Lourdes FañanÃ;s Saura

List of Publications by Year in descending order

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132 papers 5,522 citations

35 h-index 102487 66 g-index

145 all docs

145 docs citations

145 times ranked 6950 citing authors

#	Article	IF	CITATIONS
1	Childhood maltreatment disrupts HPA-axis activity under basal and stress conditions in a dose–response relationship in children and adolescents. Psychological Medicine, 2023, 53, 1060-1073.	4.5	14
2	<scp>ENIGMAâ€anxiety</scp> working group: Rationale for and organization of <scp>largeâ€scale</scp> neuroimaging studies of anxiety disorders. Human Brain Mapping, 2022, 43, 83-112.	3.6	31
3	Association and epistatic analysis of white matter related genes across the continuum schizophrenia and autism spectrum disorders: The joint effect of NRG1-ErbB genes. World Journal of Biological Psychiatry, 2022, 23, 208-218.	2.6	1
4	Polygenic contribution to the relationship of loneliness and social isolation with schizophrenia. Nature Communications, 2022, 13, 51.	12.8	16
5	Mapping genomic loci implicates genes and synaptic biology in schizophrenia. Nature, 2022, 604, 502-508.	27.8	929
6	Secretory immunoglobulin A (s-IgA) reactivity to acute psychosocial stress in children and adolescents: The influence of pubertal development and history of maltreatment. Brain, Behavior, and Immunity, 2022, 103, 122-129.	4.1	7
7	Prenatal exposures and behavioral epigenetics in human infants and children. , 2021, , 83-90.		1
8	COVID-19 una oportunidad $ ilde{A}^{o}$ nica para explorar la relaci $ ilde{A}^{3}$ n entre la infecci $ ilde{A}^{3}$ n prenatal materna, el desarrollo cerebral y los trastornos neuropsiqui $ ilde{A}$ ¡tricos en la descendencia. Revista De Psiquiatr $ ilde{A}$ a Y Salud Mental, 2021, 14, 1-3.	1.8	6
9	Prospective Long-Term Cohort Study of Subjects With First-Episode Psychosis Examining Eight Major Outcome Domains and Their Predictors: Study Protocol. Frontiers in Psychiatry, 2021, 12, 643112.	2.6	12
10	Monoamine oxidase A (MAOA) interaction with parenting practices on callous-unemotional traits in preschoolers. European Journal of Psychiatry, 2021, 35, 225-225.	1.3	0
11	COVID-19 una oportunidad única para explorar la relación entre la infección prenatal materna, el desarrollo cerebral y los trastornos neuropsiquiátricos en la descendencia. Revista De PsiquiatrÃa Y Salud Mental (English Edition), 2021, 14, 1-3.	0.3	O
12	Risk of Suicidal Behavior in Children and Adolescents Exposed to Maltreatment: The Mediating Role of Borderline Personality Traits and Recent Stressful Life Events. Journal of Clinical Medicine, 2021, 10, 5293.	2.4	13
13	Maltrato infantil y trastorno mental. Revista De PsiquiatrÃa Infanto-Juvenil, 2021, 38, 1-4.	0.3	O
14	Recent Stressful Life Events (SLE) and Adolescent Mental Health: Initial Validation of the LEIA, a New Checklist for SLE Assessment According to Their Severity, Interpersonal, and Dependent Nature. Assessment, 2020, 27, 1777-1795.	3.1	13
15	Utility of the MoCA for cognitive impairment screening in long-term psychosis patients. Schizophrenia Research, 2020, 216, 429-434.	2.0	20
16	Psychosocial stress and epigenetic aging. International Review of Neurobiology, 2020, 150, 107-128.	2.0	53
17	Familial aggregation analysis of cognitive performance in early-onset bipolar disorder. European Child and Adolescent Psychiatry, 2020, 29, 1705-1716.	4.7	3
18	Twin study designs as a tool to identify new candidate genes for depression: A systematic review of DNA methylation studies. Neuroscience and Biobehavioral Reviews, 2020, 112, 345-352.	6.1	11

