

Bruna Scalia

List of Publications by Year in descending order

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Version: 2024-02-01

72
papers

937
citations

516215

16
h-index

610482

24
g-index

73
all docs

73
docs citations

73
times ranked

1226
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Ataxia in children: early recognition and clinical evaluation. Italian Journal of Pediatrics, 2017, 43, 6. | 1.0 | 59 |
| 2 | Paediatric anti-N-methyl-d-aspartate receptor encephalitis: The first Italian multicenter case series. European Journal of Paediatric Neurology, 2015, 19, 453-463. | 0.7 | 56 |
| 3 | Inherited human IRAK-1 deficiency selectively impairs TLR signaling in fibroblasts. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E514-E523. | 3.3 | 49 |
| 4 | Targeting inflammation as a therapeutic strategy for drug-resistant epilepsies. Human Vaccines and Immunotherapeutics, 2014, 10, 868-875. | 1.4 | 47 |
| 5 | Ketogenic diet for infants with epilepsy: A literature review. Epilepsy and Behavior, 2020, 112, 107361. | 0.9 | 46 |
| 6 | Seizures in the neonate: A review of etiologies and outcomes. Seizure: the Journal of the British Epilepsy Association, 2021, 85, 48-56. | 0.9 | 39 |
| 7 | Current status of laboratory and imaging diagnosis of neonatal necrotizing enterocolitis. Italian Journal of Pediatrics, 2018, 44, 84. | 1.0 | 37 |
| 8 | Randomized Comparison of Helmet CPAP Versus High-Flow Nasal Cannula Oxygen in Pediatric Respiratory Distress. Respiratory Care, 2017, 62, 1036-1042. | 0.8 | 35 |
| 9 | Levetiracetam in neonatal seizures as first-line treatment: A prospective study. Journal of Pediatric Neurosciences, 2017, 12, 24. | 0.2 | 33 |
| 10 | Prognostic Challenges of SCN1A Genetic Mutations: Report on Two Children with Mild Features. Journal of Pediatric Neurology, 2016, 14, 082-088. | 0.0 | 32 |
| 11 | The usefulness of immunotherapy in pediatric neurodegenerative disorders: A systematic review of literature data. Human Vaccines and Immunotherapeutics, 2015, 11, 2749-2763. | 1.4 | 28 |
| 12 | Symptomatic seizures in preterm newborns: a review on clinical features and prognosis. Italian Journal of Pediatrics, 2018, 44, 115. | 1.0 | 27 |
| 13 | Preterm birth: seven-year retrospective study in a single centre population. Italian Journal of Pediatrics, 2019, 45, 45. | 1.0 | 25 |
| 14 | Mental Retardation and Early Onset of Weakness in a Girl With a Dystrophinopathy and a Large Xp21-23 Deletion. Journal of Child Neurology, 2003, 18, 79-81. | 0.7 | 20 |
| 15 | Short-Term Neurodevelopmental Outcome in Term Neonates Treated with Phenobarbital versus Levetiracetam: A Single-Center Experience. Behavioural Neurology, 2019, 2019, 1-8. | 1.1 | 20 |
| 16 | Acute hyperkinetic movement disorders in Italian paediatric emergency departments. Archives of Disease in Childhood, 2018, 103, 790-794. | 1.0 | 19 |
| 17 | Benign and severe early-life seizures: a round in the first year of life. Italian Journal of Pediatrics, 2018, 44, 54. | 1.0 | 19 |
| 18 | Clinical Course of N-Methyl-D-Aspartate Receptor Encephalitis and the Effectiveness of Cyclophosphamide Treatment. Journal of Pediatric Neurology, 2017, 15, 084-089. | 0.0 | 18 |

