Till F M Andlauer

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | 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 |
| 2 | Polygenic risk for schizophrenia and schizotypal traits in non-clinical subjects. Psychological Medicine, 2022, 52, 1069-1079. | 2.7 | 10 |
| 3 | Sex-Dependent Shared and Nonshared Genetic Architecture Across Mood and Psychotic Disorders. Biological Psychiatry, 2022, 91, 102-117. | 0.7 | 61 |
| 4 | <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 | 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 |
| 6 | 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 |
| 7 | 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 |
| 8 | 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 |
| 9 | A multiâ€informant and multiâ€polygenic approach to understanding predictors of peer victimisation in childhood and adolescence. JCPP Advances, 2022, 2, . | 1.4 | 3 |
| 10 | 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 |
| 11 | Childhood maltreatment and cognitive functioning: the role of depression, parental education, and polygenic predisposition. Neuropsychopharmacology, 2021, 46, 891-899. | 2.8 | 17 |
| 12 | Bipolar multiplex families have an increased burden of common risk variants for psychiatric disorders. Molecular Psychiatry, 2021, 26, 1286-1298. | 4.1 | 33 |
| 13 | Interaction of developmental factors and ordinary stressful life events on brain structure in adults. NeuroImage: Clinical, 2021, 30, 102683. | 1.4 | 5 |
| 14 | Clinical and genetic differences between bipolar disorder type 1 and 2 in multiplex families. Translational Psychiatry, 2021, 11, 31. | 2.4 | 22 |
| 15 | "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 |
| 16 | Genetic factors influencing a neurobiological substrate for psychiatric disorders. Translational Psychiatry, 2021, 11, 192. | 2.4 | 4 |
| 17 | The genetic basis of major depression. Psychological Medicine, 2021, 51, 2217-2230. | 2.7 | 65 |
| 18 | Genetic Variation in <scp><i>WNT9B</i></scp> Increases Relapse Hazard in Multiple Sclerosis. Annals of Neurology, 2021, 89, 884-894. | 2.8 | 12 |

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|----|--|-----|-----------|
| 19 | A Comparison of Ten Polygenic Score Methods for Psychiatric Disorders Applied Across Multiple Cohorts. Biological Psychiatry, 2021, 90, 611-620. | 0.7 | 103 |
| 20 | 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 |
| 21 | A genome-wide association study of the longitudinal course of executive functions. Translational Psychiatry, 2021, 11, 386. | 2.4 | 7 |
| 22 | Identification of transdiagnostic psychiatric disorder subtypes using unsupervised learning. Neuropsychopharmacology, 2021, 46, 1895-1905. | 2.8 | 24 |
| 23 | 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 |
| 24 | Characterisation of age and polarity at onset in bipolar disorder. British Journal of Psychiatry, 2021, 219, 659-669. | 1.7 | 20 |
| 25 | The Genetic Architecture of Depression in Individuals of East Asian Ancestry. JAMA Psychiatry, 2021, 78, 1258. | 6.0 | 88 |
| 26 | 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 |
| 27 | 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 |
| 28 | Polygenic risk scores across the extended psychosis spectrum. Translational Psychiatry, 2021, 11, 600. | 2.4 | 11 |
| 29 | 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 | Ο |
| 30 | Classical Human Leukocyte Antigen Alleles and C4 Haplotypes Are Not Significantly Associated With Depression. Biological Psychiatry, 2020, 87, 419-430. | 0.7 | 27 |
| 31 | 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 |
| 32 | Genotype-phenotype feasibility studies on khat abuse, traumatic experiences and psychosis in Ethiopia. Psychiatric Genetics, 2020, 30, 34-38. | 0.6 | 1 |
| 33 | Genetic determinants of the humoral immune response in MS. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, e827. | 3.1 | 7 |
| 34 | 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 |
| 35 | Gene Expression in Spontaneous Experimental Autoimmune Encephalomyelitis Is Linked to Human Multiple Sclerosis Risk Genes. Frontiers in Immunology, 2020, 11, 2165. | 2.2 | 6 |
| 36 | Treatment- and population-specific genetic risk factors for anti-drug antibodies against interferon-beta: a GWAS. BMC Medicine, 2020, 18, 298. | 2.3 | 11 |

