# Sally I Sharp

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7833577/publications.pdf

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|---|---|---|---|
| 39 papers | 3,342 citations | 394421 19 h-index | 330143 37 g-index |
| 41 all docs | 41 docs citations | 41 times ranked | 8075 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | The UK10K project identifies rare variants in health and disease. Nature, 2015, 526, 82-90. | 27.8 | 1,014 |
| 2 | Genome-wide association study of more than 40,000 bipolar disorder cases provides new insights into the underlying biology. Nature Genetics, 2021, 53, 817-829. | 21.4 | 629 |
| 3 | Association of Plasma Clusterin Concentration With Severity, Pathology, and Progression in Alzheimer Disease. Archives of General Psychiatry, 2010, 67, 739. | 12.3 | 353 |
| 4 | Rare coding variants in ten genes confer substantial risk for schizophrenia. Nature, 2022, 604, 509-516. | 27.8 | 326 |
| 5 | Hypertension is a potential risk factor for vascular dementia: systematic review. International Journal of Geriatric Psychiatry, 2011, 26, 661-669. | 2.7 | 184 |
| 6 | Genetics of attention-deficit hyperactivity disorder (ADHD). Neuropharmacology, 2009, 57, 590-600. | 4.1 | 113 |
| 7 | Neuropsychiatric symptoms in dementia: Importance and treatment considerations. International Review of Psychiatry, 2008, 20, 396-404. | 2.8 | 89 |
| 8 | A rare variant in APOC3 is associated with plasma triglyceride and VLDL levels in Europeans. Nature Communications, 2014, 5, 4871. | 12.8 | 62 |
| 9 | Cortical Serotonin 1A Receptor Levels Are Associated with Depression in Patients with Dementia with Lewy Bodies and Parkinsonâ€™s Disease Dementia. Dementia and Geriatric Cognitive Disorders, 2008, 26, 330-338. | 1.5 | 48 |
| 10 | Association of rare variation in the glutamate receptor gene SLC1A2 with susceptibility to bipolar disorder and schizophrenia. European Journal of Human Genetics, 2015, 23, 1200-1206. | 2.8 | 45 |
| 11 | Genetic Association, Mutation Screening, and Functional Analysis of a Kozak Sequence Variant in the Metabotropic Glutamate Receptor 3 Gene in Bipolar Disorder. JAMA Psychiatry, 2013, 70, 591. | 11.0 | 43 |
| 12 | Genetic association of the tachykinin receptor 1 TACR1 gene in bipolar disorder, attention deficit hyperactivity disorder, and the alcohol dependence syndrome. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2014, 165, 373-380. | 1.7 | 39 |
| 13 | Choline Acetyltransferase Activity in Vascular Dementia and Stroke. Dementia and Geriatric Cognitive Disorders, 2009, 28, 233-238. | 1.5 | 36 |
| 14 | Aggressive Behavior and Neuroleptic Medication Are Associated With Increased Number of Alpha1-Adrenoceptors in Patients With Alzheimer Disease. American Journal of Geriatric Psychiatry, 2007, 15, 435-437. | 1.2 | 35 |
| 15 | The functional GRM3 Kozak sequence variant rs148754219 affects the risk of schizophrenia and alcohol dependence as well as bipolar disorder. Psychiatric Genetics, 2014, 24, 277-278. | 1.1 | 33 |
| 16 | Genetic variants in or nearADH1BandADH1Caffect susceptibility to alcohol dependence in a British and Irish population. Addiction Biology, 2015, 20, 594-604. | 2.6 | 33 |
| 17 | Psychiatric and behavioral symptoms in Alzheimerâ€™s disease and other dementias: Etiology and management. Current Neurology and Neuroscience Reports, 2005, 5, 345-354. | 4.2 | 23 |
| 18 | Investigation of the biological relevance ofHelicobacter pylori cagElocus diversity, presence of CagA tyrosine phosphorylation motifs and vacuolating cytotoxin genotype on IL-8 induction in gastric epithelial cells. FEMS Immunology and Medical Microbiology, 2003, 36, 135-140. | 2.7 | 20 |

