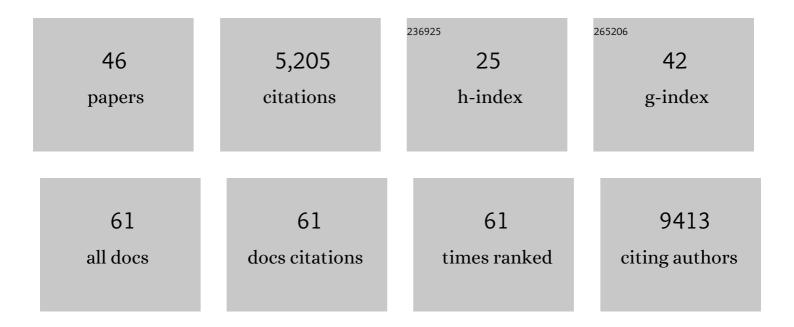
## John F Fullard

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Sex Differences in the Human Brain Transcriptome of Cases With Schizophrenia. Biological Psychiatry, 2022, 91, 92-101.	1.3	38
2	Multi-ancestry eQTL meta-analysis of human brain identifies candidate causal variants for brain-related traits. Nature Genetics, 2022, 54, 161-169.	21.4	49
3	A bidirectional competitive interaction between circHomer1 and Homer1b within the orbitofrontal cortex regulates reversal learning. Cell Reports, 2022, 38, 110282.	6.4	17
4	Non-cell-autonomous disruption of nuclear architecture as a potential cause of COVID-19-induced anosmia. Cell, 2022, 185, 1052-1064.e12.	28.9	154
5	Chromatin domain alterations linked to 3D genome organization in a large cohort of schizophrenia and bipolar disorder brains. Nature Neuroscience, 2022, 25, 474-483.	14.8	25
6	Common variants contribute to intrinsic human brain functional networks. Nature Genetics, 2022, 54, 508-517.	21.4	37
7	A complete temporal transcription factor series in the fly visual system. Nature, 2022, 604, 316-322.	27.8	60
8	The Neuroepigenome: Implications of Chemical and Physical Modifications of Genomic DNA in Schizophrenia. Biological Psychiatry, 2022, 92, 443-449.	1.3	6
9	Impact of schizophrenia GWAS loci converge onto distinct pathways in cortical interneurons vs glutamatergic neurons during development. Molecular Psychiatry, 2022, 27, 4218-4233.	7.9	6
10	ATAC-seq and psychiatric disorders. , 2021, , 143-162.		0
11	Integration of Alzheimer's disease genetics and myeloid genomics identifies disease risk regulatory elements and genes. Nature Communications, 2021, 12, 1610.	12.8	118
12	Common genetic variation influencing human white matter microstructure. Science, 2021, 372, .	12.6	106
13	Single-nucleus transcriptome analysis of human brain immune response in patients with severe COVID-19. Genome Medicine, 2021, 13, 118.	8.2	81
14	Unbiased identification of novel transcription factors in striatal compartmentation and striosome maturation. ELife, 2021, 10, .	6.0	9
15	Large-Scale Integrative Brain Transcriptome and Epigenome Imputation for PTSD Identifies Implicated Genes and Pathways. Biological Psychiatry, 2020, 87, S52.	1.3	0
16	Dissecting Hes-centred transcriptional networks in neural stem cell maintenance and tumorigenesis in <i>Drosophila</i> . Development (Cambridge), 2020, 147, .	2.5	9
17	Chromatin accessibility mapping of the striatum identifies tyrosine kinase FYN as a therapeutic target for heroin use disorder. Nature Communications, 2020, 11, 4634.	12.8	21
18	Genetic studies of Alzheimer's disease risk implicate clearance of lipid rich debris in myeloid cells. Alzheimer's and Dementia, 2020, 16, e040601.	0.8	0

