347	Alzheimer disease risk factor mutations in patients diagnosed with Creutzfelt-Jakob disease. <i>Alzheimermand Dementia</i> , <b>2020</b> , 16, e045035	1.2	
346	Assessing whole genome sequencing variation for Alzheimer disease in 4707 individuals from the Alzheimer Disease Sequencing Project (ADSP). <i>Alzheimer and Dementia</i> , <b>2020</b> , 16, e045548	1.2	
345	Mapping Alzheimer disease\( \text{Bssociated regions in the African American population. } \) Alzheimer\( \text{mand} \) Dementia, \( \text{2020}, 16, e046072 \)	1.2	

344	Early-mid adulthood measures of HDL, triglycerides and fasting glucose are associated with late-onset Alzheimer disease. <i>Alzheimermand Dementia</i> , <b>2020</b> , 16, e046125	1.2	0
343	Genome-wide interaction study of smoking in Alzheimer disease. <i>Alzheimer mand Dementia</i> , <b>2020</b> , 16, e046149	1.2	
342	Structural characterization of rare missense variants within known neurodegenerative disease proteins. <i>Alzheimermand Dementia</i> , <b>2020</b> , 16, e046405	1.2	
341	Defining Alzheimer disease subtypes using polygenic risk scores integrated with genomic and brain transcriptomic profiles. <i>Alzheimer</i> and <i>Dementia</i> , <b>2020</b> , 16, e046449	1.2	1
340	Genome wide association study of chronic traumatic encephalopathy. <i>Alzheimerm and Dementia</i> , <b>2020</b> , 16, e046505	1.2	
339	Expanding the genetic architecture of nicotine dependence and its shared genetics with multiple traits. <i>Nature Communications</i> , <b>2020</b> , 11, 5562	17.4	25
338	Genome-wide association study of rate of cognitive decline in Alzheimer's disease patients identifies novel genes and pathways. <i>Alzheimermand Dementia</i> , <b>2020</b> , 16, 1134-1145	1.2	9
337	Association of OPRM1 Functional Coding Variant With Opioid Use Disorder: A Genome-Wide Association Study. <i>JAMA Psychiatry</i> , <b>2020</b> , 77, 1072-1080	14.5	46
336	Prefrontal cortex eQTLs/mQTLs enriched in genetic variants associated with alcohol use disorder and other diseases. <i>Epigenomics</i> , <b>2020</b> , 12, 789-800	4.4	5
335	Sex-dependent autosomal effects on clinical progression of Alzheimer's disease. <i>Brain</i> , <b>2020</b> , 143, 2272	2-2280	23
335 334	Sex-dependent autosomal effects on clinical progression of Alzheimer's disease. <i>Brain</i> , <b>2020</b> , 143, 2272  An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> , 20, 672-680	2- <b>228</b> 0 3.5	23
	An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> ,		
334	An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> , 20, 672-680	3.5	6
334	An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> , 20, 672-680  Genomic influences on self-reported childhood maltreatment. <i>Translational Psychiatry</i> , <b>2020</b> , 10, 38  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> ,	3·5 8.6	6
334 333 332	An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> , 20, 672-680  Genomic influences on self-reported childhood maltreatment. <i>Translational Psychiatry</i> , <b>2020</b> , 10, 38  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> , <b>2020</b> , 25, 1673-1687	3.5 8.6 15.1	6 20 30
334 333 332 331	An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> , 20, 672-680  Genomic influences on self-reported childhood maltreatment. <i>Translational Psychiatry</i> , <b>2020</b> , 10, 38  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> , <b>2020</b> , 25, 1673-1687  Genome-Wide Association Study of Opioid Cessation. <i>Journal of Clinical Medicine</i> , <b>2020</b> , 9,  Exceptionally low likelihood of Alzheimer's dementia in APOE2 homozygotes from a 5,000-person	3.5 8.6 15.1 5.1	6 20 30 10
334 333 332 331 330	An analysis of the effect of mu-opioid receptor gene (OPRM1) promoter region DNA methylation on the response of naltrexone treatment of alcohol dependence. <i>Pharmacogenomics Journal</i> , <b>2020</b> , 20, 672-680  Genomic influences on self-reported childhood maltreatment. <i>Translational Psychiatry</i> , <b>2020</b> , 10, 38  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> , <b>2020</b> , 25, 1673-1687  Genome-Wide Association Study of Opioid Cessation. <i>Journal of Clinical Medicine</i> , <b>2020</b> , 9,  Exceptionally low likelihood of Alzheimer's dementia in APOE2 homozygotes from a 5,000-person neuropathological study. <i>Nature Communications</i> , <b>2020</b> , 11, 667	3.5 8.6 15.1 5.1	6 20 30 10

## (2018-2020)

326	Analysis of brain region-specific co-expression networks reveals clustering of established and novel genes associated with Alzheimer disease. <i>Alzheimern Research and Therapy</i> , <b>2020</b> , 12, 103	9	2
325	Genetic variants and functional pathways associated with resilience to Alzheimer's disease. <i>Brain</i> , <b>2020</b> , 143, 2561-2575	11.2	25
324	Whole exome sequencing study identifies novel rare and common Alzheimer's-Associated variants involved in immune response and transcriptional regulation. <i>Molecular Psychiatry</i> , <b>2020</b> , 25, 1859-1875	15.1	106
323	Comparison of methods for multivariate gene-based association tests for complex diseases using common variants. <i>European Journal of Human Genetics</i> , <b>2019</b> , 27, 811-823	5.3	8
322	The genetics and epigenetics of Neonatal Abstinence Syndrome. <i>Seminars in Fetal and Neonatal Medicine</i> , <b>2019</b> , 24, 105-110	3.7	25
321	Analysis of Whole-Exome Sequencing Data for Alzheimer Disease Stratified by APOE Genotype. <i>JAMA Neurology</i> , <b>2019</b> , 76, 1099-1108	17.2	18
320	Salivary microRNAs identified by small RNA sequencing and machine learning as potential biomarkers of alcohol dependence. <i>Epigenomics</i> , <b>2019</b> , 11, 739-749	4.4	7
319	CpG-related SNPs in the MS4A region have a dose-dependent effect on risk of late-onset Alzheimer disease. <i>Aging Cell</i> , <b>2019</b> , 18, e12964	9.9	6
318	The Utah Protocol for Postmortem Eye Phenotyping and Molecular Biochemical Analysis <b>2019</b> , 60, 1204	l-1212	11
317	Association of Rare Coding Mutations With Alzheimer Disease and Other Dementias Among Adults of European Ancestry. <i>JAMA Network Open</i> , <b>2019</b> , 2, e191350	10.4	37
316	Promoter Polymorphism-219T/G is an Effect Modifier of the Influence of 🛭 on Alzheimer's Disease Risk in a Multiracial Sample. <i>Journal of Clinical Medicine</i> , <b>2019</b> , 8,	5.1	20
315	A systems biology approach towards understanding and treating non-neovascular age-related macular degeneration. <i>Nature Communications</i> , <b>2019</b> , 10, 3347	17.4	104
314	International meta-analysis of PTSD genome-wide association studies identifies sex- and ancestry-specific genetic risk loci. <i>Nature Communications</i> , <b>2019</b> , 10, 4558	17.4	151
313	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates All tau, immunity and lipid processing. <i>Nature Genetics</i> , <b>2019</b> , 51, 414-430	36.3	917
312	A regulatory variant of CHRM3 is associated with cannabis-induced hallucinations in European Americans. <i>Translational Psychiatry</i> , <b>2019</b> , 9, 309	8.6	0
311	Dissecting the genetic relationship between cardiovascular risk factors and Alzheimer's disease. <i>Acta Neuropathologica</i> , <b>2019</b> , 137, 209-226	14.3	54
310	A rare missense variant of CASP7 is associated with familial late-onset Alzheimer's disease. <i>Alzheimermand Dementia</i> , <b>2019</b> , 15, 441-452	1.2	22
309	Genome-wide Association Study Identifies a Regulatory Variant of RGMA Associated With Opioid Dependence in European Americans. <i>Biological Psychiatry</i> , <b>2018</b> , 84, 762-770	7.9	41

