

Aiden P Corvin

List of Publications by Citations

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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.

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|--------------------|--------------------------|-----------------|-----------------|
| 187 papers | 20,752 citations | 56 h-index | 143 g-index |
| 202 ext. papers | 25,973 ext. citations | 10.2 avg, IF | 5.24 L-index |

| # | Paper | IF | Citations |
|-----|--|------|-----------|
| 187 | Genetic risk and a primary role for cell-mediated immune mechanisms in multiple sclerosis. <i>Nature</i> , 2011 , 476, 214-9 | 50.4 | 1948 |
| 186 | Genetic relationship between five psychiatric disorders estimated from genome-wide SNPs. <i>Nature Genetics</i> , 2013 , 45, 984-94 | 36.3 | 1628 |
| 185 | Genome-wide association analysis identifies 13 new risk loci for schizophrenia. <i>Nature Genetics</i> , 2013 , 45, 1150-9 | 36.3 | 1153 |
| 184 | Collaborative genome-wide association analysis supports a role for ANK3 and CACNA1C in bipolar disorder. <i>Nature Genetics</i> , 2008 , 40, 1056-8 | 36.3 | 949 |
| 183 | Identification of loci associated with schizophrenia by genome-wide association and follow-up. <i>Nature Genetics</i> , 2008 , 40, 1053-5 | 36.3 | 877 |
| 182 | A genome-wide association study identifies new psoriasis susceptibility loci and an interaction between HLA-C and ERAP1. <i>Nature Genetics</i> , 2010 , 42, 985-90 | 36.3 | 773 |
| 181 | Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018 , 360, | 33.3 | 666 |
| 180 | Genome-wide association study identifies 30 loci associated with bipolar disorder. <i>Nature Genetics</i> , 2019 , 51, 793-803 | 36.3 | 662 |
| 179 | Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. <i>American Journal of Human Genetics</i> , 2015 , 97, 576-92 | 11 | 649 |
| 178 | Interaction between ERAP1 and HLA-B27 in ankylosing spondylitis implicates peptide handling in the mechanism for HLA-B27 in disease susceptibility. <i>Nature Genetics</i> , 2011 , 43, 761-7 | 36.3 | 646 |
| 177 | Common genetic variants influence human subcortical brain structures. <i>Nature</i> , 2015 , 520, 224-9 | 50.4 | 601 |
| 176 | Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. <i>Nature Genetics</i> , 2017 , 49, 27-35 | 36.3 | 530 |
| 175 | Identification of common variants associated with human hippocampal and intracranial volumes. <i>Nature Genetics</i> , 2012 , 44, 552-61 | 36.3 | 498 |
| 174 | Genome-wide association meta-analysis in 269,867 individuals identifies new genetic and functional links to intelligence. <i>Nature Genetics</i> , 2018 , 50, 912-919 | 36.3 | 475 |
| 173 | Partitioning heritability of regulatory and cell-type-specific variants across 11 common diseases. <i>American Journal of Human Genetics</i> , 2014 , 95, 535-52 | 11 | 411 |
| 172 | Genome-wide association study of ulcerative colitis identifies three new susceptibility loci, including the HNF4A region. <i>Nature Genetics</i> , 2009 , 41, 1330-4 | 36.3 | 411 |
| 171 | Cortical Brain Abnormalities in 4474 Individuals With Schizophrenia and 5098 Control Subjects via the Enhancing Neuro Imaging Genetics Through Meta Analysis (ENIGMA) Consortium. <i>Biological Psychiatry</i> , 2018 , 84, 644-654 | 7.9 | 325 |

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| 170 | Common variants near ATM are associated with glycemic response to metformin in type 2 diabetes. <i>Nature Genetics</i> , 2011 , 43, 117-20 | 36.3 | 319 |
