Araceli Rosa

List of Publications by Year
in descending order

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65 papers	1,959 citations	279798 23 h-index	254184 43 g-index
69 all docs	69 docs citations	69 times ranked	2856 citing authors

#	Article	IF	Citations
1	Toxoplasma gondii Seropositivity Interacts with Catechol-O-methyltransferase Val105/158Met Variation Increasing the Risk of Schizophrenia. Genes, 2022, 13, 1088.	2.4	3
2	Clinical, cognitive and neuroanatomical associations of serum NMDAR autoantibodies in people at clinical high risk for psychosis. Molecular Psychiatry, 2021, 26, 2590-2604.	7.9	16
3	Dysregulated Lipid Metabolism Precedes Onset of Psychosis. Biological Psychiatry, 2021, 89, 288-297.	1.3	42
4	Influence of the BDNF Val66Met polymorphism on weight loss after bariatric surgery: a 24-month follow-up. Surgery for Obesity and Related Diseases, 2021, 17, 185-192.	1.2	3
5	Telomere length in patients with obesity submitted to bariatric surgery: A systematic review. European Eating Disorders Review, 2021, 29, 842-853.	4.1	6
6	Role of the FKBP5 polymorphism rs1360780, age, sex, and type of surgery in weight loss after bariatric surgery: a follow-up study. Surgery for Obesity and Related Diseases, 2020, 16, 581-589.	1.2	11
7	Longitudinal changes in telomere length in a cohort of obese patients submitted to bariatric surgery: a 2-year follow-up. Surgery for Obesity and Related Diseases, 2020, 16, 1794-1801.	1.2	2
8	Efficacy, cost-utility and physiological effects of Acceptance and Commitment Therapy (ACT) and Behavioural Activation Treatment for Depression (BATD) in patients with chronic low back pain and depression: study protocol of a randomised, controlled trial including mobile-technology-based ecological momentary assessment (IMPACT study). BMJ Open, 2020, 10, e038107.	1.9	9
9	Oxytocin receptor gene polymorphism (rs53576) and digit ratio associates with aggression: comparison in seven ethnic groups. Journal of Physiological Anthropology, 2020, 39, 20.	2.6	10
10	Dermatoglyphic fluctuating asymmetry and total a-b ridge count as biomarkers of Foetal Alcohol Syndrome: Analysis in children adopted from Eastern Europe. Early Human Development, 2020, 143, 104999.	1.8	2
11	Response to the letter to the editor: FKBP5 polymorphism rs1360780 and weight loss after bariatric surgery. Surgery for Obesity and Related Diseases, 2020, 16, 974-975.	1.2	2
12	Interaction of both positive and negative daily-life experiences with FKBP5 haplotype on psychosis risk. European Psychiatry, 2020, 63, e11.	0.2	3
13	Dermatoglyphics in children prenatally exposed to alcohol: Fluctuating asymmetry (FA) as a biomarker of alcohol exposure. Early Human Development, 2018, 127, 90-95.	1.8	7
14	Interaction between FKBP5 variability and recent life events in the anxiety spectrum: Evidence for the differential susceptibility model. PLoS ONE, 2018, 13, e0193044.	2.5	13
15	Association between RGS4 variants and psychotic-like experiences in nonclinical individuals. European Archives of Psychiatry and Clinical Neuroscience, 2017, 267, 19-24.	3.2	7
16	Interaction between FKBP5 gene and childhood trauma on psychosis, depression and anxiety symptoms in a non-clinical sample. Psychoneuroendocrinology, 2017, 85, 200-209.	2.7	28
17	The role of stressâ€regulation genes in moderating the association of stress and dailyâ€life psychotic experiences. Acta Psychiatrica Scandinavica, 2017, 136, 389-399.	4.5	24
18	Digital dermatoglyphic study in three west Algerian populations: Reguibates, Zenata, Oran. Journal of the Canadian Society of Forensic Science, 2017, 50, 164-174.	0.9	5

