Richard Sherva

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8202168/publications.pdf

Version: 2024-02-01

56 papers 3,966 citations

30 h-index 51 g-index

58 all docs

58 docs citations

58 times ranked 7422 citing authors

#	Article	IF	CITATIONS
1	Transancestral GWAS of alcohol dependence reveals common genetic underpinnings with psychiatric disorders. Nature Neuroscience, 2018, 21, 1656-1669.	14.8	490
2	Common variants in WFS1 confer risk of type 2 diabetes. Nature Genetics, 2007, 39, 951-953.	21.4	333
3	Multiple Independent Loci at Chromosome 15q25.1 Affect Smoking Quantity: a Meta-Analysis and Comparison with Lung Cancer and COPD. PLoS Genetics, 2010, 6, e1001053.	3.5	332
4	A large-scale genome-wide association study meta-analysis of cannabis use disorder. Lancet Psychiatry,the, 2020, 7, 1032-1045.	7.4	200
5	Genome-Wide Association Study of Opioid Dependence: Multiple Associations Mapped to Calcium and Potassium Pathways. Biological Psychiatry, 2014, 76, 66-74.	1.3	192
6	Variants in the CD36 gene associate with the metabolic syndrome and high-density lipoprotein cholesterol. Human Molecular Genetics, 2008, 17, 1695-1704.	2.9	164
7	Genome-wide Association Study of Cannabis Dependence Severity, Novel Risk Variants, and Shared Genetic Risks. JAMA Psychiatry, 2016, 73, 472.	11.0	148
8	Genomeâ€wide association study of the rate of cognitive decline in Alzheimer's disease. Alzheimer's and Dementia, 2014, 10, 45-52.	0.8	147
9	Fetal hemoglobin in sickle cell anemia: genome-wide association studies suggest a regulatory region in the $5\hat{a} \in \mathbb{Z}^2$ olfactory receptor gene cluster. Blood, 2010, 115, 1815-1822.	1.4	146
10	Association of Granulomatosis With Polyangiitis (Wegener's) With ⟨i⟩HLA–DPB1*04⟨ i⟩ and ⟨i⟩SEMA6A⟨ i⟩ Gene Variants: Evidence From Genomeâ€Wide Analysis. Arthritis and Rheumatism, 2013, 65, 2457-2468.	6.7	138
11	Association of a single nucleotide polymorphism in neuronal acetylcholine receptor subunit alpha 5 (CHRNA5) with smoking status and with †pleasurable buzz†during early experimentation with smoking. Addiction, 2008, 103, 1544-1552.	3.3	129
12	Common CD36 SNPs reduce protein expression and may contribute to a protective atherogenic profile. Human Molecular Genetics, 2011, 20, 193-201.	2.9	126
13	A 3-bp deletion in the HBS1L-MYB intergenic region on chromosome 6q23 is associated with HbF expression. Blood, 2011, 117, 4935-4945.	1.4	116
14	Variation in Nicotinic Acetylcholine Receptor Genes is Associated with Multiple Substance Dependence Phenotypes. Neuropsychopharmacology, 2010, 35, 1921-1931.	5.4	103
15	Two novel loci, <i>COBL</i> and <i>SLC10A2</i> for Alzheimer's disease in African Americans. Alzheimer's and Dementia, 2017, 13, 119-129.	0.8	87
16	Expanding the genetic architecture of nicotine dependence and its shared genetics with multiple traits. Nature Communications, 2020, 11, 5562.	12.8	80
17	Genome-Wide Association Study of Nicotine Dependence in American Populations: Identification of Novel Risk Loci in Both African-Americans and European-Americans. Biological Psychiatry, 2015, 77, 493-503.	1.3	78
18	Genetic Risk Variants Associated With Comorbid Alcohol Dependence and Major Depression. JAMA Psychiatry, 2017, 74, 1234.	11.0	74

