Till F M Andlauer

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genome-wide association analyses identify 44 risk variants and refine the genetic architecture of major depression. Nature Genetics, 2018, 50, 668-681.	9.4	2,224
2	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	6.0	1,085
3	Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. Cell, 2019, 179, 1469-1482.e11.	13.5	935
4	Multiple sclerosis genomic map implicates peripheral immune cells and microglia in susceptibility. Science, 2019, 365, .	6.0	710
5	Minimal phenotyping yields genome-wide association signals of low specificity for major depression. Nature Genetics, 2020, 52, 437-447.	9.4	207
6	DNA methylation as a mediator of HLA-DRB1*15:01 and a protective variant in multiple sclerosis. Nature Communications, 2018, 9, 2397.	5.8	147
7	The Genetics of the Mood Disorder Spectrum: Genome-wide Association Analyses of More Than 185,000 Cases and 439,000 Controls. Biological Psychiatry, 2020, 88, 169-184.	0.7	137
8	Novel multiple sclerosis susceptibility loci implicated in epigenetic regulation. Science Advances, 2016, 2, e1501678.	4.7	133
9	Low-Frequency and Rare-Coding Variation Contributes to Multiple Sclerosis Risk. Cell, 2018, 175, 1679-1687.e7.	13.5	115
10	A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. Biological Psychiatry, 2021, 90, 611-620.	0.7	103
11	Structural Long-Term Changes at Mushroom Body Input Synapses. Current Biology, 2010, 20, 1938-1944.	1.8	93
12	The Genetic Architecture of Depression in Individuals of East Asian Ancestry. JAMA Psychiatry, 2021, 78, 1258.	6.0	88
13	Does Childhood Trauma Moderate Polygenic Risk for Depression? A Meta-analysis of 5765 Subjects From the Psychiatric Genomics Consortium. Biological Psychiatry, 2018, 84, 138-147.	0.7	87
14	A nonsynonymous mutation in PLCG2 reduces the risk of Alzheimer's disease, dementia with Lewy bodies and frontotemporal dementia, and increases the likelihood of longevity. Acta Neuropathologica, 2019, 138, 237-250.	3.9	87
15	Presynapses in Kenyon Cell Dendrites in the Mushroom Body Calyx of Drosophila. Journal of Neuroscience, 2011, 31, 9696-9707.	1.7	83
16	Spermidine Suppresses Age-Associated Memory Impairment by Preventing Adverse Increase of Presynaptic Active Zone Size and Release. PLoS Biology, 2016, 14, e1002563.	2.6	82
17	Genome-wide association scan identifies new variants associated with a cognitive predictor of dyslexia. Translational Psychiatry, 2019, 9, 77.	2.4	82
18	A phenome-wide association and Mendelian Randomisation study of polygenic risk for depression in UK Biobank. Nature Communications, 2020, 11, 2301.	5.8	81

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19	Piccolo Regulates the Dynamic Assembly of Presynaptic F-Actin. Journal of Neuroscience, 2011, 31, 14250-14263.	1.7	69
20	A systems biology approach uncovers cell-specific gene regulatory effects of genetic associations in multiple sclerosis. Nature Communications, 2019, 10, 2236.	5.8	65
21	The genetic basis of major depression. Psychological Medicine, 2021, 51, 2217-2230.	2.7	65
22	Genetic effects influencing risk for major depressive disorder in China and Europe. Translational Psychiatry, 2017, 7, e1074-e1074.	2.4	64
23	Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. Biological Psychiatry, 2022, 91, 102-117.	0.7	61
24	Genome-wide association study reveals new insights into the heritability and genetic correlates of developmental dyslexia. Molecular Psychiatry, 2021, 26, 3004-3017.	4.1	56
25	DeepWAS: Multivariate genotype-phenotype associations by directly integrating regulatory information using deep learning. PLoS Computational Biology, 2020, 16, e1007616.	1.5	54
26	The Irre Cell Recognition Module (IRM) Proteins. Journal of Neurogenetics, 2009, 23, 48-67.	0.6	53