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19	The genome-wide associated candidate gene ZNF804A and psychosis-proneness: Evidence of sex-modulated association. PLoS ONE, 2017, 12, e0185072.	2.5	14
20	Polymorphisms of two loci at the oxytocin receptor gene in populations of Africa, Asia and South Europe. BMC Genetics, 2016, 17, 17.	2.7	27
21	Childhood trauma, BDNF Val66Met and subclinical psychotic experiences. Attempt at replication in two independent samples. Journal of Psychiatric Research, 2016, 83, 121-129.	3.1	19
22	Influence of DAOA and RGS4 genes on the risk for psychotic disorders and their associated executive dysfunctions: A family-based study. European Psychiatry, 2016, 32, 42-47.	0.2	4
23	The Interaction between Childhood Bullying and the FKBP5 Gene on Psychotic-Like Experiences and Stress Reactivity in Real Life. PLoS ONE, 2016, 11, e0158809.	2.5	21
24	COMT-by-Sex Interaction Effect on Psychosis Proneness. BioMed Research International, 2015, 2015, 1-7.	1.9	20
25	Ten-year stability of self-reported schizotypal personality features in patients with psychosis and their healthy siblings. Psychiatry Research, 2015, 227, 283-289.	3.3	17
26	Executive functioning in schizophrenia spectrum disorder patients and their unaffected siblings: A ten-year follow-up study. Schizophrenia Research, 2013, 143, 291-296.	2.0	19
27	Lifetime cannabis use and cognition in patients with schizophrenia spectrum disorders and their unaffected siblings. European Archives of Psychiatry and Clinical Neuroscience, 2013, 263, 643-653.	3.2	25
28	Substantial genetic link between iq and working memory: Implications for molecular genetic studies on schizophrenia. the european twin study of schizophrenia (EUTwinsS). American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 413-418.	1.7	18
29	Influence of genetic variability at the COMT gene on TMT-B performance in psychotic patients and their healthy siblings. Psychiatric Genetics, 2012, 22, 92-95.	1.1	3
30	Ectodermal markers of early developmental impairment in very preterm individuals. Psychiatry Research, 2012, 200, 715-718.	3.3	3
31	Convergent evidence of the contribution of TP53 genetic variation (Pro72Arg) to metabolic activity and white matter volume in the frontal lobe in schizophrenia patients. NeuroImage, 2011, 56, 45-51.	4.2	19
32	Hidalgo Borrajo, R., et al., Validity of maternal recall of obstetric complications in mothers of patients with schizophrenia spectrum disorders and their healthy siblings, Schizophr. Res. (2010), doi:10.1016/j.schres.2010.09.017. Schizophrenia Research, 2011, 126, 308-309.	2.0	5
33	Evaluation of TMT-B performance in patients with psychosis and their healthy sib-pairs. International Clinical Psychopharmacology, 2011, 26, e109-e110.	1.7	0
34	Genetic variation in dysbindin gene influences both the risk for functional psychosis and the cognitive functioning in a Spanish family based study. International Clinical Psychopharmacology, 2011, 26, e59-e60.	1.7	0
35	Dysbindinâ€1 gene contributes differentially to earlyâ€and adultâ€onset forms of functional psychosis. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 322-333.	1.7	22
36	Baseline Brain Perfusion and the Serotonin Transporter Promoter Polymorphism. Biological Psychiatry, 2010, 67, 317-322.	1.3	33

