Renato Borgatti

List of Publications by Year in descending order

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211 papers

7,036 citations

71102 41 h-index 70 g-index

217 all docs

217 docs citations

times ranked

217

8753 citing authors

#	Article	IF	CITATIONS
1	Prenatal maternal stress during the COVID-19 pandemic and infant regulatory capacity at 3 months: A longitudinal study. Development and Psychopathology, 2023, 35, 35-43.	2.3	60
2	Cognitive predictors of Social processing in congenital atypical development. Journal of Autism and Developmental Disorders, 2023, 53, 3343-3355.	2.7	2
3	Refining the mutational spectrum and gene–phenotype correlates in pontocerebellar hypoplasia: results of a multicentric study. Journal of Medical Genetics, 2022, 59, 399-409.	3.2	13
4	<i>SUFU</i> haploinsufficiency causes a recognisable neurodevelopmental phenotype at the mild end of the Joubert syndrome spectrum. Journal of Medical Genetics, 2022, 59, 888-894.	3.2	19
5	Impact of the inversion time on regional brain perfusion estimation with clinical arterial spin labeling protocols. Magnetic Resonance Materials in Physics, Biology, and Medicine, 2022, 35, 349-363.	2.0	3
6	Get Your Molar Tooth Right: Joubert Syndrome Misdiagnosis Unmasked by Whole-Exome Sequencing. Cerebellum, 2022, 21, 1144-1150.	2. 5	7
7	Cerebellar Agenesis. , 2022, , 2113-2134.		0
8	Perceived Family Functioning Profile in Adolescents at Clinical High Risk for Psychosis: Rigidity as a Possible Preventive Target. Frontiers in Psychiatry, 2022, 13, 861201.	2.6	1
9	Prognostic accuracy and clinical utility of psychometric instruments for individuals at clinical high-risk of psychosis: a systematic review and meta-analysis. Molecular Psychiatry, 2022, 27, 3670-3678.	7.9	13
10	Superior Cerebellar Atrophy: An Imaging Clue to Diagnose ITPR1-Related Disorders. International Journal of Molecular Sciences, 2022, 23, 6723.	4.1	4
11	Family Dysfunctional Interactive Patterns and Alexithymia in Adolescent Patients with Restrictive Eating Disorders. Children, 2022, 9, 1038.	1.5	3
12	Heterozygous <i>KIF1A</i> variants underlie a wide spectrum of neurodevelopmental and neurodegenerative disorders. Journal of Medical Genetics, 2021, 58, 475-483.	3.2	21
13	Early parenting intervention promotes 24â€month psychomotor development in preterm children. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 101-108.	1.5	6
14	Basal Ganglia Dysmorphism in Patients With Aicardi Syndrome. Neurology, 2021, 96, e1319-e1333.	1.1	6
15	Rehabilitation services lockdown during the COVID-19 emergency: the mental health response of caregivers of children with neurodevelopmental disabilities. Disability and Rehabilitation, 2021, 43, 27-32.	1.8	48
16	Italian parents welcomed a telehealth familyâ€centred rehabilitation programme for children with disability during COVIDâ€19 lockdown. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 194-196.	1.5	39
17	Potentials of Telerehabilitation for Families of Children With Special Health Care Needs During the Coronavirus Disease 2019 Emergency. JAMA Pediatrics, 2021, 175, 105.	6.2	12
18	How social is the cerebellum? Exploring the effects of cerebellar transcranial direct current stimulation on the prediction of social and physical events. Brain Structure and Function, 2021, 226, 671-684.	2.3	26