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|----|---|-----|-----------|
| 19 | Pyridoxine-dependent epilepsies: an observational study on clinical, diagnostic, therapeutic and prognostic features in a pediatric cohort. <i>Metabolic Brain Disease</i> , 2018, 33, 261-269. | 1.4 | 18 |
| 20 | Epileptic seizures as a manifestation of cow's milk allergy: a studied relationship and description of our pediatric experience. <i>Expert Review of Clinical Immunology</i> , 2014, 10, 1597-1609. | 1.3 | 15 |
| 21 | Biological Drugs in Guillain-Barré Syndrome: An Update. <i>Current Neuropharmacology</i> , 2017, 15, 938-950. | 1.4 | 15 |
| 22 | Ketogenic diet in pediatric seizures: a randomized controlled trial review and meta-analysis. <i>Expert Review of Neurotherapeutics</i> , 2022, 22, 169-177. | 1.4 | 15 |
| 23 | Graph theory in paediatric epilepsy: A systematic review. <i>Dialogues in Clinical Neuroscience</i> , 2021, 23, 3-13. | 1.8 | 15 |
| 24 | PRRT2 gene variant in a child with dysmorphic features, congenital microcephaly, and severe epileptic seizures: genotype-phenotype correlation?. <i>Italian Journal of Pediatrics</i> , 2019, 45, 159. | 1.0 | 13 |
| 25 | Treating the symptom or treating the disease in neonatal seizures: a systematic review of the literature. <i>Italian Journal of Pediatrics</i> , 2021, 47, 85. | 1.0 | 13 |
| 26 | A New Patient with Potocki-Lupski Syndrome: A Literature Review. <i>Journal of Pediatric Genetics</i> , 2018, 07, 029-034. | 0.3 | 11 |
| 27 | Polysomnographic evaluation of non-invasive ventilation in children with neuromuscular disease. <i>Respirology</i> , 2014, 19, 80-84. | 1.3 | 10 |
| 28 | Incidence of Mediterranean Spotted Fever in Sicilian children: a clinical-epidemiological observational retrospective study from 1987 to 2010. <i>International Journal of Infectious Diseases</i> , 2015, 31, 35-40. | 1.5 | 10 |
| 29 | Therapeutic approaches to pediatric pseudotumor cerebri: New insights from literature data. <i>International Journal of Immunopathology and Pharmacology</i> , 2017, 30, 94-97. | 1.0 | 10 |
| 30 | Ictal blinking, an under-recognized phenomenon: our experience and literature review. <i>Neuropsychiatric Disease and Treatment</i> , 2017, Volume 13, 1435-1439. | 1.0 | 10 |
| 31 | Spinal motor neuron involvement in a patient with homozygous PRUNE mutation. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 541-543. | 0.7 | 10 |
| 32 | Cardiorespiratory fitness and sports activities in children and adolescents with solitary functioning kidney. <i>Italian Journal of Pediatrics</i> , 2016, 42, 43. | 1.0 | 9 |
| 33 | Clinical spectrum of woolly hair: indications for cerebral involvement. <i>Italian Journal of Pediatrics</i> , 2017, 43, 99. | 1.0 | 9 |
| 34 | Chilblains-like lesions and SARS-CoV-2 in children: An overview in therapeutic approach. <i>Dermatologic Therapy</i> , 2021, 34, e14502. | 0.8 | 9 |
| 35 | Neonates Born to COVID-19 Mother and Risk in Management within 4 Weeks of Life: A Single-Center Experience, Systematic Review, and Meta-Analysis. <i>American Journal of Perinatology</i> , 2021, 38, 1010-1022. | 0.6 | 8 |
| 36 | Cerebral White Matter Lesions and Dysmorphisms: Signs Suggestive of 6p25 Deletion Syndrome—Literature Review. <i>Journal of Pediatric Genetics</i> , 2019, 08, 205-211. | 0.3 | 7 |