308	Genetic Variation in Genes Underlying Diverse Dementias May Explain a Small Proportion of Cases in the Alzheimer's Disease Sequencing Project. <i>Dementia and Geriatric Cognitive Disorders</i> , <b>2018</b> , 45, 1-	17 <sup>2.6</sup>	16
307	FUS Phase Separation Is Modulated by a Molecular Chaperone and Methylation of Arginine Cation-Interactions. <i>Cell</i> , <b>2018</b> , 173, 720-734.e15	56.2	409
306	Genome-wide association study of Alzheimer's disease endophenotypes at prediagnosis stages. <i>Alzheimer nand Dementia</i> , <b>2018</b> , 14, 623-633	1.2	35
305	Tau Phosphorylation is Impacted by Rare AKAP9 Mutations Associated with Alzheimer Disease in African Americans. <i>Journal of NeuroImmune Pharmacology</i> , <b>2018</b> , 13, 254-264	6.9	13
304	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360,	33.3	666
303	Genome-wide pleiotropy analysis of neuropathological traits related to Alzheimer's disease. <i>Alzheimer Research and Therapy</i> , <b>2018</b> , 10, 22	9	13
302	Genome-wide association study of cognitive flexibility assessed by the Wisconsin Card Sorting Test. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2018</b> , 177, 511-519	3.5	0
301	Male-specific epistasis between WWC1 and TLN2 genes is associated with Alzheimer's disease. <i>Neurobiology of Aging</i> , <b>2018</b> , 72, 188.e3-188.e12	5.6	13
300	Targeted Sequencing of Alzheimer Disease Genes in African Americans Implicates Novel Risk Variants. <i>Frontiers in Neuroscience</i> , <b>2018</b> , 12, 592	5.1	16
299	Translational studies support a role for serotonin 2B receptor (HTR2B) gene in aggression-related cannabis response. <i>Molecular Psychiatry</i> , <b>2018</b> , 23, 2277-2286	15.1	14
298	Rare genetic variation implicated in non-Hispanic white families with Alzheimer disease. <i>Neurology: Genetics</i> , <b>2018</b> , 4, e286	3.8	15
297	Transancestral GWAS of alcohol dependence reveals common genetic underpinnings with psychiatric disorders. <i>Nature Neuroscience</i> , <b>2018</b> , 21, 1656-1669	25.5	257
296	Ancestral origin of ApoE Alzheimer disease risk in Puerto Rican and African American populations. <i>PLoS Genetics</i> , <b>2018</b> , 14, e1007791	6	56
295	Genome-wide association study identifies glutamate ionotropic receptor GRIA4 as a risk gene for comorbid nicotine dependence and major depression. <i>Translational Psychiatry</i> , <b>2018</b> , 8, 208	8.6	8
294	Risk Locus Identification Ties Alcohol Withdrawal Symptoms to SORCS2. <i>Alcoholism: Clinical and Experimental Research</i> , <b>2018</b> , 42, 2337-2348	3.7	8
293	Variation in TMEM106B in chronic traumatic encephalopathy. <i>Acta Neuropathologica Communications</i> , <b>2018</b> , 6, 115	7.3	21
292	One for all and all for One: Improving replication of genetic studies through network diffusion. <i>PLoS Genetics</i> , <b>2018</b> , 14, e1007306	6	18
291	Genome-wide association study of body mass index in subjects with alcohol dependence. <i>Addiction Biology</i> , <b>2017</b> , 22, 535-549	4.6	18

## (2017-2017)

290	Association of maternal and infant variants in PNOC and COMT genes with neonatal abstinence syndrome severity. <i>American Journal on Addictions</i> , <b>2017</b> , 26, 42-49	3.7	25
289	A Common Variant of IL-6R is Associated with Elevated IL-6 Pathway Activity in Alzheimer's Disease Brains. <i>Journal of Alzheimer</i> Disease, <b>2017</b> , 56, 1037-1054	4.3	29
288	Genome-wide association study identifies four novel loci associated with Alzheimer's endophenotypes and disease modifiers. <i>Acta Neuropathologica</i> , <b>2017</b> , 133, 839-856	14.3	107
287	Transethnic genome-wide scan identifies novel Alzheimer's disease loci. <i>Alzheimer</i> and <i>Dementia</i> , <b>2017</b> , 13, 727-738	1.2	106
286	A common haplotype lowers PU.1 expression in myeloid cells and delays onset of Alzheimer's disease. <i>Nature Neuroscience</i> , <b>2017</b> , 20, 1052-1061	25.5	228
285	Genetic Risk Variants Associated With Comorbid Alcohol Dependence and Major Depression. <i>JAMA Psychiatry</i> , <b>2017</b> , 74, 1234-1241	14.5	47
284	Genome-Wide Association Studies of a Broad Spectrum of Antisocial Behavior. <i>JAMA Psychiatry</i> , <b>2017</b> , 74, 1242-1250	14.5	124
283	A phased SNP-based classification of sickle cell anemia HBB haplotypes. <i>BMC Genomics</i> , <b>2017</b> , 18, 608	4.5	19
282	Caspase-8, association with Alzheimer's Disease and functional analysis of rare variants. <i>PLoS ONE</i> , <b>2017</b> , 12, e0185777	3.7	23
281	Genetic assessment of age-associated Alzheimer disease risk: Development and validation of a polygenic hazard score. <i>PLoS Medicine</i> , <b>2017</b> , 14, e1002258	11.6	209
280	A putative causal relationship between genetically determined female body shape and posttraumatic stress disorder. <i>Genome Medicine</i> , <b>2017</b> , 9, 99	14.4	23
279	identified in a genome-wide gene Lannabis dependence interaction analysis of risky sexual behaviours. <i>Journal of Psychiatry and Neuroscience</i> , <b>2017</b> , 42, 252-261	4.5	5
278	Oxytocin receptor gene polymorphisms, attachment, and PTSD: Results from the National Health and Resilience in Veterans Study. <i>Journal of Psychiatric Research</i> , <b>2017</b> , 94, 139-147	5.2	34
277	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. <i>Nature Genetics</i> , <b>2017</b> , 49, 1373-1384	36.3	508
276	Early-Onset Alzheimer Disease and Candidate Risk Genes Involved in Endolysosomal Transport. <i>JAMA Neurology</i> , <b>2017</b> , 74, 1113-1122	17.2	30
275	[P3월31]: DEEP LEARNING APPLICATION IN IDENTIFYING PROTEOMIC RISK MARKERS FOR ALZHEIMER's DISEASE <b>2017</b> , 13, P1133-P1133		2
274	[O1D3D1]: GENOME-WIDE RARE VARIANT IMPUTATION AND TISSUE-SPECIFIC TRANSCRIPTOMIC ANALYSIS IDENTIFY NOVEL RARE VARIANT CANDIDATE LOCI IN LATE-ONSET ALZHEIMER'S DISEASE: THE ALZHEIMER'S DISEASE GENETICS CONSORTIUM <b>2017</b> , 13, P189		3
273	[O20804]: NOVEL GENETIC VARIANTS ASSOCIATED WITH FAMILIAL LATE-ONSET ALZHEIMER DISEASE IN THE ALZHEIMER's DISEASE SEQUENCING PROJECT 2017, 13, P572		

272	Genetic factor common to schizophrenia and HIV infection is associated with risky sexual behavior: antagonistic vs. synergistic pleiotropic SNPs enriched for distinctly different biological functions. <i>Human Genetics</i> , <b>2017</b> , 136, 75-83	6.3	11
271	Genetics of age-related macular degeneration (AMD). Human Molecular Genetics, 2017, 26, R45-R50	5.6	53
270	Two novel loci, COBL and SLC10A2, for Alzheimer's disease in African Americans. <i>Alzheimer and Dementia</i> , <b>2017</b> , 13, 119-129	1.2	48
269	The Interplay Between Risky Sexual Behaviors and Alcohol Dependence: Genome-Wide Association and Neuroimaging Support for LHPP as a Risk Gene. <i>Neuropsychopharmacology</i> , <b>2017</b> , 42, 598-605	8.7	28
268	[P1🛮42]: GENOME-WIDE ASSOCIATION STUDY OF ALZHEIMER DISEASE ENDOPHENOTYPES AT PRECLINICAL AND MCI STAGES <b>2017</b> , 13, P337-P338		
267	[P3 <b>0</b> 92]: TAU PHOSPHORYLATION IS IMPACTED BY RARE AD-ASSOCIATED AKAP9 MUTATIONS SPECIFIC TO AFRICAN AMERICANS <b>2017</b> , 13, P969-P969		
266	[O20802]: SEX-SPECIFIC ANALYSIS OF THE ADSP CASE-CONTROL WHOLE-EXOME SEQUENCING DATASET <b>2017</b> , 13, P571		
265	Global and local ancestry in African-Americans: Implications for Alzheimer's disease risk. <i>Alzheimern</i> s and Dementia, <b>2016</b> , 12, 233-43	1.2	27
264	ABCA7 frameshift deletion associated with Alzheimer disease in African Americans. <i>Neurology: Genetics</i> , <b>2016</b> , 2, e79	3.8	43
263	Homozygosity for a haplotype in the HBG2-OR51B4 region is exclusive to Arab-Indian haplotype sickle cell anemia. <i>American Journal of Hematology</i> , <b>2016</b> , 91, E308-11	7.1	13
262	Discovery of gene-gene interactions across multiple independent data sets of late onset Alzheimer disease from the Alzheimer Disease Genetics Consortium. <i>Neurobiology of Aging</i> , <b>2016</b> , 38, 141-150	5.6	31
261	A large genome-wide association study of age-related macular degeneration highlights contributions of rare and common variants. <i>Nature Genetics</i> , <b>2016</b> , 48, 134-43	36.3	769
260	Genome-wide Association Study of Cannabis Dependence Severity, Novel Risk Variants, and Shared Genetic Risks. <i>JAMA Psychiatry</i> , <b>2016</b> , 73, 472-80	14.5	113
259	Genome-wide linkage analyses of non-Hispanic white families identify novel loci for familial late-onset Alzheimer's disease. <i>Alzheimermand Dementia</i> , <b>2016</b> , 12, 2-10	1.2	18
258	Assessment of the genetic variance of late-onset Alzheimer's disease. <i>Neurobiology of Aging</i> , <b>2016</b> , 41, 200.e13-200.e20	5.6	119
257	Segregation of a rare TTC3 variant in an extended family with late-onset Alzheimer disease. <i>Neurology: Genetics</i> , <b>2016</b> , 2, e41	3.8	31
256	Network-driven plasma proteomics expose molecular changes in the Alzheimer's brain. <i>Molecular Neurodegeneration</i> , <b>2016</b> , 11, 31	19	21
255	A candidate transacting modulator of fetal hemoglobin gene expression in the Arab-Indian haplotype of sickle cell anemia. <i>American Journal of Hematology</i> , <b>2016</b> , 91, 1118-1122	7.1	12