| 169 | Genome-wide association study identifies a variant in HDAC9 associated with large vessel ischemic stroke. <i>Nature Genetics</i> , 2012 , 44, 328-33 | 36.3 | 314 |
| 168 | Polygenic dissection of diagnosis and clinical dimensions of bipolar disorder and schizophrenia. <i>Molecular Psychiatry</i> , 2014 , 19, 1017-1024 | 15.1 | 258 |
| 167 | Duplications of the neuropeptide receptor gene VIPR2 confer significant risk for schizophrenia. <i>Nature</i> , 2011 , 471, 499-503 | 50.4 | 257 |
| 166 | Study of 300,486 individuals identifies 148 independent genetic loci influencing general cognitive function. <i>Nature Communications</i> , 2018 , 9, 2098 | 17.4 | 254 |
| 165 | High frequencies of de novo CNVs in bipolar disorder and schizophrenia. <i>Neuron</i> , 2011 , 72, 951-63 | 13.9 | 240 |
| 164 | Dissection of the genetics of Parkinson's disease identifies an additional association 5' of SNCA and multiple associated haplotypes at 17q21. <i>Human Molecular Genetics</i> , 2011 , 20, 345-53 | 5.6 | 178 |
| 163 | Novel genetic loci associated with hippocampal volume. <i>Nature Communications</i> , 2017 , 8, 13624 | 17.4 | 173 |
| 162 | Genomewide linkage scan in schizoaffective disorder: significant evidence for linkage at 1q42 close to DISC1, and suggestive evidence at 22q11 and 19p13. <i>Archives of General Psychiatry</i> , 2005 , 62, 1081-8 | | 164 |
| 161 | Joint analysis of psychiatric disorders increases accuracy of risk prediction for schizophrenia, bipolar disorder, and major depressive disorder. <i>American Journal of Human Genetics</i> , 2015 , 96, 283-94 | 11 | 161 |
| 160 | The genetic architecture of the human cerebral cortex. <i>Science</i> , 2020 , 367, | 33.3 | 156 |
| 159 | Novel genetic loci underlying human intracranial volume identified through genome-wide association. <i>Nature Neuroscience</i> , 2016 , 19, 1569-1582 | 25.5 | 147 |
| 158 | Evidence that interaction between neuregulin 1 and its receptor erbB4 increases susceptibility to schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2006 , 141B, 96-103 | 3.5 | 143 |
| 157 | Common variants at the MHC locus and at chromosome 16q24.1 predispose to Barrett's esophagus. <i>Nature Genetics</i> , 2012 , 44, 1131-6 | 36.3 | 139 |
| 156 | Familial patterns and the origins of individual differences in synaesthesia. <i>Cognition</i> , 2008 , 106, 871-93 | 3.5 | 123 |
| 155 | Psychosis susceptibility gene ZNF804A and cognitive performance in schizophrenia. <i>Archives of General Psychiatry</i> , 2010 , 67, 692-700 | | 120 |
| 154 | The SNP ratio test: pathway analysis of genome-wide association datasets. <i>Bioinformatics</i> , 2009 , 25, 2762-3 | 2.3 | 116 |
| 153 | A comprehensive family-based replication study of schizophrenia genes. <i>JAMA Psychiatry</i> , 2013 , 70, 573-81 | 1.5 | 115 |

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| 152 | Confirming RGS4 as a susceptibility gene for schizophrenia. <i>American Journal of Medical Genetics Part A</i> , 2004 , 125B, 50-3 | | 114 |
| 151 | Genome-wide comparative analysis of atopic dermatitis and psoriasis gives insight into opposing genetic mechanisms. <i>American Journal of Human Genetics</i> , 2015 , 96, 104-20 | 11 | 113 |
| 150 | GWAS of Suicide Attempt in Psychiatric Disorders and Association With Major Depression Polygenic Risk Scores. <i>American Journal of Psychiatry</i> , 2019 , 176, 651-660 | 11.9 | 103 |
| 149 | Copy-number variants in neurodevelopmental disorders: promises and challenges. <i>Trends in Genetics</i> , 2009 , 25, 536-44 | 8.5 | 102 |