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19	The black sheep of the family- whole-exome sequencing in family of lithium response discordant bipolar monozygotic twins. European Neuropsychopharmacology, 2020, 34, 19-27.	0.7	8
20	Violent aggression predicted by multiple pre-adult environmental hits. Molecular Psychiatry, 2019, 24, 1549-1564.	7.9	23
21	The interaction between the ZNF804A gene and cannabis use on the risk of psychosis in a non-clinical sample. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2019, 89, 174-180.	4.8	11
22	"A circle and a triangle dancing together― Alteration of social cognition in schizophrenia compared to autism spectrum disorders. Schizophrenia Research, 2019, 210, 94-100.	2.0	34
23	Prenatal adverse environment is associated with epigenetic age deceleration at birth and hypomethylation at the hypoxia-responsive EP300 gene. Clinical Epigenetics, 2019, 11, 73.	4.1	39
24	Transcriptomic metaanalyses of autistic brains reveals shared gene expression and biological pathway abnormalities with cancer. Molecular Autism, 2019, 10, 17.	4.9	30
25	Association of OXTR rs53576 with the Developmental Trajectories of Callous-Unemotional Traits and Stressful Life Events in 3- to 9-Year-Old Community Children. Journal of Abnormal Child Psychology, 2019, 47, 1651-1662.	3.5	7
26	Epigenetics-by-sex interaction for somatization conferred by methylation at the promoter region of SLC6A4 gene. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2019, 89, 125-131.	4.8	16
27	Diez años de investigación traslacional colaborativa en enfermedades mentales: el CIBERSAM. Revista De PsiquiatrÃa Y Salud Mental, 2019, 12, 1-8.	1.8	68
28	Increased methylation at an unexplored glucocorticoid responsive element within exon 1D of NR3C1 gene is related to anxious-depressive disorders and decreased hippocampal connectivity. European Neuropsychopharmacology, 2018, 28, 579-588.	0.7	44
29	The impact of prenatal insults on the human placental epigenome: A systematic review. Neurotoxicology and Teratology, 2018, 66, 80-93.	2.4	25
30	Cortisol, cortisone, and BDNF in amniotic fluid in the second trimester of pregnancy: Effect of early life and current maternal stress and socioeconomic status. Development and Psychopathology, 2018, 30, 971-980.	2.3	24
31	Dissecting the catatonia phenotype in psychotic and mood disorders on the basis of familial-genetic factors. Schizophrenia Research, 2018, 200, 20-25.	2.0	13
32	Genetic variability in scaffolding proteins and risk for schizophrenia and autism-spectrum disorders: a systematic review. Journal of Psychiatry and Neuroscience, 2018, 43, 223-244.	2.4	34
33	Epigenetic outlier profiles in depression: A genome-wide DNA methylation analysis of monozygotic twins. PLoS ONE, 2018, 13, e0207754.	2.5	14
34	Psychometric Properties of Drinking Motives Questionnaire-Revised (DMQ-R) in Spanish Adolescents. European Journal of Psychological Assessment, 2018, 34, 145-153.	3.0	12
35	FKBP5 modulates the hippocampal connectivity deficits in depression: a study in twins. Brain Imaging and Behavior, 2017, 11, 62-75.	2.1	34
36	Stressful life events during adolescence and risk for externalizing and internalizing psychopathology: a meta-analysis. European Child and Adolescent Psychiatry, 2017, 26, 1409-1422.	4.7	67

