Coskun Yarar

List of Publications by Year in descending order

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759233 888059 50 387 12 17 h-index citations g-index papers 50 50 50 621 times ranked docs citations citing authors all docs

| # | Article | IF | CITATIONS |
|----|---|---------|-----------|
| 1 | Evaluation of immunization status in patients with cerebral palsy: a multicenter CP-VACC study. European Journal of Pediatrics, 2022, 181, 383-391. | 2.7 | 2 |
| 2 | Evaluation of micronutrient levels in children with cerebral palsy. Pediatrics International, 2022, 64, . | 0.5 | 3 |
| 3 | Wiedemann–Steiner Syndrome: A Rare Differential Diagnosis of Neurodevelopmental Delay and Dysmorphic Features. Journal of Pediatric Genetics, 2022, 11, 162-164. | 0.7 | 0 |
| 4 | Acute flaccid myelitis outbreak through 2016–2018: A multicenter experience from Turkey. European Journal of Paediatric Neurology, 2021, 30, 113-120. | 1.6 | 1 |
| 5 | Rituximab Treatment in Acute Disseminated Encephalomyelitis Associated with Salmonella Infection. Case Reports in Pediatrics, 2021, 2021, 1-3. | 0.4 | 2 |
| 6 | Brain Abscess in a Patient with Osteopetrosis: A Rare Complication. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 358-361. | 0.9 | 0 |
| 7 | NDE1 â€related disorders: A recurrent NDE1 pathogenic variant causing Lissencephaly 4 can also be associated with microhydranencephaly. American Journal of Medical Genetics, Part A, 2021, , . | 1.2 | 0 |
| 8 | Metoclopramide-Induced Acute Dystonia. Pediatric Emergency Care, 2021, 37, e528-e533. | 0.9 | 5 |
| 9 | Niemann–Pick type C disease with a novel intronic mutation: three Turkish cases from the same family. Journal of Pediatric Endocrinology and Metabolism, 2021, . | 0.9 | 2 |
| 10 | Screening for functional gastrointestinal disorders in children with epilepsy. Epilepsy and Behavior, 2020, 111, 107267. | 1.7 | 9 |
| 11 | Viral etiological causes of febrile seizures for respiratory pathogens (EFES Study). Human Vaccines and Immunotherapeutics, 2019, 15, 496-502. | 3.3 | 38 |
| 12 | Quality-of-Life Evaluation of Healthy Siblings of Children with Chronic Illness. Balkan Medical Journal, 2019, 37, 34-42. | 0.8 | 21 |
| 13 | Çocukluk Çağı Epilepsilerine Karşı Toplumsal Tutum Ölçeği (ÇEKTÖL) Geliştirilmesi: Geçerlik ve Çalışması. Turk Pediatri Arsivi, 2019, 55, 23-29. | Gývenii | rlik 1 |
| 14 | Optic Nerve Sheath Diameter Measurement in Children Presenting to a Pediatric Emergency Department with Head Trauma. Journal of Pediatric Emergency and Intensive Care Medicine, 2019, 6, 7-12. | 0.1 | 1 |
| 15 | The immunization status of children with chronic neurological disease and serological assessment of vaccine-preventable diseases. Human Vaccines and Immunotherapeutics, 2018, 14, 1970-1976. | 3.3 | 13 |
| 16 | Menstruationâ€related headache in adolescents: Point prevalence and associated factors. Pediatrics International, 2018, 60, 576-580. | 0.5 | 5 |
| 17 | Neurocognitive Consequences of Childhood Leukemia and Its Treatment. Indian Journal of Hematology and Blood Transfusion, 2018, 34, 62-69. | 0.6 | 1 |
| 18 | A multicenter cross-sectional study to evaluate the clinical characteristics and nutritional status of children with cerebral palsy. Clinical Nutrition ESPEN, 2018, 26, 27-34. | 1.2 | 31 |