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|---|---|---|---|
| 19 | Characterisation of age and polarity at onset in bipolar disorder. British Journal of Psychiatry, 2021, 219, 659-669. | 2.8 | 20 |
| 20 | Genetic Associations of Autopsy-Confirmed Vascular Dementia Subtypes. Dementia and Geriatric Cognitive Disorders, 2011, 31, 247-253. | 1.5 | 19 |
| 21 | Determining the Association of the 5HTTLPR Polymorphism with Delusions and Hallucinations in Lewy Body Dementias. American Journal of Geriatric Psychiatry, 2014, 22, 580-586. | 1.2 | 19 |
| 22 | Mutation intolerant genes and targets of FMRP are enriched for nonsynonymous alleles in schizophrenia. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2017, 174, 724-731. | 1.7 | 19 |
| 23 | A nonconservative amino acid change in the UPF3B gene in a patient with schizophrenia. Psychiatric Genetics, 2012, 22, 150-151. | 1.1 | 18 |
| 24 | Exome sequence analysis and follow up genotyping implicates rare ULK1 variants to be involved in susceptibility to schizophrenia. Annals of Human Genetics, 2018, 82, 88-92. | 0.8 | 16 |
| 25 | The KINGS Ins2+/G32S Mouse: A Novel Model of Î²-Cell Endoplasmic Reticulum Stress and Human Diabetes. Diabetes, 2020, 69, 2667-2677. | 0.6 | 16 |
| 26 | Identification ofcagATyrosine Phosphorylation DNA Motifs inHelicobacter pyloriIsolates from Peptic Ulcer Patients by Novel PCR-Restriction Fragment Length Polymorphism and Real-Time Fluorescence PCR Assays. Journal of Clinical Microbiology, 2003, 41, 3112-3118. | 3.9 | 14 |
| 27 | Sequencing of the ANKYRIN 3 gene (ANK3) encoding ankyrin G in bipolar disorder reveals a nonâ€conservative amino acid change in a short isoform of ankyrin G. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 328-335. | 1.7 | 14 |
| 28 | Genetic variation in the miRâ€708 gene and its binding targets in bipolar disorder. Bipolar Disorders, 2016, 18, 650-656. | 1.9 | 14 |
| 29 | Rare variant analysis in multiply affected families, association studies and functional analysis suggest a role for the ITGÎ²4 gene in schizophrenia and bipolar disorder. Schizophrenia Research, 2018, 199, 181-188. | 2.0 | 11 |
| 30 | The effect of clozapine on mRNA expression for genes encoding G protein-coupled receptors and the protein components of clathrin-mediated endocytosis. Psychiatric Genetics, 2013, 23, 153-162. | 1.1 | 10 |
| 31 | No association of COMT val158met polymorphism and psychotic symptoms in Lewy body dementias. Neuroscience Letters, 2012, 531, 1-4. | 2.1 | 7 |
| 32 | Lack of allelic association between markers at the DRD2 and ANKK1 gene loci with the alcohol-dependence syndrome and criminal activity. Psychiatric Genetics, 2011, 21, 323-324. | 1.1 | 6 |
| 33 | Identification of rare nonsynonymous variants in SYNE1/CPG2 in bipolar affective disorder. Psychiatric Genetics, 2017, 27, 81-88. | 1.1 | 6 |
| 34 | Evidence for genetic susceptibility to the alcohol dependence syndrome from the thiamine transporter 2 gene solute carrier SLC19A3. Psychiatric Genetics, 2014, 24, 122-123. | 1.1 | 2 |
| 35 | Genetic variant analysis of the putative regulatory regions of the LRRC7 gene in bipolar disorder. Psychiatric Genetics, 2016, 26, 99-100. | 1.1 | 2 |
| 36 | Genetic association and functional characterization of MCPH1 gene variation in bipolar disorder and schizophrenia. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, 258-265. | 1.7 | 2 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | Neurochemistry of severe dementia. Reviews in Clinical Gerontology, 2005, 15, 105-123. | 0.5 | 1 |
| 38 | F130EXOME SEQUENCE ANALYSIS IDENTIFY RARE GENETIC VARIANT IMPLICATED IN SUSCEPTIBILITY TO SCHIZOPHRENIA. European Neuropsychopharmacology, 2019, 29, S1181. | 0.7 | 0 |
| 39 | GENETIC ASSOCIATION AND FUNCTIONAL CHARACTERIZATION OF A VARIANT IN THE MCPH1 GENE IN BIPOLAR DISORDER AND SCHIZOPHRENIA. European Neuropsychopharmacology, 2019, 29, S966-S967. | 0.7 | 0 |