27	Treatment response classes in major depressive disorder identified by model-based clustering and validated by clinical prediction models. Translational Psychiatry, 2019, 9, 187.	2.4	51
28	Hair Cortisol in Twins: Heritability and Genetic Overlap with Psychological Variables and Stress-System Genes. Scientific Reports, 2017, 7, 15351.	1.6	50
29	PALS1 Is Essential for Retinal Pigment Epithelium Structure and Neural Retina Stratification. Journal of Neuroscience, 2011, 31, 17230-17241.	1.7	48
30	Active Zone Scaffold Protein Ratios Tune Functional Diversity across Brain Synapses. Cell Reports, 2018, 23, 1259-1274.	2.9	47
31	A longitudinal approach to biological psychiatric research: The PsyCourse study. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, 89-102.	1.1	47
32	An Investigation of Psychosis Subgroups With Prognostic Validation and Exploration of Genetic Underpinnings. JAMA Psychiatry, 2020, 77, 523.	6.0	39
33	Drep-2 is a novel synaptic protein important for learning and memory. ELife, 2014, 3, .	2.8	39
34	Sunlight exposure exerts immunomodulatory effects to reduce multiple sclerosis severity. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	3.3	38
35	Fatigue, depression, and pain in multiple sclerosis: How neuroinflammation translates into dysfunctional reward processing and anhedonic symptoms. Multiple Sclerosis Journal, 2022, 28, 1020-1027.	1.4	37
36	Genetic comorbidity between major depression and cardioâ€metabolic traits, stratified by age at onset of major depression. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2020, 183, 309-330.	1.1	33

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37	Bipolar multiplex families have an increased burden of common risk variants for psychiatric disorders. Molecular Psychiatry, 2021, 26, 1286-1298.	4.1	33
38	Quantitative Analysis of <i>Drosophila</i> Larval Neuromuscular Junction Morphology. Cold Spring Harbor Protocols, 2012, 2012, pdb.prot068601.	0.2	29
39	Classical Human Leukocyte Antigen Alleles and C4 Haplotypes Are Not Significantly Associated With Depression. Biological Psychiatry, 2020, 87, 419-430.	0.7	27
40	A high affinity RIM-binding protein/Aplip1 interaction prevents the formation of ectopic axonal active zones. ELife, 2015, 4, .	2.8	26
41	The genetic relationship between educational attainment and cognitive performance in major psychiatric disorders. Translational Psychiatry, 2019, 9, 210.	2.4	24
42	Identification of transdiagnostic psychiatric disorder subtypes using unsupervised learning. Neuropsychopharmacology, 2021, 46, 1895-1905.	2.8	24
43	Loss of the Coffin-Lowry syndrome associated gene <i>RSK2</i> alters ERK activity, synaptic function and axonal transport in <i>Drosophila</i> motoneurons. DMM Disease Models and Mechanisms, 2015, 8, 1389-400.	1.2	23
44	Clinical and genetic differences between bipolar disorder type 1 and 2 in multiplex families. Translational Psychiatry, 2021, 11, 31.	2.4	22
45	Inner retinal layer thinning in radiologically isolated syndrome predicts conversion to multiple sclerosis. European Journal of Neurology, 2020, 27, 2217-2224.	1.7	21
46	Investigating polygenic burden in age at disease onset in bipolar disorder: Findings from an international multicentric study. Bipolar Disorders, 2019, 21, 68-75.	1.1	20
47	Associations of schizophrenia risk genes ZNF804A and CACNA1C with schizotypy and modulation of attention in healthy subjects. Schizophrenia Research, 2019, 208, 67-75.	1.1	20
48	Advanced paternal age as a risk factor for neurodevelopmental disorders: a translational study. Molecular Autism, 2020, 11, 54.	2.6	20
49	Characterisation of age and polarity at onset in bipolar disorder. British Journal of Psychiatry, 2021, 219, 659-669.	1.7	20
50	In Vivo Imaging of the <i>Drosophila</i> Larval Neuromuscular Junction. Cold Spring Harbor Protocols, 2012, 2012, pdb.prot068593.	0.2	18