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37	An integrative review of methylation at the serotonin transporter gene and its dialogue with environmental risk factors, psychopathology and 5-HTTLPR. Neuroscience and Biobehavioral Reviews, 2017, 72, 190-209.	6.1	58
38	Cannabis use in male and female first episode of non-affective psychosis patients: Long-term clinical, neuropsychological and functional differences. PLoS ONE, 2017, 12, e0183613.	2.5	12
39	Environmental factors linked to depression vulnerability are associated with altered cerebellar resting-state synchronization. Scientific Reports, 2016, 6, 37384.	3.3	21
40	Familiality of Psychotic Disorders: A Polynosologic Study in Multiplex Families. Schizophrenia Bulletin, 2016, 42, 975-983.	4.3	23
41	Involvement of NRN1 gene in schizophrenia-spectrum and bipolar disorders and its impact on age at onset and cognitive functioning. World Journal of Biological Psychiatry, 2016, 17, 129-139.	2.6	18
42	Variations in Disrupted-in-Schizophrenia 1 gene modulate long-term longitudinal differences in cortical thickness in patients with a first-episode of psychosis. Brain Imaging and Behavior, 2016, 10, 629-635.	2.1	6
43	Altered amygdalar restingâ€state connectivity in depression is explained by both genes and environment. Human Brain Mapping, 2015, 36, 3761-3776.	3.6	8
44	Further Evidence of Depdc7 Dna Hypomethylation in Depression: a Study in Adult Twins. European Psychiatry, 2015, 30, 715-718.	0.2	14
45	Five-factor model and internalizing and externalizing syndromes: A 5-year prospective study. Personality and Individual Differences, 2015, 79, 98-103.	2.9	29
46	Season of birth and subclinical psychosis: Systematic review and meta-analysis of new and existing data. Psychiatry Research, 2015, 225, 227-235.	3.3	8
47	Desarrollo profesional en investigación traslacional en neurociencias y salud mental: educación y formación dentro del Centro de Investigación Biomédica en Red en Salud Mental. Revista De PsiquiatrÃa Y Salud Mental, 2015, 8, 65-74.	1.8	6
48	Polymorphic Variation in the Epigenetic Gene DNMT3B Modulates the Environmental Impact on Cognitive Ability: A Twin Study. European Psychiatry, 2015, 30, 303-308.	0.2	10
49	Glucocorticoid receptor gene (NR3C1) methylation processes as mediators of early adversity in stress-related disorders causality: A critical review. Neuroscience and Biobehavioral Reviews, 2015, 55, 520-535.	6.1	262
50	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
51	Brain structural correlates of schizotypy and psychosis proneness in a non-clinical healthy volunteer sample. Schizophrenia Research, 2015, 168, 37-43.	2.0	45
52	Progressive Structural Brain Changes and NRG1 Gene Variants in First-Episode Nonaffective Psychosis. Neuropsychobiology, 2015, 71, 103-111.	1.9	9
53	Maternal psychosocial stress during pregnancy alters the epigenetic signature of the glucocorticoid receptor gene promoter in their offspring: a meta-analysis. Epigenetics, 2015, 10, 893-902.	2.7	172
54	A cross-sectional and longitudinal structural magnetic resonance imaging study of the post-central gyrus in first-episode schizophrenia patients. Psychiatry Research - Neuroimaging, 2015, 231, 42-49.	1.8	16

