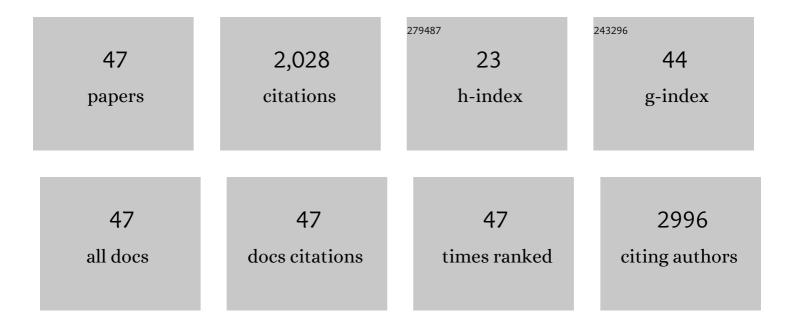
Isabelle Massat

List of Publications by Year in descending order

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Version: 2024-02-01



#	Article	IF	CITATIONS
1	Clinical Factors Associated With Treatment Resistance in Major Depressive Disorder. Journal of Clinical Psychiatry, 2007, 68, 1062-1070.	1.1	407
2	Association between COMT (Val158Met) functional polymorphism and early onset in patients with major depressive disorder in a European multicenter genetic association study. Molecular Psychiatry, 2005, 10, 598-605.	4.1	134
3	Switching Antidepressant Class Does Not Improve Response or Remission in Treatment-Resistant Depression. Journal of Clinical Psychopharmacology, 2011, 31, 512-516.	0.7	83
4	Executive and attentional contributions to Theory of Mind deficit in attention deficit/hyperactivity disorder (ADHD). Child Neuropsychology, 2016, 22, 345-365.	0.8	79
5	Serotonin transporter 5HTTLPR polymorphism and affective disorders: no evidence of association in a large European multicenter study. European Journal of Human Genetics, 2004, 12, 377-382.	1.4	78
6	Structural and functional neuroimaging in attentionâ€deficit/hyperactivity disorder. Developmental Medicine and Child Neurology, 2019, 61, 399-405.	1.1	78
7	Consensus paper of the WFSBP Task Force on Genetics: Genetics, epigenetics and gene expression markers of major depressive disorder and antidepressant response. World Journal of Biological Psychiatry, 2017, 18, 5-28.	1.3	75
8	Cytochrome P450 CYP1A2, CYP2C9, CYP2C19 and CYP2D6 genes are not associated with response and remission in a sample of depressive patients. International Clinical Psychopharmacology, 2009, 24, 250-256.	0.9	69
9	Brain-derived neurotrophic factor gene polymorphisms. International Clinical Psychopharmacology, 2011, 26, 1-10.	0.9	67
10	Tryptophan hydroxylase polymorphism and suicidality in unipolar and bipolar affective disorders: a multicenter association study. Biological Psychiatry, 2001, 49, 405-409.	0.7	66
11	Non-replication of the brain-derived neurotrophic factor (BDNF) association in bipolar affective disorder: A Belgian patient-control study. American Journal of Medical Genetics Part A, 2004, 129B, 34-35.	2.4	62
12	Working Memory-Related Functional Brain Patterns in Never Medicated Children with ADHD. PLoS ONE, 2012, 7, e49392.	1.1	61
13	Excess of allele1 for α3 subunit GABA receptor gene (GABRA3) in bipolar patients: a multicentric association study. Molecular Psychiatry, 2002, 7, 201-207.	4.1	51
14	The impact of catechol-O-methyltransferase SNPs and haplotypes on treatment response phenotypes in major depressive disorder: a case–control association study. International Clinical Psychopharmacology, 2010, 25, 218-227.	0.9	51
15	Positive association of dopamine D2 receptor polymorphism with bipolar affective disorder in a European multicenter association study of affective disorders. American Journal of Medical Genetics Part A, 2002, 114, 177-185.	2.4	50
16	A preliminary investigation of the influence of CREB1 gene on treatment resistance in major depression. Journal of Affective Disorders, 2011, 128, 56-63.	2.0	45
17	Gene-based SNP genetic association study of the corticotropin-releasing hormone receptor-2 (CRHR2) in major depression. American Journal of Medical Genetics Part A, 2002, 114, 222-226.	2.4	41
18	Citalopram versus desipramine in treatment resistant depression: Effect of continuation or switching strategies. A randomized open study. World Journal of Biological Psychiatry, 2011, 12, 364-375.	1.3	40