| 148 | Variance in neurocognitive performance is associated with dysbindin-1 in schizophrenia: a preliminary study. <i>Neuropsychologia</i> , 2007 , 45, 454-8 | 3.2 | 101 |
| 147 | Genome-wide association studies: findings at the major histocompatibility complex locus in psychosis. <i>Biological Psychiatry</i> , 2014 , 75, 276-83 | 7.9 | 95 |
| 146 | Development of strategies for SNP detection in RNA-seq data: application to lymphoblastoid cell lines and evaluation using 1000 Genomes data. <i>PLoS ONE</i> , 2013 , 8, e58815 | 3.7 | 94 |
| 145 | Human subcortical brain asymmetries in 15,847 people worldwide reveal effects of age and sex. <i>Brain Imaging and Behavior</i> , 2017 , 11, 1497-1514 | 4.1 | 87 |
| 144 | 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 |
| 143 | Evaluation of a susceptibility gene for schizophrenia: genotype based meta-analysis of RGS4 polymorphisms from thirteen independent samples. <i>Biological Psychiatry</i> , 2006 , 60, 152-62 | 7.9 | 80 |
| 142 | Fluorescent nanodiamond tracking reveals intraneuronal transport abnormalities induced by brain-disease-related genetic risk factors. <i>Nature Nanotechnology</i> , 2017 , 12, 322-328 | 28.7 | 79 |
| 141 | Cigarette smoking and psychotic symptoms in bipolar affective disorder. <i>British Journal of Psychiatry</i> , 2001 , 179, 35-8 | 5.4 | 77 |
| 140 | Common variants in the HLA-DRB1-HLA-DQA1 HLA class II region are associated with susceptibility to visceral leishmaniasis. <i>Nature Genetics</i> , 2013 , 45, 208-13 | 36.3 | 76 |
| 139 | ZNF804A risk allele is associated with relatively intact gray matter volume in patients with schizophrenia. <i>NeuroImage</i> , 2011 , 54, 2132-7 | 7.9 | 74 |
| 138 | No evidence for association of the dysbindin gene [DTNBP1] with schizophrenia in an Irish population-based study. <i>Schizophrenia Research</i> , 2003 , 60, 167-72 | 3.6 | 74 |
| 137 | Large-Scale Cognitive GWAS Meta-Analysis Reveals Tissue-Specific Neural Expression and Potential Nootropic Drug Targets. <i>Cell Reports</i> , 2017 , 21, 2597-2613 | 10.6 | 71 |
| 136 | Cortical patterning of abnormal morphometric similarity in psychosis is associated with brain expression of schizophrenia-related genes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 9604-9609 | 11.5 | 70 |
| 135 | Occurrence and co-occurrence of hallucinations by modality in schizophrenia-spectrum disorders. <i>Psychiatry Research</i> , 2017 , 252, 154-160 | 9.9 | 62 |

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| 134 | CNV analysis in a large schizophrenia sample implicates deletions at 16p12.1 and SLC1A1 and duplications at 1p36.33 and CGNL1. <i>Human Molecular Genetics</i> , 2014 , 23, 1669-76 | 5.6 | 61 |
| 133 | Early visual processing deficits in dysbindin-associated schizophrenia. <i>Biological Psychiatry</i> , 2008 , 63, 484-9 | 7.9 | 59 |
| 132 | Mental state decoding v. mental state reasoning as a mediator between cognitive and social function in psychosis. <i>British Journal of Psychiatry</i> , 2008 , 193, 77-8 | 5.4 | 58 |
| 131 | Implication of a rare deletion at distal 16p11.2 in schizophrenia. <i>JAMA Psychiatry</i> , 2013 , 70, 253-60 | 14.5 | 56 |
| 130 | Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. <i>American Journal of Human Genetics</i> , 2018 , 102, 1185-1194 | 11 | 55 |
| 129 | The correlation between reading and mathematics ability at age twelve has a substantial genetic component. <i>Nature Communications</i> , 2014 , 5, 4204 | 17.4 | 54 |
| 128 | Neurocognition and suicidal behaviour in an Irish population with major psychotic disorders. <i>Schizophrenia Research</i> , 2006 , 85, 196-200 | 3.6 | 53 |