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|----|---|-----|-----------|
| 37 | A phenome-wide association and Mendelian Randomisation study of polygenic risk for depression in UK Biobank. Nature Communications, 2020, 11, 2301. | 5.8 | 81 |
| 38 | 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 |
| 39 | Minimal phenotyping yields genome-wide association signals of low specificity for major depression. Nature Genetics, 2020, 52, 437-447. | 9.4 | 207 |
| 40 | Advanced paternal age as a risk factor for neurodevelopmental disorders: a translational study. Molecular Autism, 2020, 11, 54. | 2.6 | 20 |
| 41 | Inner retinal layer thinning in radiologically isolated syndrome predicts conversion to multiple sclerosis. European Journal of Neurology, 2020, 27, 2217-2224. | 1.7 | 21 |
| 42 | DeepWAS: Multivariate genotype-phenotype associations by directly integrating regulatory information using deep learning. PLoS Computational Biology, 2020, 16, e1007616. | 1.5 | 54 |
| 43 | An Investigation of Psychosis Subgroups With Prognostic Validation and Exploration of Genetic Underpinnings. JAMA Psychiatry, 2020, 77, 523. | 6.0 | 39 |
| 44 | 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 |
| 45 | Polygenic scores for psychiatric disease: from research tool to clinical application. Medizinische Genetik, 2020, 32, 39-45. | 0.1 | 14 |
| 46 | 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 |
| 47 | 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 |
| 48 | 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 |
| 49 | F96POLYGENIC RISK SCORE ANALYSIS OF TRAJECTORIES OF COGNITIVE PERFORMANCE IN PSYCHIATRIC PATIENTS. European Neuropsychopharmacology, 2019, 29, S1161. | 0.3 | 0 |
| 50 | The genetic relationship between educational attainment and cognitive performance in major psychiatric disorders. Translational Psychiatry, 2019, 9, 210. | 2.4 | 24 |
| 51 | Multiple sclerosis genomic map implicates peripheral immune cells and microglia in susceptibility. Science, 2019, 365, . | 6.0 | 710 |
| 52 | 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 |
| 53 | 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 |
| 54 | A systems biology approach uncovers cell-specific gene regulatory effects of genetic associations in multiple sclerosis. Nature Communications, 2019, 10, 2236. | 5.8 | 65 |

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|----|---|------|-----------|
| 55 | Cover Image, Volume 180B, Number 2, March 2019. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2019, 180, i. | 1.1 | 0 |
| 56 | EFFECTS OF SCHIZOPHRENIA AND BIPOLAR POLYGENIC RISK SCORES ON AGE AT ONSET IN BIPOLAR DISORDER. European Neuropsychopharmacology, 2019, 29, S967. | 0.3 | 1 |
| 57 | Genome-wide association scan identifies new variants associated with a cognitive predictor of dyslexia. Translational Psychiatry, 2019, 9, 77. | 2.4 | 82 |
| 58 | SU62THE ROLE OF ENVIRONMENTAL STRESS AND DNA METHYLATION IN THE LONGITUDINAL COURSE OF BIPOLAR DISORDER. European Neuropsychopharmacology, 2019, 29, S1300-S1301. | 0.3 | 1 |
| 59 | Genomic Relationships, Novel Loci, and Pleiotropic Mechanisms across Eight Psychiatric Disorders. Cell, 2019, 179, 1469-1482.e11. | 13.5 | 935 |
| 60 | 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 |
| 61 | The influence of religious activity and polygenic schizophrenia risk on religious delusions in schizophrenia. Schizophrenia Research, 2019, 210, 255-261. | 1.1 | 9 |
| 62 | 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 |
| 63 | 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 |
| 64 | Active Zone Scaffold Protein Ratios Tune Functional Diversity across Brain Synapses. Cell Reports, 2018, 23, 1259-1274. | 2.9 | 47 |
| 65 | 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 |
| 66 | 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 |
| 67 | Exome sequencing in large, multiplex bipolar disorder families from Cuba. PLoS ONE, 2018, 13, e0205895. | 1.1 | 13 |
| 68 | Low-Frequency and Rare-Coding Variation Contributes to Multiple Sclerosis Risk. Cell, 2018, 175, 1679-1687.e7. | 13.5 | 115 |
| 69 | Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, . | 6.0 | 1,085 |
| 70 | 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 |
| 71 | Genetic effects influencing risk for major depressive disorder in China and Europe. Translational Psychiatry, 2017, 7, e1074-e1074. | 2.4 | 64 |
| 72 | Polygenic Risk For BIP, MDD, And SCZ In Andalusian Multiplex Families. European Neuropsychopharmacology, 2017, 27, S385-S386. | 0.3 | 0 |