#	Article	IF	CITATIONS
19	Genome-wide Association Study Identifies a Regulatory Variant of RGMA Associated With Opioid Dependence in European Americans. Biological Psychiatry, 2018, 84, 762-770.	1.3	64
20	Genomewide Association Study for Maximum Number of Alcoholic Drinks in European Americans and African Americans. Alcoholism: Clinical and Experimental Research, 2015, 39, 1137-1147.	2.4	58
21	The genetics of alcohol dependence: Twin and SNPâ€based heritability, and genomeâ€wide association study based on AUDIT scores. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2015, 168, 739-748.	1.7	56
22	Variations in opioid receptor genes in neonatal abstinence syndrome. Drug and Alcohol Dependence, 2015, 155, 253-259.	3.2	55
23	Associations and Interactions Between SNPs in the Alcohol Metabolizing Genes and Alcoholism Phenotypes in European Americans. Alcoholism: Clinical and Experimental Research, 2009, 33, 848-857.	2.4	46
24	Identification of Novel Candidate Genes for Alzheimer's Disease by Autozygosity Mapping using Genome Wide SNP Data. Journal of Alzheimer's Disease, 2011, 23, 349-359.	2.6	46
25	Oxytocin receptor gene polymorphisms, attachment, and PTSD: Results from the National Health and Resilience in Veterans Study. Journal of Psychiatric Research, 2017, 94, 139-147.	3.1	46
26	Calibrating Longitudinal Cognition in Alzheimer's Disease Across Diverse Test Batteries and Datasets. Neuroepidemiology, 2014, 43, 194-205.	2.3	43
27	Genomewide Association Study of Alcohol Dependence Identifies Risk Loci Altering Ethanolâ€Response Behaviors in Model Organisms. Alcoholism: Clinical and Experimental Research, 2017, 41, 911-928.	2.4	43
28	Association of maternal and infant variants in <i>PNOC</i> and <i>COMT</i> genes with neonatal abstinence syndrome severity. American Journal on Addictions, 2017, 26, 42-49.	1.4	39
29	Population-Specific Risk of Type 2 Diabetes Conferred by HNF4A P2 Promoter Variants: A Lesson for Replication Studies. Diabetes, 2008, 57, 3161-3165.	0.6	37
30	Metaâ€analysis of genetic polymorphisms in granulomatosis with polyangiitis (Wegener's) reveals shared susceptibility loci with rheumatoid arthritis. Arthritis and Rheumatism, 2012, 64, 3463-3471.	6.7	33
31	Genomeâ€wide association study of rate of cognitive decline in Alzheimer's disease patients identifies novel genes and pathways. Alzheimer's and Dementia, 2020, 16, 1134-1145.	0.8	28
32	Shared genetic risk between eating disorder―and substanceâ€useâ€related phenotypes: Evidence from genomeâ€wide association studies. Addiction Biology, 2021, 26, e12880.	2.6	28
33	Power and Pitfalls of the Genome-Wide Association Study Approach to Identify Genes for Alzheimer's Disease. Current Psychiatry Reports, 2011, 13, 138-146.	4.5	27
34	Pharmacogenetic Effect of the Stromelysin (MMP3) Polymorphism on Stroke Risk in Relation to Antihypertensive Treatment. Stroke, 2011, 42, 330-335.	2.0	26
35	A Whole Genome Scan for Pulse Pressure/Stroke Volume Ratio in African Americans: The HyperGEN Study. American Journal of Hypertension, 2007, 20, 398-402.	2.0	25
36	Genetic modifiers of Hb E/ \hat{l}^2 O thalassemia identified by a two-stage genome-wide association study. BMC Medical Genetics, 2010, 11, 51.	2.1	25

#	Article	IF	Citations
37	Genomeâ€wide association metaâ€analysis of age at first cannabis use. Addiction, 2018, 113, 2073-2086.	3.3	24
38	Post-GWAS analysis of six substance use traits improves the identification and functional interpretation of genetic risk loci. Drug and Alcohol Dependence, 2020, 206, 107703.	3.2	19
39	Genome-Wide Meta-Analyses of FTND and TTFC Phenotypes. Nicotine and Tobacco Research, 2020, 22, 900-909.	2.6	17
40	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. Biological Psychiatry, 2015, 77, 870-879.	1.3	14
41	ACSL6 Is Associated with the Number of Cigarettes Smoked and Its Expression Is Altered by Chronic Nicotine Exposure. PLoS ONE, 2011, 6, e28790.	2.5	11
42	The executive prominent/memory prominent spectrum in Alzheimer's disease is highly heritable. Neurobiology of Aging, 2016, 41, 115-121.	3.1	11
43	Evidence for a quantitative trait locus affecting low levels of apolipoprotein B and low density lipoprotein on chromosome 10 in Caucasian families. Journal of Lipid Research, 2007, 48, 2632-2639.	4.2	9
44	A Whole-Genome Scan for Stroke or Myocardial Infarction in Family Blood Pressure Program Families. Stroke, 2008, 39, 1115-1120.	2.0	9
45	S100A10 identified in a genome-wide gene $ ilde{A}-$ cannabis dependence interaction analysis of risky sexual behaviours. Journal of Psychiatry and Neuroscience, 2017, 42, 252-261.	2.4	9
46	Genome-wide association study of phenotypes measuring progression from first cocaine or opioid use to dependence reveals novel risk genes. Exploration of Medicine, 2021, 2, 60-73.	1.5	6
47	Genome-wide association study of stimulant dependence. Translational Psychiatry, 2021, 11, 363.	4.8	4
48	Using linkage and association to identify and model genetic effects: summary of GAW15 Group 4. Genetic Epidemiology, 2007, 31, S34-S42.	1.3	3
49	Genome-Wide Studies in Sickle Cell Anemia Show Associations Between SNPs in the Olfactory Receptor Gene Cluster and Fetal Hemoglobin Concentration Blood, 2009, 114, 821-821.	1.4	2
50	No evidence for multiple loci affecting rheumatoid arthritis risk on chromosome 6p21. BMC Proceedings, 2007, 1, S42.	1.6	1
51	POLYGENIC RISK BURDEN AND CANNABIS USE COMORBIDITY IN PATIENTS WITH SCHIZOPHRENIA AND BIPOLAR DISORDER. European Neuropsychopharmacology, 2019, 29, S951.	0.7	1
52	P2-016: Identification of genetic variants associated with Alzheimer's disease: Progression rate., 2015, 11, P487-P487.		0
53	EXPLORING THE ROLE OF GENETIC REGULATION OF GENE EXPRESSION IN SUBSTANCE USE AND DEPENDENCE. European Neuropsychopharmacology, 2019, 29, S803-S804.	0.7	0
54	GENOME-WIDE ASSOCIATION STUDY OF COMORBID ALCOHOL DEPENDENCE AND MAJOR DEPRESSION. European Neuropsychopharmacology, 2019, 29, S971.	0.7	0

#	Article	IF	CITATIONS
55	Genome-wide association study of phenotypes measuring progression from first cocaine or opioid use to dependence reveals novel risk genes. Exploration of Medicine, 0, , .	1.5	O
56	P90. A Genetically Informed Examination of Posttraumatic Stress Disorder and Traumatic Brain Injury's Impact on Dementia Risk in US Veterans. Biological Psychiatry, 2022, 91, S123-S124.	1.3	0