| 127 | The role of the major histocompatibility complex region in cognition and brain structure: a schizophrenia GWAS follow-up. <i>American Journal of Psychiatry</i> , 2013 , 170, 877-85 | 11.9 | 51 |
| 126 | Dysbindin (DTNBP1) and the biogenesis of lysosome-related organelles complex 1 (BLOC-1): main and epistatic gene effects are potential contributors to schizophrenia susceptibility. <i>Biological Psychiatry</i> , 2008 , 63, 24-31 | 7.9 | 51 |
| 125 | Genetic modifiers and subtypes in schizophrenia: investigations of age at onset, severity, sex and family history. <i>Schizophrenia Research</i> , 2014 , 154, 48-53 | 3.6 | 49 |
| 124 | The psychosis susceptibility gene ZNF804A: associations, functions, and phenotypes. <i>Schizophrenia Bulletin</i> , 2010 , 36, 904-9 | 1.3 | 44 |
| 123 | New data and an old puzzle: the negative association between schizophrenia and rheumatoid arthritis. <i>International Journal of Epidemiology</i> , 2015 , 44, 1706-21 | 7.8 | 43 |
| 122 | Insulin-like growth factor 1 (IGF1) and its active peptide (1-3)IGF1 enhance the expression of synaptic markers in neuronal circuits through different cellular mechanisms. <i>Neuroscience Letters</i> , 2012 , 520, 51-6 | 3.3 | 42 |
| 121 | Influence of NOS1 on verbal intelligence and working memory in both patients with schizophrenia and healthy control subjects. <i>Archives of General Psychiatry</i> , 2009 , 66, 1045-54 | | 42 |
| 120 | Evidence that specific executive functions predict symptom variance among schizophrenia patients with a predominantly negative symptom profile. <i>Cognitive Neuropsychiatry</i> , 2006 , 11, 13-32 | 2 | 41 |
| 119 | The Relationship Between Polygenic Risk Scores and Cognition in Schizophrenia. <i>Schizophrenia Bulletin</i> , 2020 , 46, 336-344 | 1.3 | 38 |
| 118 | Functional genomics and schizophrenia: endophenotypes and mutant models. <i>Psychiatric Clinics of North America</i> , 2007 , 30, 365-99 | 3.1 | 38 |
| 117 | Pleiotropic Meta-Analysis of Cognition, Education, and Schizophrenia Differentiates Roles of Early Neurodevelopmental and Adult Synaptic Pathways. <i>American Journal of Human Genetics</i> , 2019 , 105, 334-350 | 11 | 37 |

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| 116 | Social cognition in bipolar disorder versus schizophrenia: comparability in mental state decoding deficits. <i>Bipolar Disorders</i> , 2012 , 14, 743-8 | 3.8 | 37 |
| 115 | Are the cognitive deficits associated with impaired insight in schizophrenia specific to executive task performance?. <i>Journal of Nervous and Mental Disease</i> , 2005 , 193, 803-8 | 1.8 | 37 |
| 114 | A genome-wide association analysis of a broad psychosis phenotype identifies three loci for further investigation. <i>Biological Psychiatry</i> , 2014 , 75, 386-97 | 7.9 | 36 |
| 113 | Effects of MIR137 on fronto-amygdala functional connectivity. <i>NeuroImage</i> , 2014 , 90, 189-95 | 7.9 | 36 |
| 112 | Population structure and genome-wide patterns of variation in Ireland and Britain. <i>European Journal of Human Genetics</i> , 2010 , 18, 1248-54 | 5.3 | 36 |
| 111 | Genetics of Schizophrenia: Ready to Translate?. <i>Current Psychiatry Reports</i> , 2017 , 19, 61 | 9.1 | 36 |
| 110 | Multiplex target enrichment using DNA indexing for ultra-high throughput SNP detection. <i>DNA Research</i> , 2011 , 18, 31-8 | 4.5 | 35 |
| 109 | Mapping genomic loci implicates genes and synaptic biology in schizophrenia.. <i>Nature</i> , 2022 , | 50.4 | 35 |