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55	Two-year follow-up of treated adolescents with early-onset bipolar disorder: Changes in neurocognition. Journal of Affective Disorders, 2015, 172, 48-54.	4.1	22
56	Birth Weight and Adult IQ, but Not Anxious-Depressive Psychopathology, Are Associated with Cortical Surface Area: A Study in Twins. PLoS ONE, 2015, 10, e0129616.	2.5	6
57	Psychosisâ€inducing effects of cannabis are related to both childhood abuse and <scp>COMT</scp> genotypes. Acta Psychiatrica Scandinavica, 2014, 129, 54-62.	4.5	54
58	The BDNF-Val66Met polymorphism modulates parental rearing effects on adult psychiatric symptoms: A community twin-based study. European Psychiatry, 2014, 29, 293-300.	0.2	14
59	Family-based association study of common variants, rare mutation study and epistatic interaction detection in HDAC genes in schizophrenia. Schizophrenia Research, 2014, 160, 97-103.	2.0	23
60	DISC1-TSNAX and DAOA genes in major depression and citalopram efficacy. Journal of Affective Disorders, 2014, 168, 91-97.	4.1	15
61	Poster #M1 CHILDHOOD MALTREATMENT, THE BDNF-VAL66MET POLYMORPHISM AND HIPPOCAMPAL VOLUME: FURTHER EVIDENCES FROM A MRI-TWIN STUDY. Schizophrenia Research, 2014, 153, S189.	2.0	O
62	Cortical thickness correlates of psychotic experiences: Examining the effect of season of birth using a genetically informative design. Journal of Psychiatric Research, 2014, 56, 144-149.	3.1	7
63	Birth Weight, Working Memory and Epigenetic Signatures in IGF2 and Related Genes: A MZ Twin Study. PLoS ONE, 2014, 9, e103639.	2.5	14
64	A systematic review of the complex organization of human cognitive domains and their heritability. Psicothema, 2014, 26, 1-9.	0.9	37
65	Screening genetic variability at the CNR1 gene in both major depression etiology and clinical response to citalopram treatment. Psychopharmacology, 2013, 227, 509-519.	3.1	51
66	Twin-based study of the complex interplay between childhood maltreatment, socioeconomic status and adult memory. European Archives of Psychiatry and Clinical Neuroscience, 2013, 263, 435-440.	3.2	9
67	Association between symptomatic profile and remission following antidepressant treatment in unipolar major depression. Journal of Affective Disorders, 2013, 150, 209-215.	4.1	6
68	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
69	Gene–environment interaction on cognition: A twin study of childhood maltreatment and COMT variability. Journal of Psychiatric Research, 2013, 47, 989-994.	3.1	18
70	BDNF Val66Met variants and brain volume changes in non-affective psychosis patients and healthy controls: A 3year follow-up study. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2013, 45, 201-206.	4.8	6
71	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
72	Regional gray matter reductions are associated with genetic liability for anxiety and depression: An MRI twin study. Journal of Affective Disorders, 2013, 149, 175-181.	4.1	26

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73	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
74	TPH1, MAOA, Serotonin Receptor 2A and 2C Genes in Citalopram Response: Possible Effect in Melancholic and Psychotic Depression. Neuropsychobiology, 2013, 67, 41-47.	1.9	30
75	Genetic origin of the relationship between parental negativity and behavior problems from early childhood to adolescence: A longitudinal genetically sensitive study. Development and Psychopathology, 2013, 25, 487-500.	2.3	11
76	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
77	Increased familiarity of intellectual deficits in early-onset schizophrenia spectrum disorders. World Journal of Biological Psychiatry, 2012, 13, 493-500.	2.6	6
78	Genetic variability in the endocannabinoid system and 12-week clinical response to citalopram treatment: the role of the CNR1, CNR2 and FAAH genes. Journal of Psychopharmacology, 2012, 26, 1391-1398.	4.0	26
79	Poster #2 CHILDHOOD ADVERSITY AND CANNABIS USE IN THE DEVELOPMENT OF POSITIVE PSYCHOTIC-LIKE EXPERIENCES: MODERATION EFFECTS OF THE COMT GENE. Schizophrenia Research, 2012, 136, S91.	2.0	O
80	Effect of the Interleukin- $1\hat{l}^2$ Gene on Dorsolateral Prefrontal Cortex Function in Schizophrenia: A Genetic Neuroimaging Study. Biological Psychiatry, 2012, 72, 758-765.	1.3	28
81	Acquisition and generalization of fear conditioning are not modulated by the BDNFâ€val66met polymorphism in humans. Psychophysiology, 2012, 49, 713-719.	2.4	23
82	Convergent evidence of the contribution of TP53 genetic variation (Pro72Arg) to metabolic activity and white matter volume in the frontal lobe in schizophrenia patients. Neurolmage, 2011, 56, 45-51.	4.2	19
83	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
84	Neurodevelopmental liability to schizophrenia: The complex mediating role of age at onset and premorbid adjustment. Schizophrenia Research, 2011, 133, 143-149.	2.0	16
85	Early-onset bipolar disorder: how about visual-spatial skills and executive functions?. European Archives of Psychiatry and Clinical Neuroscience, 2011, 261, 195-203.	3.2	23
86	Changes in plasma and platelet BDNF levels induced by S-citalopram in major depression. Psychopharmacology, 2011, 216, 1-8.	3.1	58
87	Dysbindinâ€1 gene contributes differentially to early―and adult―nset forms of functional psychosis. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 322-333.	1.7	22
88	Childhood abuse, the BDNF-Val66Met polymorphism and adult psychotic-like experiences. British Journal of Psychiatry, 2011, 199, 38-42.	2.8	103
89	GENETIC VARIABILITY IN DYSBINDIN-1 GENE (DTNBP1) CONTRIBUTES DIFFERENTIALLY TO EARLY AND ADULT ONSET FUNCTIONAL PSYCHOSES AND IT IS ASSOCIATED WITH THE FAMILIAL TRANSMISSION OF IQ AND PREFRONTAL COGNITIVE DEFICITS. Schizophrenia Research, 2010, 117, 220-221.	2.0	O
90	NEUROLOGICAL ABNORMALITIES AND FLUCTUATING ASYMMETRY: THE ROLE OF PRENATAL ENVIRONMENT. Schizophrenia Research, 2010, 117, 320.	2.0	0