## (2015-2016)

254	Shared genetic contribution to Ischaemic Stroke and Alzheimer's Disease. <i>Annals of Neurology</i> , <b>2016</b> , 79, 739-747	9.4	42
253	Polygenic risk for alcohol dependence associates with alcohol consumption, cognitive function and social deprivation in a population-based cohort. <i>Addiction Biology</i> , <b>2016</b> , 21, 469-80	4.6	21
252	The ticking clock of Cayo Santiago macaques and its implications for understanding human circadian rhythm disorders. <i>American Journal of Primatology</i> , <b>2016</b> , 78, 117-26	2.5	2
251	Expanding the genomic roadmap of Alzheimer's disease. <i>Lancet Neurology, The</i> , <b>2015</b> , 14, 783-785	24.1	9
250	Eye color: A potential indicator of alcohol dependence risk in European Americans. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2015</b> , 168B, 347-53	3.5	6
249	Polygenic Overlap Between C-Reactive Protein, Plasma Lipids, and Alzheimer Disease. <i>Circulation</i> , <b>2015</b> , 131, 2061-2069	16.7	100
248	Association of Long Runs of Homozygosity With Alzheimer Disease Among African American Individuals. <i>JAMA Neurology</i> , <b>2015</b> , 72, 1313-23	17.2	27
247	Convergent genetic and expression data implicate immunity in Alzheimer's disease. <i>Alzheimermand Dementia</i> , <b>2015</b> , 11, 658-71	1.2	146
246	Genome-wide association study of copy number variations (CNVs) with opioid dependence. <i>Neuropsychopharmacology</i> , <b>2015</b> , 40, 1016-26	8.7	30
245	Nf1 regulates alcohol dependence-associated excessive drinking and gamma-aminobutyric acid release in the central amygdala in mice and is associated with alcohol dependence in humans. <i>Biological Psychiatry</i> , <b>2015</b> , 77, 870-879	7.9	11
244	Estimating the risk for conversion from mild cognitive impairment to Alzheimer's disease in an elderly Arab community. <i>Journal of Alzheimerm Disease</i> , <b>2015</b> , 45, 865-71	4.3	11
243	Genomewide Association Study for Maximum Number of Alcoholic Drinks in European Americans and African Americans. <i>Alcoholism: Clinical and Experimental Research</i> , <b>2015</b> , 39, 1137-47	3.7	46
242	Further analyses support the association between light eye color and alcohol dependence. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2015</b> , 168, 757-60	3.5	1
241	Reply: To PMID 25559091. <i>Annals of Neurology</i> , <b>2015</b> , 78, 836-7	9.4	
240	Ancestry of the Timorese: age-related macular degeneration associated genotype and allele sharing among human populations from throughout the world. <i>Frontiers in Genetics</i> , <b>2015</b> , 6, 238	4.5	4
239	Rarity of the Alzheimer disease-protective APP A673T variant in the United States. <i>JAMA Neurology</i> , <b>2015</b> , 72, 209-16	17.2	31
238	Linkage analyses in Caribbean Hispanic families identify novel loci associated with familial late-onset Alzheimer's disease. <i>Alzheimerm and Dementia</i> , <b>2015</b> , 11, 1397-1406	1.2	18
237	Variations in opioid receptor genes in neonatal abstinence syndrome. <i>Drug and Alcohol Dependence</i> , <b>2015</b> , 155, 253-9	4.9	44

236	Genome-wide association study of nicotine dependence in American populations: identification of novel risk loci in both African-Americans and European-Americans. <i>Biological Psychiatry</i> , <b>2015</b> , 77, 493-	5 <b>0</b> 3 <sup>9</sup>	56
235	Protective variant for hippocampal atrophy identified by whole exome sequencing. <i>Annals of Neurology</i> , <b>2015</b> , 77, 547-52	9.4	43
234	Genetic analysis of quantitative phenotypes in AD and MCI: imaging, cognition and biomarkers. <i>Brain Imaging and Behavior</i> , <b>2014</b> , 8, 183-207	4.1	111
233	Genome-wide association study of the rate of cognitive decline in Alzheimer's disease. <i>Alzheimeri</i> n and Dementia, <b>2014</b> , 10, 45-52	1.2	98
232	PLXNA4 is associated with Alzheimer disease and modulates tau phosphorylation. <i>Annals of Neurology</i> , <b>2014</b> , 76, 379-92	9.4	48
231	ABCC9 gene polymorphism is associated with hippocampal sclerosis of aging pathology. <i>Acta Neuropathologica</i> , <b>2014</b> , 127, 825-43	14.3	51
230	A case of inappropriate apolipoprotein e testing in Alzheimer's disease due to lack of an informed consent discussion. <i>American Journal of Alzheimern Disease and Other Dementias</i> , <b>2014</b> , 29, 590-5	2.5	
229	Two rare AKAP9 variants are associated with Alzheimer's disease in African Americans. <i>Alzheimern</i> a and Dementia, <b>2014</b> , 10, 609-618.e11	1.2	83
228	Search for age-related macular degeneration risk variants in Alzheimer disease genes and pathways. <i>Neurobiology of Aging</i> , <b>2014</b> , 35, 1510.e7-18	5.6	34
227	Genome-wide association study of opioid dependence: multiple associations mapped to calcium and potassium pathways. <i>Biological Psychiatry</i> , <b>2014</b> , 76, 66-74	7.9	143
226	Missense variant in TREML2 protects against Alzheimer's disease. <i>Neurobiology of Aging</i> , <b>2014</b> , 35, 151	0. <b>ę</b> .169-2	2 <b>6</b> 84
225	Rare and common variants in extracellular matrix gene Fibrillin 2 (FBN2) are associated with macular degeneration. <i>Human Molecular Genetics</i> , <b>2014</b> , 23, 5827-37	5.6	34
224	Data compatibility in the addiction sciences: an examination of measure commonality. <i>Drug and Alcohol Dependence</i> , <b>2014</b> , 141, 153-8	4.9	24
223	P2-031: A VARIANT IN STK24 ACHIEVES GENOME-WIDE SIGNIFICANCE IN AFRICAN AMERICANS USING A LIABILITY MODEL <b>2014</b> , 10, P481-P481		
222	O1-04-03: LOW-FREQUENCY VARIANT IMPUTATION IDENTIFIES NOVEL DISEASE-ASSOCIATED LOCI IN A GENOME-WIDE ASSOCIATION STUDY OF LATE-ONSET ALZHEIMER'S DISEASE <b>2014</b> , 10, P135-P13	5	
221	P1-045: EXOME ARRAY ANALYSIS IDENTIFIES NOVEL RISK VARIANTS FOR ALZHEIMER'S DISEASE WITH ONSET BEFORE 65 YEARS <b>2014</b> , 10, P319-P319		1
220	P1-054: LINKAGE ANALYSES OF EXTENDED CARIBBEAN HISPANIC FAMILIES INDICATES NOVEL LOCI ASSOCIATED WITH FAMILIAL LATE-ONSET ALZHEIMER'S DISEASE <b>2014</b> , 10, P323-P323		
219	P2-125: GENOME-WIDE LINKAGE ANALYSES IDENTIFY NOVEL LOCI FOR FAMILIAL LATE-ONSET ALZHEIMER'S DISEASE <b>2014</b> , 10, P517-P517		