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19	Phonological memory updating and developmental dyslexia: The role of long-term knowledge. Child Neuropsychology, 2021, 27, 718-733.	1.3	3
20	Self-Report and Biological Indexes of Work-Related Stress in Neonatal Healthcare Professionals. Advances in Neonatal Care, 2021, 21, E120-E128.	1.1	4
21	CASK related disorder: Epilepsy and developmental outcome. European Journal of Paediatric Neurology, 2021, 31, 61-69.	1.6	7
22	COVIDâ€19â€related psychiatric impact on Italian adolescent population: A crossâ€sectional cohort study. Journal of Community Psychology, 2021, 49, 1457-1469.	1.8	22
23	Rating behavioral problems in adolescent eating disorders: Parent-child differences. Psychiatry Research, 2021, 298, 113836.	3.3	1
24	The Experience of Child Neuropsychiatry Residents who Volunteered in Italian COVID-19-Designated Hospitals. Academic Psychiatry, 2021, 45, 587-592.	0.9	3
25	Challenges and resources in adult life with Joubert syndrome: issues from an international classification of functioning (ICF) perspective. Disability and Rehabilitation, 2021, , 1-8.	1.8	1
26	Diagnostic Yield and Cost-Effectiveness of "Dynamic―Exome Analysis in Epilepsy with Neurodevelopmental Disorders: A Tertiary-Center Experience in Northern Italy. Diagnostics, 2021, 11, 948.	2.6	6
27	Prognostic Accuracy of DSM-5 Attenuated Psychosis Syndrome in Adolescents: Prospective Real-World 5-Year Cohort Study. Schizophrenia Bulletin, 2021, 47, 1663-1673.	4.3	12
28	Reading Skills of Children with Dyslexia Improved Less Than Expected during the COVID-19 Lockdown in Italy. Children, 2021, 8, 560.	1.5	18
29	Ketogenic Dietary Therapies in Patients with Autism Spectrum Disorder: Facts or Fads? A Scoping Review and a Proposal for a Shared Protocol. Nutrients, 2021, 13, 2057.	4.1	16
30	Lactobacillus plantarum PS128 and Other Probiotics in Children and Adolescents with Autism Spectrum Disorder: A Real-World Experience. Nutrients, 2021, 13, 2036.	4.1	16
31	Assessing Family Functioning Before and After an Integrated Multidisciplinary Family Treatment for Adolescents With Restrictive Eating Disorders. Frontiers in Psychiatry, 2021, 12, 653047.	2.6	7
32	Longitudinal outcome of attenuated positive symptoms, negative symptoms, functioning and remission in people at clinical high risk for psychosis: a meta-analysis. EClinicalMedicine, 2021, 36, 100909.	7.1	25
33	Hidden pandemic: COVID-19-related stress, SLC6A4 methylation, and infants' temperament at 3Âmonths. Scientific Reports, 2021, 11, 15658.	3.3	32
34	Effectiveness of Short-Term Psychodynamic Psychotherapy in Preadolescents and Adolescents Affected by Psychiatric Disorders. Psychiatry Investigation, 2021, 18, 923-927.	1.6	1
35	Social prediction in pediatric patients with congenital, non-progressive malformations of the cerebellum: From deficits in predicting movements to rehabilitation in virtual reality. Cortex, 2021, 144, 82-98.	2.4	8
36	Depression and Anxiety in Mothers Who Were Pregnant During the COVID-19 Outbreak in Northern Italy: The Role of Pandemic-Related Emotional Stress and Perceived Social Support. Frontiers in Psychiatry, 2021, 12, 716488.	2.6	34