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91	Genetic polymorphisms in the dopamine-2 receptor (DRD2), dopamine-3 receptor (DRD3), and dopamine transporter (SLC6A3) genes in schizophrenia: Data from an association study. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2010, 34, 26-31.	4.8	37
92	Dysbindin gene (DTNBP1) in major depression: association with clinical response to selective serotonin reuptake inhibitors. Pharmacogenetics and Genomics, 2009, 19, 121-128.	1.5	24
93	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
94	Putative role of the COMT gene polymorphism (Val158Met) on verbal working memory functioning in a healthy population. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 898-902.	1.7	58
95	New evidences of gene and environment interactions affecting prenatal neurodevelopment in schizophrenia-spectrum disorders: A family dermatoglyphic study. Schizophrenia Research, 2008, 103, 209-217.	2.0	31
96	Study of Neurocognitive correlates of Schizotypy Personality Clusters in healthy individuals. European Journal of Psychiatry, 2008, 22, .	1.3	2
97	Working memory in siblings of schizophrenia patients. Schizophrenia Research, 2007, 95, 70-75.	2.0	51
98	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
99	Effect of interleukin- $1\hat{l}^2$ 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
100	Genetic variability at HPA axis in major depression and clinical response to antidepressant treatment. Journal of Affective Disorders, 2007, 104, 83-90.	4.1	165
101	An Experimental Study of Catechol-O-Methyltransferase Val158Met Moderation of Δ-9-Tetrahydrocannabinol-Induced Effects on Psychosis and Cognition. Neuropsychopharmacology, 2006, 31, 2748-2757.	5.4	288
102	Analysis of COMT gene (Val 158 Met polymorphism) in the clinical response to SSRIs in depressive patients of European origin. Journal of Affective Disorders, 2006, 90, 251-256.	4.1	93
103	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
104	Evidence for a combined genetic effect of the 5-HT1A receptor and serotonin transporter genes in the clinical outcome of major depressive patients treated with citalopram. Journal of Psychopharmacology, 2005, 19, 166-172.	4.0	88
105	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
106	Dermatoglyphic anomalies and neurocognitive deficits in sibling pairs discordant for schizophrenia spectrum disorders. Psychiatry Research, 2005, 137, 215-221.	3.3	17
107	Dermatoglyphics and Schizophrenia: A meta-analysis and investigation of the impact of obstetric complications upon a–b ridge count. Schizophrenia Research, 2005, 75, 399-404.	2.0	49
108	Interleukinâ€1β (<i>ILâ€1β</i>) 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.	ia 1.7	64