218	FLT1 genetic variation predisposes to neovascular AMD in ethnically diverse populations and alters systemic FLT1 expression <b>2014</b> , 55, 3543-54		14
217	Genome-wide association meta-analysis of neuropathologic features of Alzheimer's disease and related dementias. <i>PLoS Genetics</i> , <b>2014</b> , 10, e1004606	6	219
216	Evaluating the role of a galanin enhancer genotype on a range of metabolic, depressive and addictive phenotypes. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2014</b> , 165B, 654-64	3.5	4
215	P1-035: BIVARIATE GENOME-WIDE ASSOCIATION STUDY OF ALZHEIMER DISEASE ENDOPHENOTYPES IDENTIFIES NOVEL LOCI <b>2014</b> , 10, P316-P316		
214	Reply: To PMID 23740775. Arthritis and Rheumatology, 2014, 66, 1401	9.5	1
213	A rare mutation in UNC5C predisposes to late-onset Alzheimer's disease and increases neuronal cell death. <i>Nature Medicine</i> , <b>2014</b> , 20, 1452-7	50.5	97
212	Deep resequencing of 17 glutamate system genes identifies rare variants in DISC1 and GRIN2B affecting risk of opioid dependence. <i>Addiction Biology</i> , <b>2014</b> , 19, 955-64	4.6	20
211	Association of MAPT haplotypes with Alzheimer's disease risk and MAPT brain gene expression levels. <i>Alzheimer Research and Therapy</i> , <b>2014</b> , 6, 39	9	78
210	Effects of multiple genetic loci on age at onset in late-onset Alzheimer disease: a genome-wide association study. <i>JAMA Neurology</i> , <b>2014</b> , 71, 1394-404	17.2	129
209	P2-131: WHOLE-EXOME SEQUENCING OF HISPANIC EARLY-ONSET ALZHEIMER DISEASE FAMILIES IDENTIFIES RARE VARIANTS IN MULTIPLE ALZHEIMER'S-RELATED GENES <b>2014</b> , 10, P518-P519		
208	Structural interactions between inhibitor and substrate docking sites give insight into mechanisms of human PS1 complexes. <i>Structure</i> , <b>2014</b> , 22, 125-35	5.2	55
207	Exploring the genetic architecture of alcohol dependence in African-Americans via analysis of a genomewide set of common variants. <i>Human Genetics</i> , <b>2014</b> , 133, 617-24	6.3	12
206	Gene-wide analysis detects two new susceptibility genes for Alzheimer's disease. <i>PLoS ONE</i> , <b>2014</b> , 9, e94661	3.7	90
205	BCL11A enhancer Haplotypes Are Associated with the Distribution of HbF in Arab-Indian and African Haplotype Sickle Cell Anemia but Not the Different Population Levels of HbF. <i>Blood</i> , <b>2014</b> , 124, 4066-4066	2.2	
204	Fetal hemoglobin in sickle cell anemia: genetic studies of the Arab-Indian haplotype. <i>Blood Cells, Molecules, and Diseases</i> , <b>2013</b> , 51, 22-6	2.1	40
203	Meta-analysis of 74,046 individuals identifies 11 new susceptibility loci for Alzheimer's disease. <i>Nature Genetics</i> , <b>2013</b> , 45, 1452-8	36.3	2714
202	Role of p73 in Alzheimer disease: lack of association in mouse models or in human cohorts. <i>Molecular Neurodegeneration</i> , <b>2013</b> , 8, 10	19	7
201	Integrating GWASs and human protein interaction networks identifies a gene subnetwork underlying alcohol dependence. <i>American Journal of Human Genetics</i> , <b>2013</b> , 93, 1027-34	11	55

200	GWAS of cerebrospinal fluid tau levels identifies risk variants for Alzheimer's disease. <i>Neuron</i> , <b>2013</b> , 78, 256-68	13.9	255
199	Seven new loci associated with age-related macular degeneration. <i>Nature Genetics</i> , <b>2013</b> , 45, 433-9, 439	986.3	577
198	Variants in the ATP-binding cassette transporter (ABCA7), apolipoprotein E ?4,and the risk of late-onset Alzheimer disease in African Americans. <i>JAMA - Journal of the American Medical Association</i> , <b>2013</b> , 309, 1483-92	27.4	275
197	Genome-wide association study identifies new susceptibility loci for posttraumatic stress disorder. <i>Biological Psychiatry</i> , <b>2013</b> , 74, 656-63	7.9	129
196	Identification of a rare coding variant in complement 3 associated with age-related macular degeneration. <i>Nature Genetics</i> , <b>2013</b> , 45, 1375-9	36.3	130
195	Association of granulomatosis with polyangiitis (Wegener's) with HLA-DPB1*04 and SEMA6A gene variants: evidence from genome-wide analysis. <i>Arthritis and Rheumatism</i> , <b>2013</b> , 65, 2457-68		102
194	Initial assessment of the pathogenic mechanisms of the recently identified Alzheimer risk Loci. <i>Annals of Human Genetics</i> , <b>2013</b> , 77, 85-105	2.2	40
193	A complex interplay between personality domains, marital status and a variant in CHRNA5 on the risks of cocaine, nicotine dependences and cocaine-induced paranoia. <i>PLoS ONE</i> , <b>2013</b> , 8, e49368	3.7	5
192	SORL1 is genetically associated with late-onset Alzheimer's disease in Japanese, Koreans and Caucasians. <i>PLoS ONE</i> , <b>2013</b> , 8, e58618	3.7	122
191	Prayer at midlife is associated with reduced risk of cognitive decline in Arabic women. <i>Current Alzheimer Research</i> , <b>2013</b> , 10, 340-6	3	25
190	GABRG1 and GABRA2 variation associated with alcohol dependence in African Americans. <i>Alcoholism: Clinical and Experimental Research</i> , <b>2012</b> , 36, 588-93	3.7	34
189	Common genetic variants in the CLDN2 and PRSS1-PRSS2 loci alter risk for alcohol-related and sporadic pancreatitis. <i>Nature Genetics</i> , <b>2012</b> , 44, 1349-54	36.3	223
188	Serum paraoxonase activity is associated with variants in the PON gene cluster and risk of Alzheimer disease. <i>Neurobiology of Aging</i> , <b>2012</b> , 33, 1015.e7-23	5.6	22
187	Identification of Alzheimer disease-associated variants in genes that regulate retromer function. <i>Neurobiology of Aging</i> , <b>2012</b> , 33, 2231.e15-2231.e30	5.6	115
186	Novel late-onset Alzheimer disease loci variants associate with brain gene expression. <i>Neurology</i> , <b>2012</b> , 79, 221-8	6.5	124
185	Comprehensive search for Alzheimer disease susceptibility loci in the APOE region. <i>Archives of Neurology</i> , <b>2012</b> , 69, 1270-9		81
184	A genome-wide association study of plasma total IgE concentrations in the Framingham Heart Study. <i>Journal of Allergy and Clinical Immunology</i> , <b>2012</b> , 129, 840-845.e21	11.5	120
183	Association of COL25A1 with comorbid antisocial personality disorder and substance dependence. <i>Biological Psychiatry</i> , <b>2012</b> , 71, 733-40	7.9	15