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37	ITAlian Partnership for Psychosis Prevention (ITAPP): improving the mental health of young people. European Psychiatry, 2021, 64, 1-21.	0.2	4
38	Longâ€term followâ€up in a cohort of children with isolated corpus callosum agenesis at fetal MRI. Annals of Clinical and Translational Neurology, 2021, 8, 2280-2288.	3.7	4
39	Supporting Parenting at Home-Empowering Rehabilitation through Engagement (SPHERE): study protocol for a randomised control trial. BMJ Open, 2021, 11, e051817.	1.9	1
40	Impaired urinary concentration ability is a sensitive predictor of renal disease progression in Joubert syndrome. Nephrology Dialysis Transplantation, 2020, 35, 1195-1202.	0.7	15
41	Painâ€related increase in serotonin transporter gene methylation associates with emotional regulation in 4.5â€yearâ€old pretermâ€born children. Acta Paediatrica, International Journal of Paediatrics, 2020, 109, 1166-1174.	1.5	23
42	Maternal caregiving and DNA methylation in human infants and children: Systematic review. Genes, Brain and Behavior, 2020, 19, e12616.	2.2	44
43	The Little Professor and the Virus: Scaffolding Children's Meaning Making During the COVID-19 Emergency. Frontiers in Psychiatry, 2020, 11, 817.	2.6	15
44	Premature birth affects visual body representation and body schema in preterm children. Brain and Cognition, 2020, 145, 105612.	1.8	7
45	Cerebellar Damage Affects Contextual Priors for Action Prediction in Patients with Childhood Brain Tumor. Cerebellum, 2020, 19, 799-811.	2.5	12
46	Characterizing White Matter Tract Organization in Polymicrogyria and Lissencephaly: A Multifiber Diffusion MRI Modeling and Tractography Study. American Journal of Neuroradiology, 2020, 41, 1495-1502.	2.4	9
47	Migraine Symptoms Improvement During the COVID-19 Lockdown in a Cohort of Children and Adolescents. Frontiers in Neurology, 2020, 11, 579047.	2.4	33
48	Alone With the Kids: Tele-Medicine for Children With Special Healthcare Needs During COVID-19 Emergency. Frontiers in Psychology, 2020, 11, 2193.	2.1	25
49	Clinical features of adolescents diagnosed with eating disorders and at risk for psychosis. European Psychiatry, 2020, 63, e80.	0.2	5
50	Age and sex prevalence estimate of Joubert syndrome in Italy. Neurology, 2020, 94, e797-e801.	1.1	26
51	Effectiveness of Computerized Cognitive Training Programs (CCTP) with Game-like Features in Children with or without Neuropsychological Disorders: a Meta-Analytic Investigation. Neuropsychology Review, 2020, 30, 126-141.	4.9	18
52	Virtual Reality Social Prediction Improvement and Rehabilitation Intensive Training (VR-SPIRIT) for paediatric patients with congenital cerebellar diseases: study protocol of a randomised controlled trial. Trials, 2020, 21, 82.	1.6	16
53	Home-based cognitive training in pediatric patients with acquired brain injury: preliminary results on efficacy of a randomized clinical trial. Scientific Reports, 2020, 10, 1391.	3.3	22
54	Measuring the Outcomes of Maternal COVID-19-related Prenatal Exposure (MOM-COPE): study protocol for a multicentric longitudinal project. BMJ Open, 2020, 10, e044585.	1.9	22

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55	Finding a common path to the assessment of persons with intellectual development disorders. Neurology, 2020, 94, 507-508.	1.1	2
56	Cerebellar Agenesis. , 2020, , 1-23.		0
57	The spectrum of brainstem malformations associated to mutations of the tubulin genes family: MRI and DTI analysis. European Radiology, 2019, 29, 770-782.	4.5	22
58	Epilepsy in Tubulinopathy: Personal Series and Literature Review. Cells, 2019, 8, 669.	4.1	27
59	Exploring the EEG mu rhythm associated with observation and execution of a goal-directed action in 14-month-old preterm infants. Scientific Reports, 2019, 9, 8975.	3.3	12
60	Early Brain Damage Affects Body Schema and Person Perception Abilities in Children and Adolescents with Spastic Diplegia. Neural Plasticity, 2019, 2019, 1-17.	2.2	13
61	Remote Technology-Based Training Programs for Children with Acquired Brain Injury: A Systematic Review and a Meta-Analytic Exploration. Behavioural Neurology, 2019, 2019, 1-31.	2.1	29
62	Telomere length and salivary cortisol stress reactivity in very preterm infants. Early Human Development, 2019, 129, 1-4.	1.8	13
63	Non-invasive Brain Stimulation for the Rehabilitation of Children and Adolescents With Neurodevelopmental Disorders: A Systematic Review. Frontiers in Psychology, 2019, 10, 135.	2.1	63
64	Functional Evaluation of Eating Difficulties Scale to predict oral motor skills in infants with neurodevelopmental disorders: a longitudinal study. Developmental Medicine and Child Neurology, 2019, 61, 813-819.	2.1	0
65	The phenotypic spectrum of WWOX-related disorders: 20 additional cases of WOREE syndrome and review of the literature. Genetics in Medicine, 2019, 21, 1308-1318.	2.4	48
66	The dual nature of hypothalamic-pituitary-adrenal axis regulation in dyads of very preterm infants and their mothers. Psychoneuroendocrinology, 2019, 100, 172-179.	2.7	23
67	Pain exposure associates with telomere length erosion in very preterm infants. Psychoneuroendocrinology, 2018, 89, 113-119.	2.7	15
68	Predictors and outcomes of the Neonatal Oral Motor Assessment Scale (NOMAS) performance: a systematic review. European Journal of Pediatrics, 2018, 177, 665-673.	2.7	10
69	Developmental care, neonatal behavior and postnatal maternal depressive symptomatology predict internalizing problems at 18 months for very preterm children. Journal of Perinatology, 2018, 38, 191-195.	2.0	20
70	Methodological Challenges in Developmental Human Behavioral Epigenetics: Insights Into Study Design. Frontiers in Behavioral Neuroscience, 2018, 12, 286.	2.0	9
71	Cognitive functioning of pediatric patients with brain tumor: an investigation of the role of gender. Child's Nervous System, 2018, 34, 2415-2423.	1.1	11
72	Insights from perceptual, sensory, and motor functioning in autism and cerebellar primary disturbances: Are there reliable markers for these disorders?. Neuroscience and Biobehavioral Reviews, 2018, 95, 263-279.	6.1	14

