Coskun Yarar

List of Publications by Year
in descending order

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

#	Article	IF	Citations
1	Viral etiological causes of febrile seizures for respiratory pathogens (EFES Study). Human Vaccines and Immunotherapeutics, 2019, 15, 496-502.	3.3	38
2	Serum Amyloid A, Procalcitonin, Tumor Necrosis Factor-<mml:math xmlns:mml="http://www.w3.org/1998/Math/MathML"><mml:mi>Î±</mml:mi></mml:math>, and Interleukin-1<mml:math xmlns:mml="http://www.w3.org/1998/Math/MathML"><mml:mi>Î²</mml:mi></mml:math>Levels in Neonatal Late-Onset Sepsis. Mediators of Inflammation, 2008, 2008, 1-7.	3.0	37
3	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
4	Quality-of-Life Evaluation of Healthy Siblings of Children with Chronic Illness. Balkan Medical Journal, 2019, 37, 34-42.	0.8	21
5	Characteristics of pediatric multiple sclerosis: The Turkish pediatric multiple sclerosis database. European Journal of Paediatric Neurology, 2017, 21, 864-872.	1.6	18
6	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
7	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
8	Encephalocraniocutaneous lipomatosis, a rare neurocutaneous disorder: report of additional three cases. Child's Nervous System, 2016, 32, 559-562.	1.1	16
9	A patient with cystinosis presenting transient features of Bartter syndrome. Turkish Journal of Pediatrics, 2006, 48, 260-2.	0.6	14
10	MIDAS (microphthalmia, dermal aplasia, sclerocornea) syndrome with central nervous system abnormalities. Clinical Dysmorphology, 2009, 18, 234-235.	0.3	13
11	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
12	A Multinational Survey on Actual Diagnostics and Treatment of Subacute Sclerosing Panencephalitis. Neuropediatrics, 2015, 46, 377-384.	0.6	12
13	Neurodevelopment of preterm infants born after in vitro fertilization and spontaneous multiple pregnancy. Pediatrics International, 2016, 58, 1284-1290.	0.5	12
14	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
15	Screening for functional gastrointestinal disorders in children with epilepsy. Epilepsy and Behavior, 2020, 111, 107267.	1.7	9
16	Infant with chromhidrosis. Pediatrics International, 2011, 53, 283-284.	0.5	8
17	Essential Palatal Tremor Treated With Botulinum Toxin. Pediatric Neurology, 2013, 48, 415-417.	2.1	7
18	Novel mutations in PTPN11 gene in two girls with Noonan syndrome phenotype. International Journal of Cardiology, 2015, 186, 13-15.	1.7	7

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19	Bell's palsy and hepatitis infection. Pediatrics International, 2006, 48, 493-494.	0.5	6
20	Relapsing Acute Disseminated Encephalomyelitis in Children: Further Evaluation of the Diagnosis. Journal of Child Neurology, 2010, 25, 1491-1497.	1.4	6
21	Septo-Optic Dysplasia Plus: A Patient With Diabetes Insipidus. Pediatric Neurology, 2010, 43, 76-78.	2.1	6
22	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
23	GLUTARIC ACIDURIA TYPE 1 PRESENTING AS SUBDURAL HAEMATOMA. Journal of Paediatrics and Child Health, 2012, 48, 712-712.	0.8	5
24	Menstruationâ€related headache in adolescents: Point prevalence and associated factors. Pediatrics International, 2018, 60, 576-580.	0.5	5
25	Metoclopramide-Induced Acute Dystonia. Pediatric Emergency Care, 2021, 37, e528-e533.	0.9	5
26	Pericardial effusion due to hypothyroidism in Down syndrome: report of four cases. Neuroendocrinology Letters, 2007, 28, 141-4.	0.2	5
27	Antibiotic resistance in children with complicated urinary tract infection. Journal of King Abdulaziz University, Islamic Economics, 2007, 28, 1850-4.	1.1	5
28	Evaluation of hypercoagulability state in perinatal arterial ischemic stroke with rotation thromboelastometry. Child's Nervous System, 2016, 32, 2395-2401.	1.1	4
29	Apolipoprotein e allelic variants and cerebral palsy. Turkish Journal of Pediatrics, 2018, 60, 361.	0.6	4
30	Serum leptin levels in children with cerebral palsy: relationship with growth and nutritional status. Neuroendocrinology Letters, 2006, 27, 507-12.	0.2	4
31	Atypical hemolytic uremic syndrome associated with group A beta hemolytic streptococcus. Pediatric Nephrology, 2004, 19, 943-4; author reply 945.	1.7	3
32	Akathisia in association with herpes simplex encephalitis relapse and opercular syndrome in children. Brain and Development, 2014, 36, 167-170.	1.1	3
33	Congenital amegakaryocytic thrombocytopenia with severe neurological findings. Blood Coagulation and Fibrinolysis, 2016, 27, 936-939.	1.0	3
34	Evaluation of micronutrient levels in children with cerebral palsy. Pediatrics International, 2022, 64, .	0.5	3
35	Neuhauser syndrome and Peters' anomaly. Clinical Dysmorphology, 2006, 15, 249-251.	0.3	2
36	Lennox-Gastaut sendromlu hastalarÄ±n klinik Ã¶zellikleri ve uzun sÃ¼reli seyri. Turk Pediatri Arsivi, 2012, 47, 47-51.	0.9	2

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37	Rituximab Treatment in Acute Disseminated Encephalomyelitis Associated with Salmonella Infection. Case Reports in Pediatrics, 2021, 2021, 1-3.	0.4	2
38	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
39	Hemi-meningitis with hemophagocytic lymphohistiocytosis. Annals of Indian Academy of Neurology, 2016, 19, 388.	0.5	2
40	Ã‡ocuklarda idyopatik fasiyal paralizi. Medical Journal of Bakirkoy, 2012, , 107-110.	0.1	2
41	Niemannâ€“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
42	Neurocognitive Consequences of Childhood Leukemia and Its Treatment. Indian Journal of Hematology and Blood Transfusion, 2018, 34, 62-69.	0.6	1
43	Acute flaccid myelitis outbreak through 2016â€“2018: A multicenter experience from Turkey. European Journal of Paediatric Neurology, 2021, 30, 113-120.	1.6	1
44	Ã‡ocukluk Ã‡aÄŸÄ± Epilepsilerine KarÅŸÄ± Toplumsal Tutum Ã–lÃ§eÄŸi (Ã‡EKTÃ–L) GeliÅŸtirilmesi: GeÃ§erlik ve GÃ¼venirlik Ã‡alÄ±ÅŸmasÄ±. Turk Pediatri Arsivi, 2019, 55, 23-29.	0.9	1
45	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
46	Incomplete Millerâ€”Fisher syndrome with advanced stage Burkitt lymphoma. Indian Pediatrics, 2017, 54, 413-415.	0.4	0
47	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
48	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
49	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
50	Wiedemannâ€“Steiner Syndrome: A Rare Differential Diagnosis of Neurodevelopmental Delay and Dysmorphic Features. Journal of Pediatric Genetics, 2022, 11, 162-164.	0.7	0