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109	Dermatoglyphic profile in 22q deletion syndrome. , 2004, 128B, 46-49.		7
110	Analysis of polymorphisms at the tumor suppressor gene p53 (TP53) in contributing to the risk for schizophrenia and its associated neurocognitive deficits. Neuroscience Letters, 2004, 363, 78-80.	2.1	19
111	Association analysis between a functional polymorphism in the monoamine oxidase A gene promoter and severe mood disorders. Psychiatric Genetics, 2004, 14, 203-208.	1.1	69
112	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
113	Directional and fluctuating asymmetry in finger and a-b ridge counts in psychosis: a case-control study. BMC Psychiatry, 2003, 3, 3.	2.6	33
114	Relapse of major depression after complete and partial remission during a 2-year follow-up. Journal of Affective Disorders, 2003, 73, 237-244.	4.1	88
115	Neurocognitive, behavioural and neurodevelopmental correlates of schizotypy clusters in adolescents from the general population. Schizophrenia Research, 2003, 61, 293-302.	2.0	81
116	Nonreplication of the association between ab-ridge count and cerebral structural measures in schizophrenia. Comprehensive Psychiatry, 2003, 44, 459-461.	3.1	4
117	5-HTTLPR Polymorphism of the Serotonin Transporter Gene Predicts Non-Remission in Major Depression Patients Treated With Citalopram in a 12-Weeks Follow Up Study. Journal of Clinical Psychopharmacology, 2003, 23, 563-567.	1.4	156
118	Further Evidence That Congenital Dermatoglyphic Abnormalities Are Associated With Psychosis: A Twin Study. Schizophrenia Bulletin, 2002, 28, 697-701.	4.3	17
119	Human genetic variation and mental disorders. Neurotoxicity Research, 2002, 4, 523-530.	2.7	5
120	The 5-HT2Areceptor gene 102T/C polymorphism is associated with suicidal behavior in depressed patients. American Journal of Medical Genetics Part A, 2001, 105, 801-804.	2.4	74
121	Georgian and Kurd mtDNA sequence analysis shows a lack of correlation between languages and female genetic lineages., 2000, 112, 5-16.		60
122	Congenital Dermatoglyphic Malformations and Psychosis: A Twin Study. American Journal of Psychiatry, 2000, 157, 1511-1513.	7.2	31
123	Association between cerebral structural abnormalities and dermatoglyphic ridge counts in schizophrenia. Comprehensive Psychiatry, 2000, 41, 380-384.	3.1	19
124	a-b ridge count and schizophrenia. Schizophrenia Research, 2000, 46, 285-286.	2.0	13
125	Negative dimension of schizotypy associated with early developmental instability in normal adolescents. Schizophrenia Research, 2000, 41, 84.	2.0	0
126	Developmental instability and schizotypy. Schizophrenia Research, 2000, 43, 125-134.	2.0	31

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127	Congenital dermatoglyphic malformations in severe bipolar disorder. Psychiatry Research, 1998, 78, 133-140.	3.3	36
128	Serotonin Transporter Gene and Risk for Bipolar Affective Disorder: An Association Study in a Spanish Population. Biological Psychiatry, 1998, 43, 843-847.	1.3	84
129	Dermatoglyphic abnormalities in psychosis: A twin study. Biological Psychiatry, 1997, 41, 624-626.	1.3	17
130	Allelic association analysis of the 5-HT2C receptor gene in bipolar affective disorder. Neuroscience Letters, 1996, 212, 65-67.	2.1	63
131	Dermatoglyphic a-b ridge count as a possible marker for developmental disturbance in schizophrenia: replication in two samples. Schizophrenia Research, 1996, 20, 307-314.	2.0	72
132	Parental age in schizophrenia in a case-controlled study. British Journal of Psychiatry, 1993, 162, 574-574.	2.8	18