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73	A Different Brain: Anomalies of Functional and Structural Connections in Williams Syndrome. Frontiers in Neurology, 2018, 9, 721.	2.4	10
74	Visual perception and spatial transformation of the body in children and adolescents with brain tumor. Neuropsychologia, 2018, 120, 124-136.	1.6	10
75	Maternal sensitivity is associated with configural processing of infant's cues in preterm and full-term mothers. Early Human Development, 2018, 125, 35-45.	1.8	20
76	Mutations in the Neuroblastoma Amplified Sequence gene in a family affected by Acrofrontofacionasal Dysostosis type 1. Bone, 2018, 114, 125-136.	2.9	24
77	Tubulin genes and malformations of cortical development. European Journal of Medical Genetics, 2018, 61, 744-754.	1.3	93
78	Very preterm birth is associated with <i>PLAGL1</i> gene hypomethylation at birth and discharge. Epigenomics, 2018, 10, 1121-1130.	2.1	7
79	Language and the cerebellum. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 154, 181-202.	1.8	89
80	From early stress to 12-month development in very preterm infants: Preliminary findings on epigenetic mechanisms and brain growth. PLoS ONE, 2018, 13, e0190602.	2.5	60
81	Feasibility of a home-based computerized cognitive training for pediatric patients with congenital or acquired brain damage: An explorative study. PLoS ONE, 2018, 13, e0199001.	2.5	22
82	When one is Enough: Impaired Multisensory Integration in Cerebellar Agenesis. Cerebral Cortex, 2017, 27, bhw049.	2.9	37
83	Promoting Neuroprotective Care in Neonatal Intensive Care Units and Preterm Infant Development: Insights From the Neonatal Adequate Care for Quality of Life Study. Child Development Perspectives, 2017, 11, 9-15.	3.9	29
84	Very Preterm and Fullâ€Term Infants' Response to Socioâ€Emotional Stress: The Role of Postnatal Maternal Bonding. Infancy, 2017, 22, 695-712.	1.6	37
85	Greater brain response to emotional expressions of their own children in mothers of preterm infants: an fMRI study. Journal of Perinatology, 2017, 37, 716-722.	2.0	11
86	Role of the cerebellum in high stages of motor planning hierarchy. Journal of Neurophysiology, 2017, 117, 1474-1482.	1.8	10
87	Why Are Prospective Longitudinal Studies Needed in Preterm Behavioral Epigenetic Research?. JAMA Pediatrics, 2017, 171, 92.	6.2	9
88	WD40-repeat 47, a microtubule-associated protein, is essential for brain development and autophagy. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E9308-E9317.	7.1	77
89	Anterior Mesencephalic Cap Dysplasia: Novel Brain Stem Malformative Features Associated with Joubert Syndrome. American Journal of Neuroradiology, 2017, 38, 2385-2390.	2.4	15
90	Paediatricians should encourage the parents of children with special healthcare needs to disclose their use of complementary and alternative medicine. Acta Paediatrica, International Journal of Paediatrics, 2017, 106, 1883-1884.	1.5	0