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19	A European multicenter association study ofHTR2A receptor polymorphism in bipolar affective disorder. , 2000, 96, 136-140.		38
20	Evaluation of the role of MAPK1 and CREB1 polymorphisms on treatment resistance, response and remission in mood disorder patients. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2013, 44, 271-278.	2,5	38
21	5HT1A and 5HT2A receptor genes in treatment response phenotypes in major depressive disorder. International Clinical Psychopharmacology, 2010, 25, 228-231.	0.9	37
22	No implication of brain-derived neurotrophic factor (BDNF) gene in unipolar affective disorder: Evidence from Belgian first and replication patient–control studies. European Neuropsychopharmacology, 2005, 15, 491-495.	0.3	32
23	COMT and age at onset in mood disorders: A replication and extension study. Neuroscience Letters, 2011, 498, 218-221.	1.0	32
24	ADHD and ASD: distinct brain patterns of inhibition-related activation?. Translational Psychiatry, 2020, 10, 24.	2.4	28
25	Grey matter volume differences associated with gender in children with attention-deficit/hyperactivity disorder: A voxel-based morphometry study. Developmental Cognitive Neuroscience, 2015, 14, 32-37.	1.9	26
26	Genetic association between the phospholipase A2 gene and unipolar affective disorder: a multicentre case???control study. Psychiatric Genetics, 2003, 13, 211-220.	0.6	22
27	Failure to Replicate Influence of GRIK4 and GNB3 Polymorphisms on Treatment Outcome in Major Depression. Neuropsychobiology, 2012, 65, 70-75.	0.9	22
28	Influence of COX-2 and OXTR polymorphisms on treatment outcome in treatment resistant depression. Neuroscience Letters, 2012, 516, 85-88.	1.0	21
29	Lack of association between the 5HT2A receptor polymorphism (T102C) and unipolar affective disorder in a multicentric European study. European Neuropsychopharmacology, 2003, 13, 365-368.	0.3	20
30	The impact of serotonin receptor 1A and 2A gene polymorphisms and interactions on suicide attempt and suicide risk in depressed patients with insufficient response to treatment – a European multicentre study. International Clinical Psychopharmacology, 2016, 31, 1-7.	0.9	19
31	Identification of clinical factors associated with resistance to antidepressants in bipolar depression: results from an European Multicentre Study. International Clinical Psychopharmacology, 2010, 25, 297-301.	0.9	18
32	Hyperactivity in motor response inhibition networks in unmedicated children with attention deficit-hyperactivity disorder. World Journal of Biological Psychiatry, 2018, 19, 101-111.	1.3	17
33	The impact of Cytochrome P450 CYP1A2, CYP2C9, CYP2C19 and CYP2D6 genes on suicide attempt and suicide risk—a European multicentre study on treatment-resistant major depressive disorder. European Archives of Psychiatry and Clinical Neuroscience, 2013, 263, 385-391.	1.8	16
34	Attentional control of emotional interference in children with ADHD and typically developing children: An emotional N-back study. Psychiatry Research, 2017, 254, 1-7.	1.7	16
35	Dopamine transporter genotype modulates brain activity during a working memory task in children with ADHD. Research in Developmental Disabilities, 2019, 92, 103430.	1.2	13
36	No influence of PTGS2 polymorphisms on response and remission to antidepressants in major depression. Psychiatry Research, 2011, 188, 166-169.	1.7	10

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#	Article	IF	CITATIONS
37	Positive association of dopamine D2 receptor polymorphism with bipolar affective disorder in a European Multicenter Association Study of affective disorders. American Journal of Medical Genetics Part A, 2002, 114, 177-85.	2.4	10
38	Disorder-specific brain volumetric abnormalities in Attention-Deficit/Hyperactivity Disorder relative to Autism Spectrum Disorder. PLoS ONE, 2020, 15, e0241856.	1.1	9
39	No evidence for the involvement of CAC/CTG repeats from within 18q21.33–q23 in bipolar disorder. European Journal of Human Genetics, 2000, 8, 385-388.	1.4	8
40	Relationship Between White Matter Abnormalities and Neuropsychological Measures in Children With ADHD. Journal of Attention Disorders, 2020, 24, 1020-1031.	1.5	8
41	Motor Abnormalities in Attention-Deficit/Hyperactivity Disorder and Autism Spectrum Disorder Are Associated With Regional Grey Matter Volumes. Frontiers in Neurology, 2021, 12, 666980.	1.1	8
42	Association study ofCREB1polymorphisms and suicidality in MDD: results from a European multicenter study on treatment resistant depression. International Journal of Neuroscience, 2015, 125, 336-343.	0.8	7
43	Lack of genetic association between the phospholipase A2 gene and bipolar mood disorder in a European multicentre case–control study. Psychiatric Genetics, 2006, 16, 169-171.	0.6	5
44	Expanded RED products and loci containing CAG/CTG repeats on chromosome 17 (ERDA1) and chromosome 18 (CTG18.1) in trans-generational pairs with bipolar affective disorder. , 2004, 128B, 71-75.		4
45	Dysbindin gene (DTNBP1) in major depressive disorder (MDD) patients: Lack of association with clinical phenotypes. World Journal of Biological Psychiatry, 2010, 11, 985-990.	1.3	4
46	A European multicenter association study of HTR2A receptor polymorphism in bipolar affective disorder. American Journal of Medical Genetics Part A, 2000, 96, 136-140.	2.4	2
47	F55. An Image-Based Meta-Analysis of Successful and Failed Stopping in Attention Deficit/Hyperactivity Disorder Using Statistical Parametric Maps. Biological Psychiatry, 2019, 85, S234.	0.7	1