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182	Linkage analysis followed by association show NRG1 associated with cannabis dependence in African Americans. <i>Biological Psychiatry</i> , <b>2012</b> , 72, 637-44	7.9	39
181	Childhood adversity increases risk for nicotine dependence and interacts with 5 nicotinic acetylcholine receptor genotype specifically in males. <i>Neuropsychopharmacology</i> , <b>2012</b> , 37, 669-76	8.7	37
180	High prevalence of mild cognitive impairment and Alzheimer's disease in arabic villages in northern Israel: impact of gender and education. <i>Journal of Alzheimerm Disease</i> , <b>2012</b> , 29, 431-9	4.3	40
179	A functional promoter polymorphism of the Eglobin gene is a specific marker of the Arab-Indian haplotype. <i>American Journal of Hematology</i> , <b>2012</b> , 87, 824-6	7.1	10
178	Autosomal linkage scan for loci predisposing to comorbid dependence on multiple substances. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2012</b> , 159B, 361-9	3.5	12
177	Multiple loci influencing hippocampal degeneration identified by genome scan. <i>Annals of Neurology</i> , <b>2012</b> , 72, 65-75	9.4	50
176	Brain expression genome-wide association study (eGWAS) identifies human disease-associated variants. <i>PLoS Genetics</i> , <b>2012</b> , 8, e1002707	6	174
175	Evidence for a role of the rare p.A152T variant in MAPT in increasing the risk for FTD-spectrum and Alzheimer's diseases. <i>Human Molecular Genetics</i> , <b>2012</b> , 21, 3500-12	5.6	174
174	Behavioral variant frontotemporal lobar degeneration with amyotrophic lateral sclerosis with a chromosome 9p21 hexanucleotide repeat. <i>Frontiers in Neurology</i> , <b>2012</b> , 3, 136	4.1	5
173	ECatenin is genetically and biologically associated with cortical cataract and future Alzheimer-related structural and functional brain changes. <i>PLoS ONE</i> , <b>2012</b> , 7, e43728	3.7	42
172	ANKRD7 and CYTL1 are novel risk genes for alcohol drinking behavior. <i>Chinese Medical Journal</i> , <b>2012</b> , 125, 1127-34	2.9	10
171	Rare nonsynonymous variants in alpha-4 nicotinic acetylcholine receptor gene protect against nicotine dependence. <i>Biological Psychiatry</i> , <b>2011</b> , 70, 528-36	7.9	55
170	A 3-bp deletion in the HBS1L-MYB intergenic region on chromosome 6q23 is associated with HbF expression. <i>Blood</i> , <b>2011</b> , 117, 4935-45	2.2	102
169	Agitated depression in substance dependence. <i>Drug and Alcohol Dependence</i> , <b>2011</b> , 116, 163-9	4.9	9
168	Ancestry of African Americans with sickle cell disease. <i>Blood Cells, Molecules, and Diseases</i> , <b>2011</b> , 47, 47	l <b>-5</b> 2.1	23
167	A comprehensive genetic association study of Alzheimer disease in African Americans. <i>Archives of Neurology</i> , <b>2011</b> , 68, 1569-79		187
166	Association of TTR polymorphisms with hippocampal atrophy in Alzheimer disease families. <i>Neurobiology of Aging</i> , <b>2011</b> , 32, 249-56	5.6	14
165	Magnetic resonance imaging-measured atrophy and its relationship to cognitive functioning in vascular dementia and Alzheimer's disease patients. <i>Alzheimermand Dementia</i> , <b>2011</b> , 7, 493-500	1.2	9

164	Genetics and genomics of late-onset Alzheimer's disease and its endophenotypes. <i>International Journal of Alzheimer</i> Disease, <b>2011</b> , 2011, 284728	3.7	7
163	ACSL6 is associated with the number of cigarettes smoked and its expression is altered by chronic nicotine exposure. <i>PLoS ONE</i> , <b>2011</b> , 6, e28790	3.7	8
162	Hamilton et al. Respond to "Consolidating Data Harmonization". <i>American Journal of Epidemiology</i> , <b>2011</b> , 174, 265-266	3.8	10
161	Systems biology-based analysis implicates a novel role for vitamin D metabolism in the pathogenesis of age-related macular degeneration. <i>Human Genomics</i> , <b>2011</b> , 5, 538-68	6.8	50
160	Glucuronic acid epimerase is associated with plasma triglyceride and high-density lipoprotein cholesterol levels in Turks. <i>Annals of Human Genetics</i> , <b>2011</b> , 75, 398-417	2.2	14
159	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. <i>Nature Genetics</i> , <b>2011</b> , 43, 436-41	36.3	1367
158	Power and pitfalls of the genome-wide association study approach to identify genes for Alzheimer's disease. <i>Current Psychiatry Reports</i> , <b>2011</b> , 13, 138-46	9.1	19
157	Association between polymorphisms in catechol-O-methyltransferase (COMT) and cocaine-induced paranoia in European-American and African-American populations. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2011</b> , 156B, 651-60	3.5	24
156	SORCS1 alters amyloid precursor protein processing and variants may increase Alzheimer's disease risk. <i>Annals of Neurology</i> , <b>2011</b> , 69, 47-64	9.4	79
155	Identification of novel candidate genes for Alzheimer's disease by autozygosity mapping using genome wide SNP data. <i>Journal of Alzheimerm Disease</i> , <b>2011</b> , 23, 349-59	4.3	37
154	Meta-analysis of the association between variants in SORL1 and Alzheimer disease. <i>Archives of Neurology</i> , <b>2011</b> , 68, 99-106		135
153	The PhenX Toolkit: get the most from your measures. <i>American Journal of Epidemiology</i> , <b>2011</b> , 174, 253-	- <b>6</b> Ø	397
152	A genomewide linkage scan of cocaine dependence and major depressive episode in two populations. <i>Neuropsychopharmacology</i> , <b>2011</b> , 36, 2422-30	8.7	24
151	Influence of ROBO1 and RORA on risk of age-related macular degeneration reveals genetically distinct phenotypes in disease pathophysiology. <i>PLoS ONE</i> , <b>2011</b> , 6, e25775	3.7	22
150	Mild cognitive impairment is associated with mild parkinsonian signs in a door-to-door study. Journal of Alzheimerm Disease, <b>2010</b> , 22, 1005-13	4.3	15
149	Genome-wide analysis of genetic loci associated with Alzheimer disease. <i>JAMA - Journal of the American Medical Association</i> , <b>2010</b> , 303, 1832-40	27.4	888
148	Confirmation and generalization of an alcohol-dependence locus on chromosome 10q. <i>Neuropsychopharmacology</i> , <b>2010</b> , 35, 1325-32	8.7	9
147	Variation in nicotinic acetylcholine receptor genes is associated with multiple substance dependence phenotypes. <i>Neuropsychopharmacology</i> , <b>2010</b> , 35, 1921-31	8.7	90

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146	Interaction of FKBP5 with childhood adversity on risk for post-traumatic stress disorder. <i>Neuropsychopharmacology</i> , <b>2010</b> , 35, 1684-92	8.7	262	
145	Meta-analysis confirms CR1, CLU, and PICALM as alzheimer disease risk loci and reveals interactions with APOE genotypes. <i>Archives of Neurology</i> , <b>2010</b> , 67, 1473-84		330	
144	Alzheimer's Disease Neuroimaging Initiative biomarkers as quantitative phenotypes: Genetics core aims, progress, and plans. <i>Alzheimermand Dementia</i> , <b>2010</b> , 6, 265-73	1.2	279	
143	Genetic variants near TIMP3 and high-density lipoprotein-associated loci influence susceptibility to age-related macular degeneration. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2010</b> , 107, 7401-6	11.5	417	
142	Fetal hemoglobin in sickle cell anemia: genome-wide association studies suggest a regulatory region in the 5' olfactory receptor gene cluster. <i>Blood</i> , <b>2010</b> , 115, 1815-22	2.2	132	
141	Convergence of linkage, gene expression and association data demonstrates the influence of the RAR-related orphan receptor alpha (RORA) gene on neovascular AMD: a systems biology based approach. <i>Vision Research</i> , <b>2010</b> , 50, 698-715	2.1	37	
140	Genetic modifiers of Hb E/beta0 thalassemia identified by a two-stage genome-wide association study. <i>BMC Medical Genetics</i> , <b>2010</b> , 11, 51	2.1	19	
139	Fetal Hemoglobin In Sickle Cell Anemia: Molecular Characterization of Saudi Patients From the Eastern Province. <i>Blood</i> , <b>2010</b> , 116, 1627-1627	2.2		
138	A 3-Bp Deletion Between Transcription Factor Binding Motifs In the HBS1L-MYB Intergenic Region on Chromosome 6q23 Is Associated with HbF Expression. <i>Blood</i> , <b>2010</b> , 116, 1013-1013	2.2		
137	Education attenuates the effect of medial temporal lobe atrophy on cognitive function in Alzheimer's disease: the MIRAGE study. <i>Journal of Alzheimer Disease</i> , <b>2009</b> , 17, 855-62	4.3	35	
136	Performance of random forest when SNPs are in linkage disequilibrium. <i>BMC Bioinformatics</i> , <b>2009</b> , 10, 78	3.6	62	
135	Essential tremor might be less frequent than Parkinson's disease in North Israel Arab villages. <i>Movement Disorders</i> , <b>2009</b> , 24, 119-22	7	19	
134	Pro-opiomelanocortin gene variation related to alcohol or drug dependence: evidence and replications across family- and population-based studies. <i>Biological Psychiatry</i> , <b>2009</b> , 66, 128-36	7.9	27	
133	Disclosure of APOE genotype for risk of Alzheimer's disease. <i>New England Journal of Medicine</i> , <b>2009</b> , 361, 245-54	59.2	418	
132	Association of variants in MANEA with cocaine-related behaviors. <i>Archives of General Psychiatry</i> , <b>2009</b> , 66, 267-74		18	
131	The Sortilin-Related Receptor SORL1 is Functionally and Genetically Associated with Alzheimer's Disease. <i>Research and Perspectives in Alzheimer</i> Disease, <b>2009</b> , 157-165			
130	Transient cocaine-associated behavioral symptoms rated with a new instrument, the scale for assessment of positive symptoms for cocaine-induced psychosis (SAPS-CIP). <i>American Journal on Addictions</i> , <b>2009</b> , 18, 339-45	3.7	18	
129	Correlates of co-occurring ADHD in drug-dependent subjects: prevalence and features of substance dependence and psychiatric disorders. <i>Addictive Behaviors</i> , <b>2008</b> , 33, 1199-207	4.2	161	