| 108 | Data science for mental health: a UK perspective on a global challenge. <i>Lancet Psychiatry</i> , 2016 , 3, 993-998 | 23.3 | 34 |
| 107 | An inherited duplication at the gene p21 Protein-Activated Kinase 7 (PAK7) is a risk factor for psychosis. <i>Human Molecular Genetics</i> , 2014 , 23, 3316-26 | 5.6 | 32 |
| 106 | Mutation of Semaphorin-6A disrupts limbic and cortical connectivity and models neurodevelopmental psychopathology. <i>PLoS ONE</i> , 2011 , 6, e26488 | 3.7 | 32 |
| 105 | Detecting schizophrenia at the level of the individual: relative diagnostic value of whole-brain images, connectome-wide functional connectivity and graph-based metrics. <i>Psychological Medicine</i> , 2020 , 50, 1852-1861 | 6.9 | 32 |
| 104 | Are relational style and neuropsychological performance predictors of social attributions in chronic schizophrenia?. <i>Psychiatry Research</i> , 2008 , 161, 19-27 | 9.9 | 31 |
| 103 | Polymorphism in a lincRNA Associates with a Doubled Risk of Pneumococcal Bacteremia in Kenyan Children. <i>American Journal of Human Genetics</i> , 2016 , 98, 1092-1100 | 11 | 30 |
| 102 | A dysbindin risk haplotype associated with less severe manic-type symptoms in psychosis. <i>Neuroscience Letters</i> , 2008 , 431, 146-9 | 3.3 | 29 |
| 101 | Genome-wide association study identifies 30 Loci Associated with Bipolar Disorder | | 28 |
| 100 | Is "clinical" insight the same as "cognitive" insight in schizophrenia?. <i>Journal of the International Neuropsychological Society</i> , 2009 , 15, 471-5 | 3.1 | 27 |
| 99 | Neural effects of the CSMD1 genome-wide associated schizophrenia risk variant rs10503253. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2013 , 162B, 530-7 | 3.5 | 26 |

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|----|---|------|----|
| 98 | Evidence for cis-acting regulation of ANK3 and CACNA1C gene expression. <i>Bipolar Disorders</i> , 2010 , 12, 440-5 | 3.8 | 26 |
| 97 | The effect of the neurogranin schizophrenia risk variant rs12807809 on brain structure and function. <i>Twin Research and Human Genetics</i> , 2012 , 15, 296-303 | 2.2 | 26 |
| 96 | Cognitive analysis of schizophrenia risk genes that function as epigenetic regulators of gene expression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2016 , 171, 1170-1179 | 3.5 | 25 |
| 95 | Effects of a novel schizophrenia risk variant rs7914558 at CNM2 on brain structure and attributional style. <i>British Journal of Psychiatry</i> , 2014 , 204, 115-21 | 5.4 | 25 |
| 94 | Reduced occipital and prefrontal brain volumes in dysbindin-associated schizophrenia. <i>Neuropsychopharmacology</i> , 2010 , 35, 368-73 | 8.7 | 25 |
| 93 | Chitinase-3-like 1 (CHI3L1) gene and schizophrenia: genetic association and a potential functional mechanism. <i>Biological Psychiatry</i> , 2008 , 64, 98-103 | 7.9 | 25 |
| 92 | D-amino acid oxidase (DAO) genotype and mood symptomatology in schizophrenia. <i>Neuroscience Letters</i> , 2007 , 426, 97-100 | 3.3 | 25 |
| 91 | Variability in working memory performance explained by epistasis vs polygenic scores in the ZNF804A pathway. <i>JAMA Psychiatry</i> , 2014 , 71, 778-785 | 14.5 | 24 |
| 90 | Altered medial prefrontal activity during dynamic face processing in schizophrenia spectrum patients. <i>Schizophrenia Research</i> , 2014 , 157, 225-30 | 3.6 | 24 |
| 89 | Repeated insulin-like growth factor 1 treatment in a patient with rett syndrome: a single case study. <i>Frontiers in Pediatrics</i> , 2014 , 2, 52 | 3.4 | 24 |