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91	Tubulin-related cerebellar dysplasia: definition of a distinct pattern of cerebellar malformation. European Radiology, 2017, 27, 5080-5092.	4.5	36
92	Clinical Characterization, Genetics, and Long-Term Follow-up of a Large Cohort of Patients With Agenesis of the Corpus Callosum. Journal of Child Neurology, 2017, 32, 60-71.	1.4	34
93	Maternal Sensitivity Buffers the Association between SLC6A4 Methylation and Socio-Emotional Stress Response in 3-Month-Old Full Term, but not very Preterm Infants. Frontiers in Psychiatry, 2017, 8, 171.	2.6	28
94	The Functional Evaluation of Eating Difficulties Scale: Study Protocol and Validation in Infants with Neurodevelopmental Impairments and Disabilities. Frontiers in Pediatrics, 2017, 5, 273.	1.9	2
95	Learning and Using Abstract Words: Evidence from Clinical Populations. BioMed Research International, 2017, 2017, 1-8.	1.9	2
96	Relationship Between Maternal Sensitivity During Early Interaction and Maternal Ability in Perceiving Infants' Body and Face. Infancy, 2016, 21, 582-602.	1.6	8
97	COMT vall 58met polymorphism is associated with behavioral response and physiologic reactivity to socio-emotional stress in 4-month-old infants., 2016, 45, 71-82.		16
98	An Automated Function for Identifying EEG Independent Components Representing Bilateral Source Activity. IFMBE Proceedings, 2016, , 105-109.	0.3	22
99	Clinical, neuroradiological and molecular characterization of cerebellar dysplasia with cysts (Poretti–Boltshauser syndrome). European Journal of Human Genetics, 2016, 24, 1262-1267.	2.8	43
100	Language outcomes at 36 months in prematurely born children is associated with the quality of developmental care in NICUs. Journal of Perinatology, 2016, 36, 768-774.	2.0	14
101	Does quality of developmental care in NICUs affect health-related quality of life in 5-y-old children born preterm?. Pediatric Research, 2016, 80, 824-828.	2.3	12
102	Serotonin Transporter Gene (<i>SLC6A4</i>) Methylation Associates With NeonatalÂIntensive Care Unit Stay and 3â€Monthâ€Old Temperament in Preterm Infants. Child Development, 2016, 87, 38-48.	3.0	73
103	<i>SLC6A4</i> promoter region methylation and socio-emotional stress response in very preterm and full-term infants. Epigenomics, 2016, 8, 895-907.	2.1	37
104	Pain-related stress in the Neonatal Intensive Care Unit and salivary cortisol reactivity to socio-emotional stress in 3-month-old very preterm infants. Psychoneuroendocrinology, 2016, 72, 161-165.	2.7	67
105	Neonatal developmental care in infant pain management and internalizing behaviours at 18Âmonths in prematurely born children. European Journal of Pain, 2016, 20, 1010-1021.	2.8	31
106	Aberrant supracallosal longitudinal bundle: MR features, pathogenesis and associated clinical phenotype. European Radiology, 2016, 26, 2587-2596.	4.5	7
107	Complementary and alternative medicine (CAM) for children with special health care needs: A comparative usage study in Italy. European Journal of Integrative Medicine, 2016, 8, 115-121.	1.7	8
108	The Light Side of Preterm Behavioral Epigenetics. Advances in Medical Diagnosis, Treatment, and Care, 2016, , 107-127.	0.1	1

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109	Clinical profile of patients with ATP1A3 mutations in Alternating Hemiplegia of Childhoodâ€"a study of 155 patients. Orphanet Journal of Rare Diseases, 2015, 10, 123.	2.7	117
110	Learning to live without the cerebellum. NeuroReport, 2015, 26, 809-813.	1.2	17
111	Pain-related stress during the Neonatal Intensive Care Unit stay and SLC6A4 methylation in very preterm infants. Frontiers in Behavioral Neuroscience, 2015, 9, 99.	2.0	78
112	Functional genome-wide siRNA screen identifies KIAA0586 as mutated in Joubert syndrome. ELife, 2015, 4, e06602.	6.0	64
113	Social stress regulation in 4-month-old infants: Contribution of maternal social engagement and infants' 5-HTTLPR genotype. Early Human Development, 2015, 91, 173-179.	1.8	36
114	Oral-facial-digital syndrome type VI: is C5orf42 really the major gene?. Human Genetics, 2015, 134, 123-126.	3.8	30
115	A categorical approach to infants' individual differences during the Still-Face paradigm. , 2015, 38, 67-76.		29
116	Mother–infant dyadic reparation and individual differences in vagal tone affect 4-month-old infants' social stress regulation. Journal of Experimental Child Psychology, 2015, 140, 158-170.	1.4	34
117	Intersensory redundancy promotes visual rhythm discrimination in visually impaired infants., 2015, 39, 92-97.		1
118	A de-novo STXBP1 gene mutation in a patient showing the Rett syndrome phenotype. NeuroReport, 2015, 26, 254-257.	1.2	39
119	Facial emotion recognition in Williams syndrome and Down syndrome: A matching and developmental study. Child Neuropsychology, 2015, 21, 668-692.	1.3	24
120	Paediatric Biobanks: Opinions, Feelings and Attitudes of Parents towards the Specimen Donation of Their Sick Children to a Hypothetical Biobank. Pathobiology, 2015, 81, 304-308.	3.8	9
121	Neurophysiological and clinical findings on Nodding Syndrome in 21 South Sudanese children and a review of the literature. Seizure: the Journal of the British Epilepsy Association, 2015, 31, 64-71.	2.0	23
122	Neuropsychological assessment of children with epilepsy and average intelligence using NEPSY II. Journal of Clinical and Experimental Neuropsychology, 2015, 37, 1036-1051.	1.3	17
123	Cognitive rehabilitation in a child with Joubert Syndrome: Developmental trends and adaptive changes in a single case report. Research in Developmental Disabilities, 2015, 47, 375-384.	2.2	15
124	A dynamic system analysis of dyadic flexibility and stability across the Face-to-Face Still-Face procedure: Application of the State Space Grid., 2015, 38, 1-10.		34
125	Mutations in \hat{I}_{\pm} - and \hat{I}^2 -tubulin encoding genes: Implications in brain malformations. Brain and Development, 2015, 37, 273-280.	1.1	94
126	Vagal tone as a biomarker of longâ€term memory for a stressful social event at 4 months. Developmental Psychobiology, 2014, 56, 1564-1574.	1.6	18