128	BCL11A is a major HbF quantitative trait locus in three different populations with beta-hemoglobinopathies. <i>Blood Cells, Molecules, and Diseases</i> , <b>2008</b> , 41, 255-258	2.1	137
127	Association of distinct variants in SORL1 with cerebrovascular and neurodegenerative changes related to Alzheimer disease. <i>Archives of Neurology</i> , <b>2008</b> , 65, 1640-8		53
126	Further examination of the candidate genes in chromosome 12p13 locus for late-onset Alzheimer disease. <i>Neurogenetics</i> , <b>2008</b> , 9, 127-38	3	23
125	Substance dependence low-density whole genome association study in two distinct American populations. <i>Human Genetics</i> , <b>2008</b> , 123, 495-506	6.3	21
124	Variation and heritability of Hb F and F-cells among beta-thalassemia heterozygotes in Hong Kong. <i>American Journal of Hematology</i> , <b>2008</b> , 83, 458-64	7.1	26
123	Magnetic resonance imaging traits in siblings discordant for Alzheimer disease. <i>Journal of Neuroimaging</i> , <b>2008</b> , 18, 268-75	2.8	18
122	Estimated glomerular filtration rate in sickle cell anemia is associated with polymorphisms of bone morphogenetic protein receptor 1B. <i>American Journal of Hematology</i> , <b>2007</b> , 82, 179-84	7.1	42
121	The neuronal sortilin-related receptor SORL1 is genetically associated with Alzheimer disease. <i>Nature Genetics</i> , <b>2007</b> , 39, 168-77	36.3	888
120	Beta-globin gene cluster polymorphisms are strongly associated with severity of HbE/beta(0)-thalassemia. <i>Clinical Genetics</i> , <b>2007</b> , 72, 497-505	4	26
119	Comorbid psychiatric diagnoses and their association with cocaine-induced psychosis in cocaine-dependent subjects. <i>American Journal on Addictions</i> , <b>2007</b> , 16, 343-51	3.7	41
118	A network model to predict the risk of death in sickle cell disease. <i>Blood</i> , <b>2007</b> , 110, 2727-35	2.2	132
117	Heritability of magnetic resonance imaging (MRI) traits in Alzheimer disease cases and their siblings in the MIRAGE study. <i>Alzheimer Disease and Associated Disorders</i> , <b>2007</b> , 21, 85-91	2.5	25
116	Association between SORL1 and Alzheimer's disease in a genome-wide study. <i>NeuroReport</i> , <b>2007</b> , 18, 1761-4	1.7	76
115	Education effects on cognitive function in a healthy aged Arab population. <i>International Psychogeriatrics</i> , <b>2007</b> , 19, 593-603	3.4	35
114	Potential ethnic modifiers in the assessment and treatment of Alzheimer's disease: challenges for the future. <i>International Psychogeriatrics</i> , <b>2007</b> , 19, 539-58	3.4	48
113	Association studies between the plasmin genes and late-onset Alzheimer's disease. <i>Neurobiology of Aging</i> , <b>2007</b> , 28, 1041-3	5.6	9
112	Serum heat shock protein 70 level as a biomarker of exceptional longevity. <i>Mechanisms of Ageing and Development</i> , <b>2006</b> , 127, 862-8	5.6	58
111	Predictors of subjective memory complaint in cognitively normal relatives of patients with Alzheimer's disease. <i>Journal of Neuropsychiatry and Clinical Neurosciences</i> , <b>2006</b> , 18, 384-8	2.7	13

#### (2005-2006)

110	Diabetes mellitus and risk of developing Alzheimer disease: results from the Framingham Study. <i>Archives of Neurology</i> , <b>2006</b> , 63, 1551-5		218
109	Association of polymorphisms of IGF1R and genes in the transforming growth factor- beta /bone morphogenetic protein pathway with bacteremia in sickle cell anemia. <i>Clinical Infectious Diseases</i> , <b>2006</b> , 43, 593-8	11.6	45
108	Intronic variants in the dopa decarboxylase (DDC) gene are associated with smoking behavior in European-Americans and African-Americans. <i>Human Molecular Genetics</i> , <b>2006</b> , 15, 2192-9	5.6	43
107	Polymorphisms in the PON gene cluster are associated with Alzheimer disease. <i>Human Molecular Genetics</i> , <b>2006</b> , 15, 77-85	5.6	78
106	Association of polymorphisms in the Angiotensin-converting enzyme gene with Alzheimer disease in an Israeli Arab community. <i>American Journal of Human Genetics</i> , <b>2006</b> , 78, 871-877	11	60
105	Statin use and the risk of Alzheimer's disease: the MIRAGE study. Alzheimermand Dementia, <b>2006</b> , 2, 96-	-11023	38
104	Association of asthma with a functional promoter polymorphism in the IL16 gene. <i>Journal of Allergy and Clinical Immunology</i> , <b>2006</b> , 117, 86-91	11.5	30
103	Genetic association between endothelial nitric oxide synthase and Alzheimer disease. <i>Clinical Genetics</i> , <b>2006</b> , 70, 49-56	4	23
102	Sickle cell leg ulcers: associations with haemolysis and SNPs in Klotho, TEK and genes of the TGF-beta/BMP pathway. <i>British Journal of Haematology</i> , <b>2006</b> , 133, 570-8	4.5	130
101	Fetal Hemoglobin in Sickle Cell Anemia: Associations with Single Nucleotide Polymorphisms in Quantitative Trait Loci on Chromsomes 8q12 and Xp22 <i>Blood</i> , <b>2006</b> , 108, 1222-1222	2.2	1
100	Severity of Sickle Cell Disease: Modeling Interrelationships among Hemolysis, Pulmonary Hypertension and Risk of Death <i>Blood</i> , <b>2006</b> , 108, 786-786	2.2	
99	Complement factor H polymorphism and age-related macular degeneration. <i>Science</i> , <b>2005</b> , 308, 421-4	33.3	1996
98	Rating the severity and character of transient cocaine-induced delusions and hallucinations with a new instrument, the Scale for Assessment of Positive Symptoms for Cocaine-Induced Psychosis (SAPS-CIP). <i>Drug and Alcohol Dependence</i> , <b>2005</b> , 80, 23-33	4.9	50
97	Hemolysis-associated priapism in sickle cell disease. <i>Blood</i> , <b>2005</b> , 106, 3264-7	2.2	160
96	Association of single nucleotide polymorphisms in klotho with priapism in sickle cell anaemia. <i>British Journal of Haematology</i> , <b>2005</b> , 128, 266-72	4.5	63
95	Relation between atherogenic dyslipidemia and the Adult Treatment Program-III definition of metabolic syndrome (Genetic Epidemiology of Metabolic Syndrome Project). <i>American Journal of Cardiology</i> , <b>2005</b> , 95, 194-8	3	60
94	Analysis of the glucocerebrosidase gene in Parkinson's disease. <i>Movement Disorders</i> , <b>2005</b> , 20, 367-70	7	93
93	Multifactor-dimensionality reduction versus family-based association tests in detecting susceptibility loci in discordant sib-pair studies. <i>BMC Genetics</i> , <b>2005</b> , 6 Suppl 1, S146	2.6	4