51	Evidence for increased genetic risk load for major depression in patients assigned to electroconvulsive therapy. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, 35-45.	1.1	18
52	Higher frequencies of HLA DQB1*05:01 and anti-glycosphingolipid antibodies in a cluster of severe Guillain–Barré syndrome. Journal of Neurology, 2016, 263, 2105-2113.	1.8	17
53	Childhood maltreatment and cognitive functioning: the role of depression, parental education, and polygenic predisposition. Neuropsychopharmacology, 2021, 46, 891-899.	2.8	17
54	Association of Whole-Genome and NETRIN1 Signaling Pathway–Derived Polygenic Risk Scores for Major Depressive Disorder and White Matter Microstructure in the UK Biobank. Biological Psychiatry: Cognitive Neuroscience and Neuroimaging, 2019, 4, 91-100.	1.1	16

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55	Successful Replication of GWAS Hits for Multiple Sclerosis in 10,000 Germans Using the Exome Array. Genetic Epidemiology, 2015, 39, 601-608.	0.6	15
56	<i>Cis</i> -epistasis at the <i>LPA</i> locus and risk of cardiovascular diseases. Cardiovascular Research, 2022, 118, 1088-1102.	1.8	14
5 7	Polygenic scores for psychiatric disease: from research tool to clinical application. Medizinische Genetik, 2020, 32, 39-45.	0.1	14
58	In Vivo Imaging of Drosophila Larval Neuromuscular Junctions to Study Synapse Assembly. Cold Spring Harbor Protocols, 2012, 2012, pdb.top068577-pdb.top068577.	0.2	13
59	Exome sequencing in large, multiplex bipolar disorder families from Cuba. PLoS ONE, 2018, 13, e0205895.	1.1	13
60	The role of environmental stress and DNA methylation in the longitudinal course of bipolar disorder. International Journal of Bipolar Disorders, 2020, 8, 9.	0.8	13
61	Genetic Variation in <scp><i>WNT9B</i></scp> Increases Relapse Hazard in Multiple Sclerosis. Annals of Neurology, 2021, 89, 884-894.	2.8	12
62	The Aryl Hydrocarbon Receptor–Dependent TGF-α/VEGF-B Ratio Correlates With Disease Subtype and Prognosis in Multiple Sclerosis. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, .	3.1	12
63	Treatment- and population-specific genetic risk factors for anti-drug antibodies against interferon-beta: a GWAS. BMC Medicine, 2020, 18, 298.	2.3	11
64	Polygenic risk scores across the extended psychosis spectrum. Translational Psychiatry, 2021, 11, 600.	2.4	11
65	Polygenic risk for schizophrenia and schizotypal traits in non-clinical subjects. Psychological Medicine, 2022, 52, 1069-1079.	2.7	10
66	The influence of religious activity and polygenic schizophrenia risk on religious delusions in schizophrenia. Schizophrenia Research, 2019, 210, 255-261.	1.1	9
67	Effect of <i>HLA-DRB1</i> alleles and genetic variants on the development of neutralizing antibodies to interferon beta in the BEYOND and BENEFIT trials. Multiple Sclerosis Journal, 2019, 25, 565-573.	1.4	9
68	HLA Genetic Risk Burden in Multiple Sclerosis. JAMA Neurology, 2016, 73, 1500.	4.5	8
69	Building an Imaging Chamber for In Vivo Imaging of <i>Drosophila</i> Larvae. Cold Spring Harbor Protocols, 2012, 2012, pdb.prot068585.	0.2	7
70	MS susceptibility is not affected by single nucleotide polymorphisms in the MMP9 gene. Journal of Neuroimmunology, 2015, 279, 46-49.	1.1	7
71	Genetic determinants of the humoral immune response in MS. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, e827.	3.1	7
72	A genome-wide association study of the longitudinal course of executive functions. Translational Psychiatry, 2021, 11, 386.	2.4	7

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73	Fighting the famine with an amine: synaptic strategies for smart search. Nature Neuroscience, 2011, 14, 124-126.	7.1	6
74	Gene Expression in Spontaneous Experimental Autoimmune Encephalomyelitis Is Linked to Human Multiple Sclerosis Risk Genes. Frontiers in Immunology, 2020, 11, 2165.	2.2	6
