Maria Pontillo

List of Publications by Citations

40 papers	707 citations	15 h-index	25 g-index
43 ext. papers	901 ext. citations	4.6 avg, IF	3.67 L-index

#	Paper	IF	Citations
40	Cognitive decline preceding the onset of psychosis in patients with 22q11.2 deletion syndrome. *JAMA Psychiatry*, **2015**, 72, 377-85	14.5	139
39	Adolescents at ultra-high risk for psychosis with and without 22q11 deletion syndrome: a comparison of prodromal psychotic symptoms and general functioning. *Schizophrenia Research*, **2012**, 139, 151-6	3.6	42
38	Ultra high risk status and transition to psychosis in 22q11.2 deletion syndrome. *World Psychiatry*, **2016**, 15, 259-265	14.4	41
37	Multiple stimulus presentation yields larger deficits in children with developmental dyslexia: a study with reading and RAN-type tasks. *Child Neuropsychology*, **2013**, 19, 639-47	2.7	37
36	Neurodevelopmental and psychiatric issues in Down s syndrome: assessment and intervention. *Psychiatric Genetics*, **2013**, 23, 95-107	2.9	37
35	Twelve-month psychosis-predictive value of the ultra-high risk criteria in children and adolescents. *Schizophrenia Research*, **2015**, 169, 186-192	3.6	35
34	Subthreshold Psychosis in 22q11.2 Deletion Syndrome: Multisite Naturalistic Study. *Schizophrenia Bulletin*, **2017**, 43, 1079-1089	1.3	32
33	Peer Victimization and Onset of Social Anxiety Disorder in Children and Adolescents. *Brain Sciences*, **2019**, 9,	3.4	28
32	Variations in Dysbindin-1 are associated with cognitive response to antipsychotic drug treatment. *Nature Communications*, **2018**, 9, 2265	17.4	28
31	Comorbid Personality Disorders in Individuals With an At-Risk Mental State for Psychosis: A Meta-Analytic Review. *Frontiers in Psychiatry*, **2019**, 10, 429	5	26
30	Adolescence is the starting point of sex-dichotomous COMT genetic effects. *Translational Psychiatry*, **2017**, 7, e1141	8.6	24
29	Complete Sequence of the 22q11.2 Allele in 1,053 Subjects with 22q11.2 Deletion Syndrome Reveals Modifiers of Conotruncal Heart Defects. *American Journal of Human Genetics*, **2020**, 106, 26-40	11	24
28	The eye-voice lead during oral reading in developmental dyslexia. *Frontiers in Human Neuroscience*, **2013**, 7, 696	3.3	22
27	Psychosocial interventions for very early and early-onset schizophrenia: a review of treatment efficacy. *Current Opinion in Psychiatry*, **2015**, 28, 312-23	4.9	19
26	Variance of IQ is partially dependent on deletion type among 1,427 22q11.2 deletion syndrome subjects. *American Journal of Medical Genetics, Part A*, **2018**, 176, 2172-2181	2.5	18
25	Is it still correct to differentiate between early and very early onset psychosis?. *Schizophrenia Research*, **2016**, 170, 211-6	3.6	15
24	Clinical presentation of Attenuated Psychosis Syndrome in children and adolescents: Is there an age effect?. *Psychiatry Research*, **2017**, 252, 169-174	9.9	14