92	Whole-genome variance components linkage analysis using single-nucleotide polymorphisms versus microsatellites on quantitative traits of derived phenotypes from factor analysis of electroencephalogram waves. <i>BMC Genetics</i> , <b>2005</b> , 6 Suppl 1, S15	2.6	3
91	Genome-wide linkage analysis for alcohol dependence: a comparison between single-nucleotide polymorphism and microsatellite marker assays. <i>BMC Genetics</i> , <b>2005</b> , 6 Suppl 1, S8	2.6	4
90	Nonsteroidal anti-inflammatory drug use and Alzheimer's disease risk: the MIRAGE Study. <i>BMC Geriatrics</i> , <b>2005</b> , 5, 2	4.1	56
89	Polymorphisms in the promoter region of catalase gene and essential hypertension. <i>Disease Markers</i> , <b>2005</b> , 21, 3-7	3.2	40
88	Multiple QTLs influencing triglyceride and HDL and total cholesterol levels identified in families with atherogenic dyslipidemia. <i>Journal of Lipid Research</i> , <b>2005</b> , 46, 2202-13	6.3	37
87	Gene-Gene Interactions and the Pathophysiology of Sickle Cell Disease: Modeling the Effects of SNPs on Sickle Cell-Associated Vasoocclusive Events Using Classification and Regression Trees and Stochastic Gradient Boosting <i>Blood</i> , <b>2005</b> , 106, 3183-3183	2.2	2
86	Lack of association between angiotensin-converting enzyme and dementia of the Alzheimer's type in an elderly Arab population in Wadi Ara, Israel. <i>Neuropsychiatric Disease and Treatment</i> , <b>2005</b> , 1, 73-6	3.1	2
85	Leg Ulcers in Sickle Cell Anemia Are Associated with Laboratory Markers of Hemolysis and SNPs in KL and Genes of the TGF- <b>/</b> BMP Pathway <i>Blood</i> , <b>2005</b> , 106, 2317-2317	2.2	1
84	Polymorphisms (Snps) in Multiple Genes of the Tgf-[/Bmp Pathway Are Associated with a Global Measure of Sickle Cell Disease Severity <i>Blood</i> , <b>2005</b> , 106, 74-74	2.2	
83	Association of Polymorphisms of the Transforming Growth Factor- <b>/</b> Bone Morphogenetic Protein (TGF- <b>/</b> BMP) Pathway with Sickle Cell Bacteremia <i>Blood</i> , <b>2005</b> , 106, 3170-3170	2.2	
82	Who seeks genetic susceptibility testing for Alzheimer's disease? Findings from a multisite, randomized clinical trial. <i>Genetics in Medicine</i> , <b>2004</b> , 6, 197-203	8.1	86
81	Differential modulation of endotoxin responsiveness by human caspase-12 polymorphisms. <i>Nature</i> , <b>2004</b> , 429, 75-9	50.4	344
80	Estimating risk curves for first-degree relatives of patients with Alzheimer's disease: the REVEAL study. <i>Genetics in Medicine</i> , <b>2004</b> , 6, 192-6	8.1	129
79	Genetic Polymorphisms Associated with Fetal Hemoglobin Response to Hydroxyurea in Patients with Sickle Cell Anemia <i>Blood</i> , <b>2004</b> , 104, 108-108	2.2	5
78	Association of Single Nucleotide Polymorphisms in Klotho with Priapism in Sickle Cell Anemia <i>Blood</i> , <b>2004</b> , 104, 1673-1673	2.2	
77	The genetics of adult-onset neuropsychiatric disease: complexities and conundra?. <i>Science</i> , <b>2003</b> , 302, 822-6	33.3	147
76	Reasons for seeking genetic susceptibility testing among first-degree relatives of people with Alzheimer disease. <i>Alzheimer Disease and Associated Disorders</i> , <b>2003</b> , 17, 86-93	2.5	70
75	Genetic and environmental epidemiology of Alzheimer's disease in arabs residing in Israel. <i>Journal of Molecular Neuroscience</i> , <b>2003</b> , 20, 207-12	3.3	19

74	Search for genetic factors predisposing to atherogenic dyslipidemia. <i>BMC Genetics</i> , <b>2003</b> , 4 Suppl 1, S10	<b>ጋው</b> .6	9
73	Genome-wide screen for heavy alcohol consumption. <i>BMC Genetics</i> , <b>2003</b> , 4 Suppl 1, S106	2.6	18
72	Empirically derived phenotypic subgroups - qualitative and quantitative trait analyses. <i>BMC Genetics</i> , <b>2003</b> , 4 Suppl 1, S15	2.6	13
71	Identification of multiple loci for Alzheimer disease in a consanguineous Israeli-Arab community. <i>Human Molecular Genetics</i> , <b>2003</b> , 12, 415-22	5.6	96
70	Genetic variants of WNK4 in whites and African Americans with hypertension. <i>Hypertension</i> , <b>2003</b> , 41, 1191-5	8.5	27
69	Differences between African Americans and Whites in their attitudes toward genetic testing for Alzheimer's disease. <i>Genetic Testing and Molecular Biomarkers</i> , <b>2003</b> , 7, 39-44		33
68	Risk of dementia among white and African American relatives of patients with Alzheimer disease. JAMA - Journal of the American Medical Association, 2002, 287, 329-36	27.4	255
67	Association between apolipoprotein E genotype and Alzheimer disease in African American subjects. <i>Archives of Neurology</i> , <b>2002</b> , 59, 594-600		76
66	Reliability of information collected by proxy in family studies of Alzheimer's disease. <i>Neuroepidemiology</i> , <b>2001</b> , 20, 105-11	5.4	28
65	Distinguishable effects of presenilin-1 and APP717 mutations on amyloid plaque deposition. <i>Neurobiology of Aging</i> , <b>2001</b> , 22, 367-76	5.6	17
64	Association between angiotensin-converting enzyme and Alzheimer disease. <i>Archives of Neurology</i> , <b>2000</b> , 57, 210-4		88
63	Familial risk for Alzheimer disease in ethnic minorities: nondiscriminating genes. <i>Archives of Neurology</i> , <b>2000</b> , 57, 28-9		18
62	An alpha-2-macroglobulin insertion-deletion polymorphism in Alzheimer disease. <i>Nature Genetics</i> , <b>1999</b> , 22, 19-22	36.3	107
61	A family history study of male sexual orientation using three independent samples. <i>Behavior Genetics</i> , <b>1999</b> , 29, 79-86	3.2	65
60	No association between the HLA-A2 allele and Alzheimer disease. <i>Neurogenetics</i> , <b>1999</b> , 2, 177-82	3	15
59	Alpha-2 macroglobulin gene in early- and late-onset Alzheimer disease. <i>Neuroscience Letters</i> , <b>1999</b> , 271, 129-31	3.3	22
58	Power of concordant versus discordant sib pairs at different penetrance levels. <i>Genetic Epidemiology</i> , <b>1999</b> , 17 Suppl 1, S679-84	2.6	1
57	Locating Genetic Modifiers for Inherited Neurodegenerative Diseases. <i>Cerebral Cortex</i> , <b>1999</b> , 433-459		1

56	Association between bleomycin hydrolase and Alzheimer's disease in caucasians. <i>Annals of Neurology</i> , <b>1998</b> , 44, 808-11	9.4	39
55	Novel ATP7B mutations causing Wilson disease in several Israeli ethnic groups. <i>Human Mutation</i> , <b>1998</b> , 11, 145-51	4.7	35
54	No genetic association between the LRP receptor and sporadic or late-onset familial Alzheimer disease. <i>Neurogenetics</i> , <b>1998</b> , 1, 179-83	3	38
53	Amyloid-beta-protein isoforms in brain of subjects with PS1-linked, beta APP-linked and sporadic Alzheimer disease. <i>Molecular Brain Research</i> , <b>1998</b> , 56, 178-85		23
52	Smoking and risk of Alzheimer's disease. MIRAGE Study Group. <i>Lancet, The</i> , <b>1998</b> , 352, 819	40	4
51	Autosomal dominant orthostatic hypotensive disorder maps to chromosome 18q. <i>American Journal of Human Genetics</i> , <b>1998</b> , 63, 1425-30	11	41
50	Novel ATP7B mutations causing Wilson disease in several Israeli ethnic groups <b>1998</b> , 11, 145		2
49	Effects of Age, Sex, and Ethnicity on the Association Between Apolipoprotein E Genotype and Alzheimer Disease. <i>JAMA - Journal of the American Medical Association</i> , <b>1997</b> , 278, 1349	27.4	2208
48	GENETICS AND THE DEMENTIA PATIENT. Neurologist, 1997, 3, 13-30	1.6	22
47	Presenilin polymorphisms in Alzheimer's disease. <i>Lancet, The</i> , <b>1997</b> , 350, 959	40	17
46	Detecting linkage for a complex disease using simulated extended pedigrees. <i>Genetic Epidemiology</i> , <b>1997</b> , 14, 981-6	2.6	1
45	No association between alpha 1-antichymotrypsin and familial Alzheimer's disease. <i>Annals of the New York Academy of Sciences</i> , <b>1996</b> , 802, 35-41	6.5	8
44	Linkage disequilibrium analysis in Machado-Joseph disease patients of different ethnic origins. <i>Human Genetics</i> , <b>1996</b> , 98, 620-4	6.3	23
43	Gender equality in Machado-Joseph disease. <i>Nature Genetics</i> , <b>1995</b> , 11, 118-9	36.3	12
42	Statement on Use of Apolipoprotein E Testing for Alzheimer Disease. <i>JAMA - Journal of the American Medical Association</i> , <b>1995</b> , 274, 1627	27.4	128
41	Linkage of congenital, recessive deafness (DFNB4) to chromosome 7q31 and evidence for genetic heterogeneity in the Middle Eastern Druze population. <i>Human Molecular Genetics</i> , <b>1995</b> , 4, 1637-42	5.6	90
40	Allele epsilon 4 of apolipoprotein E shows a dose effect on age at onset of Pick disease. <i>Experimental Neurology</i> , <b>1995</b> , 136, 162-70	5.7	37
39	Apolipoprotein E genotype in patients with Alzheimer's disease: implications for the risk of dementia among relatives. <i>Annals of Neurology</i> , <b>1995</b> , 38, 797-808	9.4	65