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127	Maternal stress and depressive symptoms associated with quality of developmental care in 25 Italian Neonatal Intensive Care Units: A cross sectional observational study. International Journal of Nursing Studies, 2014, 51, 994-1002.	5.6	32
128	A Novel Mutation in <i>STXBP1</i> Gene in a Child With Epileptic Encephalopathy and an Atypical Electroclinical Pattern. Journal of Child Neurology, 2014, 29, 249-253.	1.4	18
129	Brain malformations and mutations in $\langle i \rangle \hat{l}_{\pm} \langle i \rangle \hat{a} \in \mathbb{R}$ and $\langle i \rangle \hat{l}_{\pm} \langle i \rangle \hat{a} \in \mathbb{R}$ ubulin genes: a review of the literature and description of two new cases. Developmental Medicine and Child Neurology, 2014, 56, 354-360.	2.1	42
130	Novel SETX variants in a patient with ataxia, neuropathy, and oculomotor apraxia are associated with normal sensitivity to oxidative DNA damaging agents. Brain and Development, 2014, 36, 682-689.	1.1	8
131	Sequence memory skills in Spastic Bilateral Cerebral Palsy are age independent as in normally developing children. Disability and Rehabilitation, 2013, 35, 506-512.	1.8	13
132	Cerebroretinal Microangiopathy With Calcifications and Cysts Associated With <i>CTC1</i> and <i>NDP</i> Mutations. Journal of Child Neurology, 2013, 28, 1702-1708.	1.4	11
133	Phenotypic spectrum and prevalence of INPP5E mutations in Joubert Syndrome and related disorders. European Journal of Human Genetics, 2013, 21, 1074-1078.	2.8	64
134	Four-Month-Old Infants' Long-Term Memory for a Stressful Social Event. PLoS ONE, 2013, 8, e82277.	2.5	22
135	Cerebellar Agenesis. , 2013, , 1855-1872.		4
136	The Role of Negative Maternal Affective States and Infant Temperament in Early Interactions Between Infants With Cleft Lip and Their Mothers. Journal of Pediatric Psychology, 2012, 37, 241-250.	2.1	25
137	Level of NICU Quality of Developmental Care and Neurobehavioral Performance in Very Preterm Infants. Pediatrics, 2012, 129, e1129-e1137.	2.1	148
138	Pediatric Biobanking: A Pilot Qualitative Survey of Practices, Rules, and Researcher Opinions in Ten European Countries. Biopreservation and Biobanking, 2012, 10, 29-36.	1.0	22
139	Differential distribution and lateralization of infant gestures and their relation to maternal gestures in the Face-to-Face Still-Face paradigm., 2012, 35, 819-828.		15
140	A novel mutation in the βâ€tubulin gene <i>TUBB2B</i> associated with complex malformation of cortical development and deficits in axonal guidance. Developmental Medicine and Child Neurology, 2012, 54, 765-769.	2.1	50
141	Seizures and EEG patterns in Pallister–Killian syndrome: 13ÂNew Italian patients. European Journal of Paediatric Neurology, 2012, 16, 636-641.	1.6	13
142	Electroclinical pattern in <i>MECP2</i> duplication syndrome: Eight new reported cases and review of literature. Epilepsia, 2012, 53, 1146-1155.	5.1	34
143	Measuring maternal stress and perceived support in 25 Italian NICUs. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 136-142.	1.5	97
144	Delineation and Diagnostic Criteria of Oral-Facial-Digital Syndrome Type VI. Orphanet Journal of Rare Diseases, 2012, 7, 4.	2.7	64