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|----|--|-----|-----------|
| 37 | High mobility group box 1 and markers of oxidative stress in human cord blood. <i>Pediatrics International</i> , 2019, 61, 264-270. | 0.2 | 7 |
| 38 | Inter-society consensus for the use of inhaled corticosteroids in infants, children and adolescents with airway diseases. <i>Italian Journal of Pediatrics</i> , 2021, 47, 97. | 1.0 | 7 |
| 39 | Clinical features and long-term outcomes in pediatric cyclic vomiting syndrome: A 9-year experience at three tertiary academic centers. <i>Neurogastroenterology and Motility</i> , 2021, , e14224. | 1.6 | 7 |
| 40 | Fever-Associated Seizures or Epilepsy: An Overview of Old and Recent Literature Acquisitions. <i>Frontiers in Pediatrics</i> , 2022, 10, 858945. | 0.9 | 7 |
| 41 | The Gut-brain axis: A new pathogenic view of neurologic symptoms – Description of a pediatric case. <i>Journal of Pediatric Neurosciences</i> , 2017, 12, 105. | 0.2 | 6 |
| 42 | Natalizumab in multiple sclerosis: discontinuation, progressive multifocal leukoencephalopathy and possible use in children. <i>Expert Review of Neurotherapeutics</i> , 2015, 15, 1321-1341. | 1.4 | 5 |
| 43 | Chromosome 2p15-p16.1 microduplication in a boy with congenital anomalies: Is it a distinctive syndrome?. <i>European Journal of Medical Genetics</i> , 2019, 62, 47-54. | 0.7 | 5 |
| 44 | Influence of etiology on treatment choices for neonatal seizures: A survey among pediatric neurologists. <i>Brain and Development</i> , 2019, 41, 595-599. | 0.6 | 5 |
| 45 | Severe Psychotic Symptoms in Youth with PANS/PANDAS: Case-Series. <i>Journal of Child and Adolescent Psychopharmacology</i> , 2020, 30, 567-571. | 0.7 | 5 |
| 46 | Is SARSCoV-2 nasopharyngeal swab still a gold standard in children?. <i>Medical Hypotheses</i> , 2020, 144, 110041. | 0.8 | 5 |
| 47 | Dry Eye in Systemic Sclerosis Patients: Novel Methods to Monitor Disease Activity. <i>Diagnostics</i> , 2020, 10, 404. | 1.3 | 5 |
| 48 | aEEG vs cEEG™s sensivity for seizure detection in the setting of neonatal intensive care units: a systematic review and meta-analysis. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2022, , . | 0.7 | 5 |
| 49 | Electrocardiographic Evaluation in Patients With Spinal Muscular Atrophy: A Case-Control Study. <i>Journal of Child Neurology</i> , 2018, 33, 487-492. | 0.7 | 4 |
| 50 | Nervous system involvement in clinical peripheral inflammation: A description of three pediatric cases. <i>Journal of Pediatric Neurosciences</i> , 2016, 11, 277. | 0.2 | 4 |
| 51 | Evidences of emerging pain consciousness during prenatal development: a narrative review. <i>Neurological Sciences</i> , 2022, 43, 3523-3532. | 0.9 | 4 |
| 52 | Thrombotic events in children and adolescent patients with SARS-CoV-2 infection: a systematic review with meta-analysis on incidence and management. <i>Expert Review of Hematology</i> , 2022, 15, 635-643. | 1.0 | 4 |
| 53 | Usefulness of video-EEG in the paediatric emergency department. <i>Expert Review of Neurotherapeutics</i> , 2014, 14, 769-785. | 1.4 | 3 |
| 54 | Long-term survival in a patient with muscle-eye-brain disease. <i>Neurological Sciences</i> , 2015, 36, 2147-2149. | 0.9 | 3 |