23	Bridging the gap between different measures of the reading speed deficit in developmental dyslexia. *Experimental Brain Research*, **2014**, 232, 237-52	2.3	14
22	Use of Transcranial Direct Stimulation in the Treatment of Negative Symptoms of Schizophrenia. *Clinical EEG and Neuroscience*, **2018**, 49, 18-26	2.3	12
21	Attention deficit hyperactivity disorder symptoms as antecedents of later psychotic outcomes in 22q11.2 deletion syndrome. *Schizophrenia Research*, **2019**, 204, 320-325	3.6	11
20	No age effect in the prevalence and clinical significance of ultra-high risk symptoms and criteria for psychosis in 22q11 deletion syndrome: Confirmation of the genetically driven risk for psychosis?. *PLoS ONE*, **2017**, 12, e0174797	3.7	10
19	All that glitters is not gold: prevalence and relevance of psychotic-like experiences in clinical sample of children and adolescents aged 8-17 years old. *Microbial Biotechnology*, **2018**, 12, 702-707	3.3	10
18	Prevalence, course and psychosis-predictive value of negative symptoms in 22q11.2 deletion syndrome. *Schizophrenia Research*, **2019**, 206, 386-393	3.6	10
17	An attachment perspective on the risk for psychosis: Clinical correlates and the predictive value of attachment patterns and mentalization. *Schizophrenia Research*, **2020**, 222, 209-217	3.6	8
16	An Overview of Recent Findings on Social Anxiety Disorder in Adolescents and Young Adults at Clinical High Risk for Psychosis. *Brain Sciences*, **2017**, 7,	3.4	8
15	Developmental dyslexia in a regular orthography: can the reading profile be reduced to strategic control?. *Cognitive Neuropsychology*, **2013**, 30, 147-71	2.3	7
14	Indicated prevention with long-chain polyunsaturated omega-3 fatty acids in patients with 22q11DS genetically at high risk for psychosis. Protocol of a randomized, double-blind, placebo-controlled treatment trial. *Microbial Biotechnology*, **2016**, 10, 390-6	3.3	6
13	Prevalence and treatment of psychiatric disorders other than psychosis in children and adolescents with 22q11DS: Examining associations with social and role functioning. *Psychiatry Research*, **2017**, 254, 238-243	9.9	6
12	Failure to learn a new spatial format in children with developmental dyslexia. *Scientific Reports*, **2014**, 4, 4869	4.9	5
11	Neurocognitive profile and onset of psychosis symptoms in children, adolescents and young adults with 22q11 deletion syndrome: A longitudinal study. *Schizophrenia Research*, **2019**, 208, 76-81	3.6	4
10	Antipsychotics Do Not Influence Neurological Soft Signs in Children and Adolescents at Ultra-High Risk for Psychosis: A Pilot Study. *Journal of Psychiatric Practice*, **2019**, 25, 186-191	1.3	3
9	Clinical significance of family accommodation and parental psychological distress in a sample of children and adolescents with obsessive-compulsive disorder aged 8-17 years old. *Italian Journal of Pediatrics*, **2020**, 46, 167	3.2	2
8	Prevalence and Clinical Significance of Symptoms at Ultra High Risk for Psychosis in Children and Adolescents with Obsessive?Compulsive Disorder: Is There an Association with Global, Role, and Social Functioning?. *Brain Sciences*, **2018**, 8,	3.4	2
7	Visual perception skills: a comparison between patients with Noonan syndrome and 22q11.2 deletion syndrome. *Genes, Brain and Behavior*, **2017**, 16, 627-634	3.6	1
6	Clinical profile, conversion rate, and suicidal thinking and behaviour in children and adolescents at ultra-high risk for psychosis: a theoretical perspective. *Research in Psychotherapy: Psychopathology, Process and Outcome*, **2020**, 23, 455	0.9	1

5	Psychoeducation focused on family accommodation: a practical intervention for parents of children and adolescents with obsessive-compulsive disorder. *Italian Journal of Pediatrics*, **2021**, 47, 224	3.2	1
4	Personality Traits and Disorders in Adolescents at Clinical High Risk for Psychosis: Toward a Clinically Meaningful Diagnosis. *Frontiers in Psychiatry*, **2020**, 11, 562835	5	1
3	Dopamine dysfunction in 22q11 deletion syndrome: possible cause of motor symptoms. *Psychiatric Genetics*, **2016**, 26, 187-92	2.9	1
2	Schizofrenia ad esordio in età evolutiva: aspetti clinici e interventi possibili. *Quaderni Di Psicoterapia Cognitiva*, **2016**, 25-41	0.5	
1	22q11 microdeletion syndrome and ultra-high risk for psychosis: The role of neurological soft signs as an independent marker of vulnerability for psychosis. *Microbial Biotechnology*, **2019**, 13, 1191-1198	3.3	