| 88 | Genome-wide association study of intraocular pressure identifies the GLCCI1/ICA1 region as a glaucoma susceptibility locus. <i>Human Molecular Genetics</i> , 2013 , 22, 4653-60 | 5.6 | 24 |
| 87 | Genetic differences between five European populations. <i>Human Heredity</i> , 2010 , 70, 141-9 | 1.1 | 24 |
| 86 | Does the ability to sustain attention underlie symptom severity in schizophrenia?. <i>Schizophrenia Research</i> , 2009 , 107, 319-23 | 3.6 | 24 |
| 85 | Association of Copy Number Variation of the 15q11.2 BP1-BP2 Region With Cortical and Subcortical Morphology and Cognition. <i>JAMA Psychiatry</i> , 2020 , 77, 420-430 | 14.5 | 24 |
| 84 | Dose response of the 16p11.2 distal copy number variant on intracranial volume and basal ganglia. <i>Molecular Psychiatry</i> , 2020 , 25, 584-602 | 15.1 | 24 |
| 83 | Genome-wide association study reveals greater polygenic loading for schizophrenia in cases with a family history of illness. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2016 , 171B, 276-89 | 3.5 | 23 |
| 82 | Neuronal cell adhesion genes: Key players in risk for schizophrenia, bipolar disorder and other neurodevelopmental brain disorders?. <i>Cell Adhesion and Migration</i> , 2010 , 4, 511-4 | 3.2 | 23 |
| 81 | Integrating machine learning and multimodal neuroimaging to detect schizophrenia at the level of the individual. <i>Human Brain Mapping</i> , 2020 , 41, 1119-1135 | 5.9 | 23 |

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|----|--|-----|----|
| 80 | The NOS1 variant rs6490121 is associated with variation in prefrontal function and grey matter density in healthy individuals. <i>NeuroImage</i> , 2012 , 60, 614-22 | 7.9 | 22 |
| 79 | Are deficits in executive sub-processes simply reflecting more general cognitive decline in schizophrenia?. <i>Schizophrenia Research</i> , 2006 , 85, 168-73 | 3.6 | 22 |
| 78 | Independent evidence for an association between general cognitive ability and a genetic locus for educational attainment. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2015 , 168B, 363-73 | 3.5 | 21 |
| 77 | Social dysfunction in schizophrenia: an investigation of the GAF scale's sensitivity to deficits in social cognition. <i>Schizophrenia Research</i> , 2013 , 146, 363-5 | 3.6 | 21 |
| 76 | Targeted Sequencing of 10,198 Samples Confirms Abnormalities in Neuronal Activity and Implicates Voltage-Gated Sodium Channels in Schizophrenia Pathogenesis. <i>Biological Psychiatry</i> , 2019 , 85, 554-562 | 7.9 | 21 |
| 75 | A neuropsychological investigation of the genome wide associated schizophrenia risk variant NRG1 rs12807809. <i>Schizophrenia Research</i> , 2011 , 125, 304-6 | 3.6 | 20 |
| 74 | Mood-incongruent psychosis in bipolar disorder: conditional linkage analysis shows genome-wide suggestive linkage at 1q32.3, 7p13 and 20q13.31. <i>Bipolar Disorders</i> , 2009 , 11, 610-20 | 3.8 | 20 |
| 73 | A randomized controlled trial of cognitive remediation for a national cohort of forensic patients with schizophrenia or schizoaffective disorder. <i>BMC Psychiatry</i> , 2019 , 19, 27 | 4.2 | 20 |
| 72 | What Next in Schizophrenia Genetics for the Psychiatric Genomics Consortium?. <i>Schizophrenia Bulletin</i> , 2016 , 42, 538-41 | 1.3 | 19 |
| 71 | Genes predict village of origin in rural Europe. <i>European Journal of Human Genetics</i> , 2010 , 18, 1269-70 | 5.3 | 19 |
| 70 | Common polygenic variation in coeliac disease and confirmation of ZNF335 and NIFA as disease susceptibility loci. <i>European Journal of Human Genetics</i> , 2016 , 24, 291-7 | 5.3 | 18 |