| # | Article | lF | CITATIONS |
|----|--|-----|-----------|
| 19 | Apolipoprotein e allelic variants and cerebral palsy. Turkish Journal of Pediatrics, 2018, 60, 361. | 0.6 | 4 |
| 20 | Incomplete Millerâ€"Fisher syndrome with advanced stage Burkitt lymphoma. Indian Pediatrics, 2017, 54, 413-415. | 0.4 | 0 |
| 21 | Characteristics of pediatric multiple sclerosis: The Turkish pediatric multiple sclerosis database. European Journal of Paediatric Neurology, 2017, 21, 864-872. | 1.6 | 18 |
| 22 | Congenital amegakaryocytic thrombocytopenia with severe neurological findings. Blood Coagulation and Fibrinolysis, 2016, 27, 936-939. | 1.0 | 3 |
| 23 | Neurodevelopment of preterm infants born after <i>in vitro</i> fertilization and spontaneous multiple pregnancy. Pediatrics International, 2016, 58, 1284-1290. | 0.5 | 12 |
| 24 | Evaluation of hypercoagulability state in perinatal arterial ischemic stroke with rotation thromboelastometry. Child's Nervous System, 2016, 32, 2395-2401. | 1.1 | 4 |
| 25 | Encephalocraniocutaneous lipomatosis, a rare neurocutaneous disorder: report of additional three cases. Child's Nervous System, 2016, 32, 559-562. | 1.1 | 16 |
| 26 | Hemi-meningitis with hemophagocytic lymphohistiocytosis. Annals of Indian Academy of Neurology, 2016, 19, 388. | 0.5 | 2 |
| 27 | A Multinational Survey on Actual Diagnostics and Treatment of Subacute Sclerosing Panencephalitis. Neuropediatrics, 2015, 46, 377-384. | 0.6 | 12 |
| 28 | Novel mutations in PTPN11 gene in two girls with Noonan syndrome phenotype. International Journal of Cardiology, 2015, 186, 13-15. | 1.7 | 7 |
| 29 | Akathisia in association with herpes simplex encephalitis relapse and opercular syndrome in children. Brain and Development, 2014, 36, 167-170. | 1.1 | 3 |
| 30 | Essential Palatal Tremor Treated With Botulinum Toxin. Pediatric Neurology, 2013, 48, 415-417. | 2.1 | 7 |
| 31 | Lennox-Gastaut sendromlu hastaların klinik özellikleri ve uzun sÃ⅓reli seyri. Turk Pediatri Arsivi, 2012, 47, 47-51. | 0.9 | 2 |
| 32 | GLUTARIC ACIDURIA TYPE 1 PRESENTING AS SUBDURAL HAEMATOMA. Journal of Paediatrics and Child Health, 2012, 48, 712-712. | 0.8 | 5 |
| 33 | Çocuklarda idyopatik fasiyal paralizi. Medical Journal of Bakirkoy, 2012, , 107-110. | 0.1 | 2 |
| 34 | Infant with chromhidrosis. Pediatrics International, 2011, 53, 283-284. | 0.5 | 8 |
| 35 | Serum asymmetric dimethylarginine (ADMA), homocysteine, vitamin B12, folate levels, and lipid profiles in epileptic children treated with valproic acid. European Journal of Pediatrics, 2011, 170, 873-877. | 2.7 | 16 |
| 36 | Magnetic resonance imaging at first episode in pediatric multiple sclerosis retrospective evaluation according to KIDMUS and lesion dissemination in space criteria. Brain and Development, 2010, 32, 487-494. | 1.1 | 5 |

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| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 37 | Relapsing Acute Disseminated Encephalomyelitis in Children: Further Evaluation of the Diagnosis. Journal of Child Neurology, 2010, 25, 1491-1497. | 1.4 | 6 |
| 38 | Spontaneous Acute Subdural Hematoma and Chronic Epidural Hematoma in a Child with F XIII Deficiency. Journal of Emergency Medicine, 2010, 38, 25-29. | 0.7 | 16 |
| 39 | Septo-Optic Dysplasia Plus: A Patient With Diabetes Insipidus. Pediatric Neurology, 2010, 43, 76-78. | 2.1 | 6 |
| 40 | MIDAS (microphthalmia, dermal aplasia, sclerocornea) syndrome with central nervous system abnormalities. Clinical Dysmorphology, 2009, 18, 234-235. | 0.3 | 13 |
| 41 | Serum Amyloid A, Procalcitonin, Tumor Necrosis Factor- <mml:math xmlns:mml="http://www.w3.org/1998/Math/MathML"><mml:mi>α</mml:mi></mml:math> , and Interleukin-1 <mml:math xmlns:mml="http://www.w3.org/1998/Math/MathML"><mml:mi>β</mml:mi></mml:math> Levels in Neonatal Late-Onset Sepsis: Mediators of Inflammation, 2008, 2008, 1-7. | 3.0 | 37 |
| 42 | Successful Treatment with Linezolid of Meningitis Complicated with Subdural Empyema in a 6-Month-Old Boy. Journal of Tropical Pediatrics, 2007, 53, 431-433. | 1.5 | 9 |
| 43 | Pericardial effusion due to hypothyroidism in Down syndrome: report of four cases. Neuroendocrinology Letters, 2007, 28, 141-4. | 0.2 | 5 |
| 44 | Antibiotic resistance in children with complicated urinary tract infection. Journal of King Abdulaziz University, Islamic Economics, 2007, 28, 1850-4. | 1.1 | 5 |
| 45 | Neuhauser syndrome and Peters' anomaly. Clinical Dysmorphology, 2006, 15, 249-251. | 0.3 | 2 |
| 46 | Bell's palsy and hepatitis infection. Pediatrics International, 2006, 48, 493-494. | 0.5 | 6 |
| 47 | Serum leptin levels in children with cerebral palsy: relationship with growth and nutritional status. Neuroendocrinology Letters, 2006, 27, 507-12. | 0.2 | 4 |
| 48 | A patient with cystinosis presenting transient features of Bartter syndrome. Turkish Journal of Pediatrics, 2006, 48, 260-2. | 0.6 | 14 |
| 49 | Atypical hemolytic uremic syndrome associated with group A beta hemolytic streptococcus. Pediatric Nephrology, 2004, 19, 943-4; author reply 945. | 1.7 | 3 |
| 50 | Identification of a Homozygous Deletion within FGD4 in a Charcot-Marie-Tooth type 4H Family by Exome Sequencing. Journal of Pediatric Neurology, 0, , . | 0.2 | 0 |