75	"The Heidelberg Five―personality dimensions: Genomeâ€wide associations, polygenic risk for neuroticism, and psychopathology 20 years after assessment. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2021, 186, 77-89.	1.1	6
76	Polygenic scores differentially predict developmental trajectories of subtypes of social withdrawal in childhood. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2021, 62, 1320-1329.	3.1	6
77	Genetic risk for psychiatric illness is associated with the number of hospitalizations of bipolar disorder patients. Journal of Affective Disorders, 2022, 296, 532-540.	2.0	6
78	Interaction of developmental factors and ordinary stressful life events on brain structure in adults. NeuroImage: Clinical, 2021, 30, 102683.	1.4	5
79	GWAS meta-analysis followed by Mendelian randomization revealed potential control mechanisms for circulating α-Klotho levels. Human Molecular Genetics, 2022, 31, 792-802.	1.4	5
80	Genetic factors influencing a neurobiological substrate for psychiatric disorders. Translational Psychiatry, 2021, 11, 192.	2.4	4
81	Gray matter atrophy in relapsing-remitting multiple sclerosis is associated with white matter lesions in connecting fibers. Multiple Sclerosis Journal, 2022, 28, 900-909.	1.4	4
82	A multiâ€informant and multiâ€polygenic approach to understanding predictors of peer victimisation in childhood and adolescence. JCPP Advances, 2022, 2, .	1.4	3
83	Investigating the phenotypic and genetic associations between personality traits and suicidal behavior across major mental health diagnoses. European Archives of Psychiatry and Clinical Neuroscience, 2022, , 1.	1.8	2
84	EFFECTS OF SCHIZOPHRENIA AND BIPOLAR POLYGENIC RISK SCORES ON AGE AT ONSET IN BIPOLAR DISORDER. European Neuropsychopharmacology, 2019, 29, S967.	0.3	1
85	SU62THE ROLE OF ENVIRONMENTAL STRESS AND DNA METHYLATION IN THE LONGITUDINAL COURSE OF BIPOLAR DISORDER. European Neuropsychopharmacology, 2019, 29, S1300-S1301.	0.3	1
86	Genotype-phenotype feasibility studies on khat abuse, traumatic experiences and psychosis in Ethiopia. Psychiatric Genetics, 2020, 30, 34-38.	0.6	1
87	Interplay between the genetics of personality traits, severe psychiatric disorders and COVID-19 host genetics in the susceptibility to SARS-CoV-2 infection. BJPsych Open, 2021, 7, e188.	0.3	1
88	Polygenic Risk For BIP, MDD, And SCZ In Andalusian Multiplex Families. European Neuropsychopharmacology, 2017, 27, S385-S386.	0.3	0
89	Integrating Polygenic Allele Burden Information And Phenomic Data To Characterize Complex Disease Trajectories In Severe Mental Illness. European Neuropsychopharmacology, 2017, 27, S406.	0.3	0
90	POLYGENIC BURDEN ANALYSIS OF LONGITUDINAL CLUSTERS OF QUALITY OF LIFE AND FUNCTIONING IN PATIENTS WITH SEVERE MENTAL ILLNESS. European Neuropsychopharmacology, 2017, 27, S408-S409.	0.3	0

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91	Using Machine Learning To Build Individualized Prediction Models Of Future Quality Of Life In Psychosis Patients. European Neuropsychopharmacology, 2017, 27, S464.	0.3	0
92	F96POLYGENIC RISK SCORE ANALYSIS OF TRAJECTORIES OF COGNITIVE PERFORMANCE IN PSYCHIATRIC PATIENTS. European Neuropsychopharmacology, 2019, 29, S1161.	0.3	0
93	Cover Image, Volume 180B, Number 2, March 2019. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, i.	1.1	0
94	S13. IMPACT OF POLYGENIC AND POLY-ENVIRONMENTAL RISK FACTORS ON A PSYCHOSIS RISK PHENOTYPE EXPLAINED THROUGH BRAIN STRUCTURE. Schizophrenia Bulletin, 2020, 46, S35-S36.	2.3	0
95	A Nonsynonymous Mutation in PLCG2 Reduces the Risk of Alzheimer's Disease, Dementia with Lewy-Bodies and Frontotemporal Dementia, and Increases the Likelihood of Longevity. SSRN Electronic Journal, O, , .	0.4	0
96	Interplay between the Genetics of Personality Traits, severe Psychiatric Disorders, and COVID-19 Host Genetics in the Susceptibility to SARS-CoV-2 Infection - ADDENDUM. BJPsych Open, 2021, 7, e206.	0.3	0