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37	IMPACT OF SPECIFIC TYPES OF EARLY ADVERSITY EVENTS ON ADULT PSYCHOSIS-LIKE SYMPTOMS: PRELIMINARY RESULTS BASED IN THE UB-TWIN SAMPLE. Schizophrenia Research, 2010, 117, 269-270.	2.0	1
38	Decay of linkage disequilibrium within genes across HGDP-CEPH human samples: most population isolates do not show increased LD. BMC Genomics, 2009, 10, 338.	2.8	19
39	COMT Val¹⁵⁸Met moderation of cannabisâ€induced psychosis: a momentary assessment study of â€˜switching onâ€™ hallucinations in the flow of daily life. Acta Psychiatrica Scandinavica, 2009, 119, 156-160.	4.5	106
40	Evidence that the COMTVal158Met polymorphism moderates sensitivity to stress in psychosis: An experience-sampling study. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 10-17.	1.7	104
41	Differential methylation of the Xâ€chromosome is a possible source of discordance for bipolar disorder female monozygotic twins. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 459-462.	1.7	70
42	Gray matter deficits in bipolar disorder are associated with genetic variability at interleukinâ€1 beta gene (2q13). Genes, Brain and Behavior, 2008, 7, 796-801.	2.2	54
43	Identification of two risk haplotypes for schizophrenia and bipolar disorder in the synaptic vesicle monoamine transporter gene (SVMT). American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 502-507.	1.7	19
44	Familyâ€based association study of neuregulinâ€1 gene and psychosis in a Spanish sample. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 954-957.	1.7	23
45	Effect of interleukin-1Î² gene functional polymorphism on dorsolateral prefrontal cortex activity in schizophrenic patients. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 1090-1093.	1.7	28
46	An Experimental Study of Catechol-O-Methyltransferase Val158Met Moderation of Î”-9-Tetrahydrocannabinol-Induced Effects on Psychosis and Cognition. Neuropsychopharmacology, 2006, 31, 2748-2757.	5.4	288
47	The Val66Met polymorphism of the brain-derived neurotrophic factor gene is associated with risk for psychosis: Evidence from a family-based association study. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2006, 141B, 135-138.	1.7	79
48	Ventricular enlargement in schizophrenia is associated with a genetic polymorphism at the interleukin-1 receptor antagonist gene. NeuroImage, 2005, 27, 1002-1006.	4.2	46
49	Dermatoglyphic anomalies and neurocognitive deficits in sibling pairs discordant for schizophrenia spectrum disorders. Psychiatry Research, 2005, 137, 215-221.	3.3	17
50	Interleukin-1 cluster is associated with genetic risk for schizophrenia and bipolar disorder. Journal of Medical Genetics, 2004, 41, 219-223.	3.2	67
51	Interleukinâ€1Î² (ILâ€1Î²) gene and increased risk for the depressive symptomâ€dimension in schizophrenia spectrum disorders. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2004, 124B, 10-14.	1.7	64
52	New Evidence of Association Between COMT Gene and Prefrontal Neurocognitive Function in Healthy Individuals From Sibling Pairs Discordant for Psychosis. American Journal of Psychiatry, 2004, 161, 1110-1112.	7.2	160
53	Neurocognitive, behavioural and neurodevelopmental correlates of schizotypy clusters in adolescents from the general population. Schizophrenia Research, 2003, 61, 293-302.	2.0	81
54	Nonreplication of the association between ab-ridge count and cerebral structural measures in schizophrenia. Comprehensive Psychiatry, 2003, 44, 459-461.	3.1	4

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55	Further Evidence That Congenital Dermatoglyphic Abnormalities Are Associated With Psychosis: A Twin Study. Schizophrenia Bulletin, 2002, 28, 697-701.	4.3	17
56	1q21â€q22 locus is associated with susceptibility to the realityâ€distortion syndrome of schizophrenia spectrum disorders. American Journal of Medical Genetics Part A, 2002, 114, 516-518.	2.4	26
57	Human genetic variation and mental disorders. Neurotoxicity Research, 2002, 4, 523-530.	2.7	5
58	Association study between novel promoter variants at the 5-HT2C receptor gene and human patients with bipolar affective disorder. Neuroscience Letters, 2001, 309, 135-137.	2.1	24
59	Dermatoglyphics and abnormal palmar flexion creases as markers of early prenatal stress in children with idiopathic intellectual disability. Journal of Intellectual Disability Research, 2001, 45, 416-423.	2.0	28
60	Congenital Dermatoglyphic Malformations and Psychosis: A Twin Study. American Journal of Psychiatry, 2000, 157, 1511-1513.	7.2	31
61	a-b ridge count and schizophrenia. Schizophrenia Research, 2000, 46, 285-286.	2.0	13
62	Developmental instability and schizotypy. Schizophrenia Research, 2000, 43, 125-134.	2.0	31
63	FC12.03 Recent dermatoglyphic studies in twin samples: Further evidences for an environmental risk factor in schizophrenia. European Psychiatry, 2000, 15, 305s-306s.	0.2	0
64	Variability in the serotonin transporter gene and increased risk for major depression with melancholia. Human Genetics, 1998, 103, 319-322.	3.8	92
65	Quantitative and qualitative palmar dermatoglyphics in the Mediterranean population of Delta de l'Ebre (Spain). International Journal of Anthropology, 1998, 13, 89-96.	0.1	0