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|----|---|-----|-----------|
| 73 | Integrating Polygenic Allele Burden Information And Phenomic Data To Characterize Complex Disease Trajectories In Severe Mental Illness. European Neuropsychopharmacology, 2017, 27, S406. | 0.3 | Ο |
| 74 | 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 |
| 75 | Using Machine Learning To Build Individualized Prediction Models Of Future Quality Of Life In Psychosis Patients. European Neuropsychopharmacology, 2017, 27, S464. | 0.3 | 0 |
| 76 | Hair Cortisol in Twins: Heritability and Genetic Overlap with Psychological Variables and Stress-System Genes. Scientific Reports, 2017, 7, 15351. | 1.6 | 50 |
| 77 | 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 |
| 78 | 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 |
| 79 | HLA Genetic Risk Burden in Multiple Sclerosis. JAMA Neurology, 2016, 73, 1500. | 4.5 | 8 |
| 80 | Novel multiple sclerosis susceptibility loci implicated in epigenetic regulation. Science Advances, 2016, 2, e1501678. | 4.7 | 133 |
| 81 | 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 |
| 82 | 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 |
| 83 | MS susceptibility is not affected by single nucleotide polymorphisms in the MMP9 gene. Journal of Neuroimmunology, 2015, 279, 46-49. | 1.1 | 7 |
| 84 | A high affinity RIM-binding protein/Aplip1 interaction prevents the formation of ectopic axonal active zones. ELife, 2015, 4, . | 2.8 | 26 |
| 85 | Drep-2 is a novel synaptic protein important for learning and memory. ELife, 2014, 3, . | 2.8 | 39 |
| 86 | 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 |
| 87 | In Vivo Imaging of the <i>Drosophila</i> Larval Neuromuscular Junction. Cold Spring Harbor Protocols, 2012, 2012, pdb.prot068593. | 0.2 | 18 |
| 88 | Quantitative Analysis of <i>Drosophila</i> Larval Neuromuscular Junction Morphology. Cold Spring Harbor Protocols, 2012, 2012, pdb.prot068601. | 0.2 | 29 |
| 89 | Building an Imaging Chamber for In Vivo Imaging of <i>Drosophila</i> Larvae. Cold Spring Harbor Protocols, 2012, 2012, pdb.prot068585. | 0.2 | 7 |
| 90 | Fighting the famine with an amine: synaptic strategies for smart search. Nature Neuroscience, 2011, 14, 124-126. | 7.1 | 6 |

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|----|--|-----|-----------|
| 91 | Piccolo Regulates the Dynamic Assembly of Presynaptic F-Actin. Journal of Neuroscience, 2011, 31, 14250-14263. | 1.7 | 69 |
| 92 | Presynapses in Kenyon Cell Dendrites in the Mushroom Body Calyx of Drosophila. Journal of Neuroscience, 2011, 31, 9696-9707. | 1.7 | 83 |
| 93 | PALS1 Is Essential for Retinal Pigment Epithelium Structure and Neural Retina Stratification. Journal of Neuroscience, 2011, 31, 17230-17241. | 1.7 | 48 |
| 94 | Structural Long-Term Changes at Mushroom Body Input Synapses. Current Biology, 2010, 20, 1938-1944. | 1.8 | 93 |
| 95 | The Irre Cell Recognition Module (IRM) Proteins. Journal of Neurogenetics, 2009, 23, 48-67. | 0.6 | 53 |
| 96 | 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, 0, , . | 0.4 | 0 |