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|----|---|-----|-----------|
| 55 | Delayed neonatal visual evoked potentials are associated to asymmetric growth pattern in twins. <i>Clinical Neurophysiology</i> , 2020, 131, 744-749. | 0.7 | 3 |
| 56 | Ocular Motor Paroxysmal Events in Neonates and Infants: A Review of the Literature. <i>Pediatric Neurology</i> , 2021, 117, 4-9. | 1.0 | 3 |
| 57 | Sutures ultrasound: useful diagnostic screening for posterior plagiocephaly. <i>Child's Nervous System</i> , 2021, 37, 3715-3720. | 0.6 | 3 |
| 58 | Down syndrome and arterial ischemic stroke in childhood: A potential immunologic link with selective IgG4 subclass deficiency. <i>European Journal of Paediatric Neurology</i> , 2014, 18, 520-525. | 0.7 | 2 |
| 59 | Giant Melanocytic Nevi and Soft Tissue Undergrowth in the Left Leg: Pathogenetic Hypothesis. <i>Mental Illness</i> , 2019, 11, 8184. | 0.8 | 2 |
| 60 | Monogenic Syndromes with Congenital Heart Diseases in Newborns (Diagnostic Clues for) <i>Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 547 Td</i> 2021, 10, 173-193. | 0.3 | 2 |
| 61 | Non-Epileptic Paroxysmal Events: Clinical features and diagnostic differences with epileptic seizures. A Single Tertiary Centre Study. <i>Clinical Neurology and Neurosurgery</i> , 2021, 207, 106739. | 0.6 | 2 |
| 62 | Correspondence to levetiracetam in neonatal seizures as first-line treatment. <i>Journal of Pediatric Neurosciences</i> , 2018, 13, 520. | 0.2 | 2 |
| 63 | Beyond neonatal seizures - epileptic evolution in preterm newborns: a systematic review and meta-analysis. <i>Epileptic Disorders</i> , 2022, 24, 140-150. | 0.7 | 2 |
| 64 | Neonatal ischemic limb lesions: From etiology to topical nitroglycerine. A case series analysis. <i>Dermatologic Therapy</i> , 2022, 35, e15426. | 0.8 | 2 |
| 65 | Developmental EEG hallmark or biological artifact? Glossokinetic artifact mimicking anterior slow dysrhythmia in two full term newborns. <i>Neurophysiologie Clinique</i> , 2019, 49, 377-380. | 1.0 | 1 |
| 66 | Previously Unreported &lt;b>&lt;i>COL7A1&lt;/b> Mutation in a Somali Patient with Dystrophic Epidermolysis Bullosa. <i>Molecular Syndromology</i> , 2019, 10, 332-338. | 0.3 | 1 |
| 67 | Malformations of Cortical Development, Cognitive Involvement and Epilepsy: A Single Institution Experience in 19 Young Patients. <i>Children</i> , 2021, 8, 637. | 0.6 | 1 |
| 68 | Chorioamnionitis, Inflammation and Neonatal Apnea: Effects on Preterm Neonatal Brainstem and on Peripheral Airways: Chorioamnionitis and Neonatal Respiratory Functions. <i>Children</i> , 2021, 8, 917. | 0.6 | 1 |
| 69 | Neurodevelopmental outcomes of neonatal non-epileptic paroxysmal events: a prospective study. <i>Developmental Medicine and Child Neurology</i> , 2021, 63, 343-348. | 1.1 | 1 |
| 70 | A Novel 4q32.3 Deletion in a Boy: Additional Signs and the Role of MARCH1. <i>Journal of Pediatric Genetics</i> , 2021, 10, 259-265. | 0.3 | 1 |
| 71 | The Role of Dendritic Cells in Central Nervous System Autoimmunity: Focusing on Multiple Sclerosis and Emerging Therapeutics Targeting Dendritic Cells. <i>Journal of Pediatric Biochemistry</i> , 2017, 06, 121-135. | 0.2 | 0 |
| 72 | A mild phenotype associated with a de novo microdeletion 10q23.1-q23.2: a new patient with a novel feature. <i>BMJ Case Reports</i> , 2016, 2016, bcr2016214388. | 0.2 | 0 |