| 69 | Replicated genetic evidence supports a role for HOMER2 in schizophrenia. <i>Neuroscience Letters</i> , 2010 , 468, 229-33 | 3.3 | 18 |
| 68 | A NOS1 variant implicated in cognitive performance influences evoked neural responses during a high density EEG study of early visual perception. <i>Human Brain Mapping</i> , 2012 , 33, 1202-11 | 5.9 | 17 |
| 67 | Cognitive Characterization of Schizophrenia Risk Variants Involved in Synaptic Transmission: Evidence of CACNA1C's Role in Working Memory. <i>Neuropsychopharmacology</i> , 2017 , 42, 2612-2622 | 8.7 | 16 |
| 66 | MIR137HG risk variant rs1625579 genotype is related to corpus callosum volume in schizophrenia. <i>Neuroscience Letters</i> , 2015 , 602, 44-9 | 3.3 | 16 |
| 65 | The phenotypic manifestations of rare CNVs in schizophrenia. <i>Schizophrenia Research</i> , 2014 , 158, 255-60 | 3.6 | 16 |
| 64 | Genome-wide schizophrenia variant at MIR137 does not impact white matter microstructure in healthy participants. <i>Neuroscience Letters</i> , 2014 , 574, 6-10 | 3.3 | 15 |
| 63 | Analysis of the hexanucleotide repeat expansion and founder haplotype at C9ORF72 in an Irish psychosis case-control sample. <i>Neurobiology of Aging</i> , 2014 , 35, 1510.e1-5 | 5.6 | 15 |

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|----|---|-----|----|
| 62 | The miR-137 schizophrenia susceptibility variant rs1625579 does not predict variability in brain volume in a sample of schizophrenic patients and healthy individuals. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2014 , 165B, 467-71 | 3.5 | 15 |
| 61 | The shock of the new: progress in schizophrenia genomics. <i>Current Genomics</i> , 2011 , 12, 516-24 | 2.6 | 15 |
| 60 | DNA methylation meta-analysis reveals cellular alterations in psychosis and markers of treatment-resistant schizophrenia. <i>ELife</i> , 2021 , 10, | 8.9 | 15 |
| 59 | Further evidence of alerted default network connectivity and association with theory of mind ability in schizophrenia. <i>Schizophrenia Research</i> , 2017 , 184, 52-58 | 3.6 | 14 |
| 58 | Neuroharmony: A new tool for harmonizing volumetric MRI data from unseen scanners. <i>NeuroImage</i> , 2020 , 220, 117127 | 7.9 | 14 |
| 57 | Genetic classification of populations using supervised learning. <i>PLoS ONE</i> , 2011 , 6, e14802 | 3.7 | 14 |
| 56 | Do antisaccade deficits in schizophrenia provide evidence of a specific inhibitory function?. <i>Journal of the International Neuropsychological Society</i> , 2006 , 12, 901-6 | 3.1 | 14 |
| 55 | The Genetics of Endophenotypes of Neurofunction to Understand Schizophrenia (GENUS) consortium: A collaborative cognitive and neuroimaging genetics project. <i>Schizophrenia Research</i> , 2018 , 195, 306-317 | 3.6 | 14 |
| 54 | No evidence that runs of homozygosity are associated with schizophrenia in an Irish genome-wide association dataset. <i>Schizophrenia Research</i> , 2014 , 154, 79-82 | 3.6 | 13 |
| 53 | Unlocking the treasure trove: from genes to schizophrenia biology. <i>Schizophrenia Bulletin</i> , 2014 , 40, 492-63 | 6.3 | 13 |
| 52 | Schizophrenia at a genetics crossroads: where to now?. <i>Schizophrenia Bulletin</i> , 2013 , 39, 490-5 | 1.3 | 12 |
| 51 | Functional investigation of a schizophrenia GWAS signal at the CDC42 gene. <i>World Journal of Biological Psychiatry</i> , 2012 , 13, 550-4 | 3.8 | 12 |
| 50 | Investigation of the apolipoprotein-L (APOL) gene family and schizophrenia using a novel DNA pooling strategy for public database SNPs. <i>Schizophrenia Research</i> , 2005 , 76, 231-8 | 3.6 | 12 |