38	Machado Joseph disease is not an allele of the spinocerebellar ataxia 2 locus. <i>Human Genetics</i> , <b>1994</b> , 93, 335-8	6.3	8
37	46,XY/47,XYY male with the fragile X syndrome: cytogenetic and molecular studies. <i>American Journal of Medical Genetics Part A</i> , <b>1993</b> , 45, 589-93		6
36	Susceptibility genes for familial Alzheimer's disease on chromosomes 19 and 21: a reality check. <i>Genetic Epidemiology</i> , <b>1993</b> , 10, 425-30	2.6	3
35	Identification of the genetic locus for keratosis palmaris et plantaris on chromosome 17 near the RARA and keratin type I genes. <i>Nature Genetics</i> , <b>1993</b> , 5, 158-62	36.3	38
34	The Machado-Joseph disease locus is different from the spinocerebellar ataxia locus (SCA1). <i>Genomics</i> , <b>1992</b> , 13, 852-5	4.3	9
33	The human cationic amino acid transporter (ATRC1): physical and genetic mapping to 13q12-q14. <i>Genomics</i> , <b>1992</b> , 12, 430-4	4.3	34
32	Gene Localization By Linkage Analysis. Otolaryngologic Clinics of North America, <b>1992</b> , 25, 907-922	2	2
31	Association of decreased paternal age and late-onset Alzheimer's disease. An example of genetic imprinting?. <i>Archives of Neurology</i> , <b>1991</b> , 48, 599-604		31
30	A contiguous linkage map of chromosome 13q with 39 distinct loci separated on average by 5.1 centimorgans. <i>Genomics</i> , <b>1991</b> , 11, 517-29	4.3	22
29	Estimation of familial risk in Alzheimer's disease. <i>Annals of Neurology</i> , <b>1990</b> , 27, 338-40		
	Escillation of Familiatrisk in Alzheimer's disease. Annais of Neurology, 1990, 21, 330-40	9.4	4
28	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. <i>Genomics</i> , <b>1990</b> , 7, 110-4	9·4 4·3	15
ſ	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13.		
28	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. <i>Genomics</i> , <b>1990</b> , 7, 110-4  Assessment of genetic risk for Alzheimer's disease among first-degree relatives. <i>Annals of</i>	4.3	15
28	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. Genomics, 1990, 7, 110-4  Assessment of genetic risk for Alzheimer's disease among first-degree relatives. Annals of Neurology, 1989, 25, 485-93	4·3 9·4	15 124
28 27 26	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. <i>Genomics</i> , <b>1990</b> , 7, 110-4  Assessment of genetic risk for Alzheimer's disease among first-degree relatives. <i>Annals of Neurology</i> , <b>1989</b> , 25, 485-93  Familial Alzheimer's disease: progress and problems. <i>Neurobiology of Aging</i> , <b>1989</b> , 10, 417-25  Reliability of self-reported age at onset of major depression. <i>Journal of Psychiatric Research</i> , <b>1989</b> ,	4·3 9·4 5.6	15 124 67
28 27 26 25	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. <i>Genomics</i> , 1990, 7, 110-4  Assessment of genetic risk for Alzheimer's disease among first-degree relatives. <i>Annals of Neurology</i> , 1989, 25, 485-93  Familial Alzheimer's disease: progress and problems. <i>Neurobiology of Aging</i> , 1989, 10, 417-25  Reliability of self-reported age at onset of major depression. <i>Journal of Psychiatric Research</i> , 1989, 23, 35-47  Predicting genotypes at loci for autosomal recessive disorders using linked genetic markers:	4·3 9·4 5.6	15 124 67 33
28 27 26 25 24	An alpha satellite DNA polymorphism specific for the centromeric region of chromosome 13. <i>Genomics</i> , 1990, 7, 110-4  Assessment of genetic risk for Alzheimer's disease among first-degree relatives. <i>Annals of Neurology</i> , 1989, 25, 485-93  Familial Alzheimer's disease: progress and problems. <i>Neurobiology of Aging</i> , 1989, 10, 417-25  Reliability of self-reported age at onset of major depression. <i>Journal of Psychiatric Research</i> , 1989, 23, 35-47  Predicting genotypes at loci for autosomal recessive disorders using linked genetic markers: application to Wilson's disease. <i>Human Genetics</i> , 1988, 79, 109-17	4·3 9·4 5.6 5·2 6·3	15 124 67 33 20

20	Suicide and attempted suicide in Huntington disease: implications for preclinical testing of persons at risk. <i>American Journal of Medical Genetics Part A</i> , <b>1986</b> , 24, 305-11		207
19	Clinical anthropometry and medical genetics: a compilation of body measurements in genetic and congenital disorders. <i>American Journal of Medical Genetics Part A</i> , <b>1986</b> , 25, 343-59		23
18	Evidence for linkage between Wilson disease and esterase D in three kindreds: detection of linkage for an autosomal recessive disorder by the family study method. <i>Genetic Epidemiology</i> , <b>1986</b> , 3, 201-9	2.6	31
17	Development of a map of chromosome 11p. <i>Genetic Epidemiology</i> , <b>1986</b> , 1, 153-8	2.6	17
16	Diabetes mellitus in Huntington disease. <i>Clinical Genetics</i> , <b>1985</b> , 27, 62-7	4	121
15	Anthropometric discrimination among affected, at-risk, and not-at-risk individuals in families with Huntington disease. <i>American Journal of Medical Genetics Part A</i> , <b>1985</b> , 21, 307-16		39
14	An anthropometric assessment of Huntington's disease patients and families. <i>American Journal of Physical Anthropology</i> , <b>1985</b> , 67, 185-94	2.5	48
13	Automating data manipulation for genetic analysis using a data base management system. <i>Human Heredity</i> , <b>1985</b> , 35, 296-301	1.1	5
12	The natural history of Huntington disease: possible role of "aging genes". <i>American Journal of Medical Genetics Part A</i> , <b>1984</b> , 18, 115-23		46
11	GWAS including 82,707 subjects identifies functional coding variant in OPRM1 gene associated with opioid use disorder		2
10	Expanding the Genetic Architecture of Nicotine Dependence and its Shared Genetics with Multiple Traits: Findings from the Nicotine Dependence GenOmics (iNDiGO) Consortium		3
9	Exome sequencing identifies rare damaging variants in the ATP8B4 and ABCA1 genes as novel risk factors for Alzheimer Disease		1
8	Cell-type Specific Expression Quantitative Trait Loci Associated with Alzheimer Disease in Blood and Brain Tissue		2
7	Trans-ancestral GWAS of alcohol dependence reveals common genetic underpinnings with psychiatric disorders		7
6	Largest genome-wide association study for PTSD identifies genetic risk loci in European and African ancestries and implicates novel biological pathways		6
5	Sex-dependent polygenic effects on the clinical progressions of Alzheimer disease		1
4	Causal associations between potentially modifiable risk factors and the Alzheimer disease phenome: A Mendelian randomization study		4
3	Leveraging genome-wide data to investigate differences between opioid use vs. opioid dependence in 41,176 individuals from the Psychiatric Genomics Consortium		2

2 A common haplotype lowers PU.1 expression in myeloid cells and delays onset of Alzheimer disease

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Genome-Wide Meta-Analysis of Late-Onset Alzheimer Disease Using Rare Variant Imputation in 65,602 Subjects Identifies Novel Rare Variant Locus NCK2: The International Genomics of Alzheimer Project (IGAP)

2