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19	Integration of Alzheimer's disease genetics and myeloid genomics reveals novel disease risk mechanisms. Alzheimer's and Dementia, 2020, 16, e043897.	0.8	0
20	Common schizophrenia risk variants are enriched in open chromatin regions of human glutamatergic neurons. Nature Communications, 2020, 11, 5581.	12.8	53
21	Differential activity of transcribed enhancers in the prefrontal cortex of 537 cases with schizophrenia and controls. Molecular Psychiatry, 2019, 24, 1685-1695.	7.9	40
22	Genetic Variation in Long-Range Enhancers. Current Topics in Behavioral Neurosciences, 2019, 42, 35-50.	1.7	2
23	Integrative transcriptome imputation reveals tissue-specific and shared biological mechanisms mediating susceptibility to complex traits. Nature Communications, 2019, 10, 3834.	12.8	68
24	CommonMind Consortium provides transcriptomic and epigenomic data for Schizophrenia and Bipolar Disorder. Scientific Data, 2019, 6, 180.	5.3	149
25	237. Large-Scale Gene-Trait Association Study IdentifiesÂNovel Genes Across Multiple Traits. Biological Psychiatry, 2019, 85, S98.	1.3	0
26	Assessment of somatic single-nucleotide variation in brain tissue of cases with schizophrenia. Translational Psychiatry, 2019, 9, 21.	4.8	16
27	Revealing the brain's molecular architecture. Science, 2018, 362, 1262-1263.	12.6	45
28	Integrative functional genomic analysis of human brain development and neuropsychiatric risks. Science, 2018, 362, .	12.6	516
29	Comprehensive functional genomic resource and integrative model for the human brain. Science, 2018, 362, .	12.6	618
30	Landscape of Conditional eQTL in Dorsolateral Prefrontal Cortex and Co-localization with Schizophrenia GWAS. American Journal of Human Genetics, 2018, 102, 1169-1184.	6.2	128
31	An atlas of chromatin accessibility in the adult human brain. Genome Research, 2018, 28, 1243-1252.	5.5	170
32	Evaluation of chromatin accessibility in prefrontal cortex of individuals with schizophrenia. Nature Communications, 2018, 9, 3121.	12.8	141
33	Brain Cell Type Specific Gene Expression and Co-expression Network Architectures. Scientific Reports, 2018, 8, 8868.	3.3	335
34	The Mount Sinai cohort of large-scale genomic, transcriptomic and proteomic data in Alzheimer's disease. Scientific Data, 2018, 5, 180185.	5.3	320
35	Open chromatin profiling of human postmortem brain infers functional roles for non-coding schizophrenia loci. Human Molecular Genetics, 2017, 26, 1942-1951.	2.9	69
36	Striatal H3K27 Acetylation Linked to Glutamatergic Gene Dysregulation in Human Heroin Abusers Holds Promise as Therapeutic Target. Biological Psychiatry, 2017, 81, 585-594.	1.3	77

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37	Gene expression elucidates functional impact of polygenic risk for schizophrenia. Nature Neuroscience, 2016, 19, 1442-1453.	14.8	952
38	Cardiometabolic risk loci share downstream cis- and trans-gene regulation across tissues and diseases. Science, 2016, 353, 827-830.	12.6	241
39	Understanding the genetic liability to schizophrenia through the neuroepigenome. Schizophrenia Research, 2016, 177, 115-124.	2.0	22
40	The Relationship of Common Risk Variants and Polygenic Risk for Schizophrenia to Sensorimotor Gating. Biological Psychiatry, 2016, 79, 988-996.	1.3	44
41	A Role for Noncoding Variation in Schizophrenia. Cell Reports, 2014, 9, 1417-1429.	6.4	225
42	Clearance of apoptotic corpses. Apoptosis: an International Journal on Programmed Cell Death, 2009, 14, 1029-1037.	4.9	40
43	Role of the Sc C Terminus in Transcriptional Activation and E(spl) Repressor Recruitment. Journal of Biological Chemistry, 2005, 280, 1299-1305.	3.4	24
44	The Role of the Platelet Glycoprotein IIb / IIIa in Thrombosis and Haemostasis. Current Pharmaceutical Design, 2004, 10, 1567-1576.	1.9	107
45	Characterization of a ligand-attenuated binding site on glycoprotein IIb/IIIa. Thrombosis and Haemostasis, 2002, 88, 811-6.	3.4	0
46	A Val193Met mutation in GPIIIa results in a GPIIb/IIIa receptor with a constitutively high affinity for a small ligand. British Journal of Haematology, 2001, 115, 131-139.	2.5	11