| 49 | Genome-wide association study of over 40,000 bipolar disorder cases provides new insights into the underlying biology | | 11 |
| 48 | Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. <i>Biological Psychiatry</i> , 2022 , 91, 102-117 | 7.9 | 11 |
| 47 | Dissecting the Shared Genetic Architecture of Suicide Attempt, Psychiatric Disorders, and Known Risk Factors. <i>Biological Psychiatry</i> , 2021 , | 7.9 | 11 |
| 46 | Mosaic copy number variation in schizophrenia. <i>European Journal of Human Genetics</i> , 2013 , 21, 1007-11 | 5.3 | 10 |
| 45 | Two patients walk into a clinic...a genomics perspective on the future of schizophrenia. <i>BMC Biology</i> , 2011 , 9, 77 | 7.3 | 10 |

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|----|--|-----|----|
| 44 | Expression of nuclear Methyl-CpG binding protein 2 (Mecp2) is dependent on neuronal stimulation and application of Insulin-like growth factor 1. <i>Neuroscience Letters</i> , 2016 , 621, 111-116 | 3.3 | 10 |
| 43 | Childhood trauma, parental bonding, and social cognition in patients with schizophrenia and healthy adults. <i>Journal of Clinical Psychology</i> , 2021 , 77, 241-253 | 2.8 | 10 |
| 42 | GWAS meta-analysis (N=279,930) identifies new genes and functional links to intelligence | | 9 |
| 41 | Prevalence of N-Methyl-d-Aspartate Receptor antibody (NMDAR-Ab) encephalitis in patients with first episode psychosis and treatment resistant schizophrenia on clozapine, a population based study. <i>Schizophrenia Research</i> , 2020 , 222, 455-461 | 3.6 | 8 |
| 40 | Beyond C4: Analysis of the complement gene pathway shows enrichment for IQ in patients with psychotic disorders and healthy controls. <i>Genes, Brain and Behavior</i> , 2019 , 18, e12602 | 3.6 | 8 |
| 39 | Identifying schizophrenia patients who carry pathogenic genetic copy number variants using standard clinical assessment: retrospective cohort study. <i>British Journal of Psychiatry</i> , 2020 , 216, 275-279 | 5.4 | 7 |
| 38 | A meta-analysis of deep brain structural shape and asymmetry abnormalities in 2,833 individuals with schizophrenia compared with 3,929 healthy volunteers via the ENIGMA Consortium. <i>Human Brain Mapping</i> , 2021 , | 5.9 | 7 |
| 37 | Effects of MiR-137 genetic risk score on brain volume and cortical measures in patients with schizophrenia and controls. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2018 , 177, 369-376 | 3.5 | 6 |
| 36 | Variance in facial recognition performance associated with BDNF in schizophrenia. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2007 , 144B, 578-9 | 3.5 | 6 |
| 35 | The attitudes of Irish trainees to their training and its supervision: a five-year follow up study. <i>Irish Journal of Psychological Medicine</i> , 2001 , 18, 120-125 | 3 | 6 |
| 34 | Methyl-CpG-binding protein 2 mediates overlapping mechanisms across brain disorders. <i>Scientific Reports</i> , 2020 , 10, 22255 | 4.9 | 5 |
| 33 | Childhood trauma, brain structure and emotion recognition in patients with schizophrenia and healthy participants. <i>Social Cognitive and Affective Neuroscience</i> , 2020 , 15, 1336-1350 | 4 | 5 |
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| 13 | Proof of concept: Molecular prediction of schizophrenia risk | | 1 |
| 12 | Large-scale analysis of DNA methylation identifies cellular alterations in blood from psychosis patients and molecular biomarkers of treatment-resistant schizophrenia | | 1 |
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| 1 | Schizophrenia genomics 2022 , 17-41 | | |