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145	Sequence Learning in Cerebral Palsy. Pediatric Neurology, 2011, 44, 207-213.	2.1	27
146	A Smile Enhances 3â€Monthâ€Olds' Recognition of an Individual Face. Infancy, 2011, 16, 306-317.	1.6	17
147	Duplications of FOXG1 in 14q12 are associated with developmental epilepsy, mental retardation, and severe speech impairment. European Journal of Human Genetics, 2011, 19, 102-107.	2.8	104
148	Behavioural features of Italian infants and young adults with Williams–Beuren syndrome. Journal of Intellectual Disability Research, 2011, 55, 121-131.	2.0	10
149	Studying cross-cultural differences in temperament in the first year of life: United States and Italy. International Journal of Behavioral Development, 2011, 35, 27-37.	2.4	48
150	Molecular Mechanisms Generating and Stabilizing Terminal 22q13 Deletions in 44 Subjects with Phelan/McDermid Syndrome. PLoS Genetics, 2011, 7, e1002173.	3.5	172
151	A comparison of dyadic interactions and coping with stillâ€face in healthy preâ€ŧerm and fullâ€ŧerm infants. British Journal of Developmental Psychology, 2010, 28, 347-368.	1.7	82
152	Neurological soft signs feature a double dissociation within the language system in Williams syndrome. Neuropsychologia, 2010, 48, 3298-3304.	1.6	7
153	Infant's emotional variability associated to interactive stressful situation: A novel analysis approach with Sample Entropy and Lempel–Ziv Complexity. , 2010, 33, 346-356.		14
154	Novel <i>TMEM67</i> mutations and genotype-phenotype correlates in meckelin-related ciliopathies. Human Mutation, 2010, 31, n/a-n/a.	2.5	77
155	Genotype–phenotype relationship in three cases with overlapping 19p13.12 microdeletions. European Journal of Human Genetics, 2010, 18, 1302-1309.	2.8	46
156	The Development of Dynamic Facial Expression Recognition at Different Intensities in 4- to 18-Year-Olds. Social Development, 2010, 19, 71-92.	1.3	132
157	Novel spliceâ€site mutations and a large intragenic deletion in <i>PLA2G6</i> associated with a severe and rapidly progressive form of infantile neuroaxonal dystrophy. Clinical Genetics, 2010, 78, 432-440.	2.0	30
158	Lateral asymmetries in infants' regulatory and communicative gestures. , 2010, , 103-111.		3
159	A wide spectrum of clinical, neurophysiological and neuroradiological abnormalities in a family with a novel CACNA1A mutation. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 840-843.	1.9	51
160	Narrative language in Williams Syndrome and its neuropsychological correlates. Journal of Neurolinguistics, 2010, 23, 97-111.	1,1	42
161	Evidence for a link among cognition, language and emotion in cerebellar malformations. Cortex, 2010, 46, 907-918.	2.4	64
162	Neuropsychological and behavioural aspects in children and adolescents with idiopathic epilepsy at diagnosis and after 12 months of treatment. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 540-546.	2.0	38

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163	International Classification of Functioning, Disability and Health in children with congenital central hypoventilation syndrome. Disability and Rehabilitation, 2009, 31, S144-S152.	1.8	17
164	International Classification of Functioning, Disability and Health in subjects with alternating hemiplegia of childhood. Disability and Rehabilitation, 2009, 31, S108-S115.	1.8	8
165	Expanding <i>CEP290</i> mutational spectrum in ciliopathies. American Journal of Medical Genetics, Part A, 2009, 149A, 2173-2180.	1.2	38
166	<i>MKS3/TMEM67</i> mutations are a major cause of COACH Syndrome, a Joubert Syndrome related disorder with liver involvement. Human Mutation, 2009, 30, E432-E442.	2.5	96
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