Kyung-Hee Park

List of Publications by Year
in descending order

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Version: 2024-02-01

28	123	1478505 6	1372567 10
papers	citations	h-index	g-index

29	29	29	172
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Incidence of congenital hypothyroidism by gestational age: a retrospective observational study. , 2023, 40, 30-36.		2
2	Clinical and Laboratory Findings of Nosocomial Sepsis in Extremely Low Birth Weight Infants According to Causative Organisms. Journal of Clinical Medicine, 2022, 11, 260.	2.4	2
3	Implementing the Golden Hour Protocol to Improve the Clinical Outcomes in Preterm Infants. Neonatal Medicine, 2022, 29, 10-17.	0.2	0
4	Neurodevelopmental Outcomes of Very-Low-Birth-Weight Infants without Severe Brain Lesions and Impact of Postnatal Steroid Use: A Single-Center Korean Study. Neonatal Medicine, 2022, 29, 36-45.	0.2	0
5	Perinatal Prognostic Factors for Congenital Diaphragmatic Hernia: A Korean Single-Center Study. Neonatal Medicine, 2022, 29, 76-83.	0.2	0
6	Genetic Aspects of Small for Gestational Age Infants Using Targeted-Exome Sequencing and Whole-Exome Sequencing: A Single Center Study. Journal of Clinical Medicine, 2022, 11, 3710.	2.4	1
7	Chromosomal Microarray in Children With Developmental Delay: The Experience of a Tertiary Center in Korea. Frontiers in Pediatrics, 2021, 9, 690493.	1.9	7
8	Using lactate dehydrogenase to predict the severity of respiratory distress in term newborn infants with no perinatal asphyxia. Turkish Journal of Pediatrics, 2021, 63, 393.	0.6	0
9	Association between vitamin D deficiency at one month of age and bronchopulmonary dysplasia. Medicine (United States), 2021, 100, e27966.	1.0	3
10	Lung Volume Reduction Surgery in Preterm Infant with Giant Bullae due to Bronchopulmonary Dysplasia: A Case Report. Perinatology, 2021, 32, 204.	0.1	0
11	Meconium peritonitis resulting from different etiologies in siblings: a case report. BMC Pediatrics, 2020, 20, 106.	1.7	6
12	Identification of a novel variant in the PHEX gene using targeted gene panel sequencing in a 24-month-old boy with hypophosphatemic rickets. Annals of Pediatric Endocrinology and Metabolism, 2020, 25, 63-67.	2.3	2
13	An A627V-activating mutation in the thyroid-stimulating hormone receptor gene in familial nonautoimmune hyperthyroidism. Annals of Pediatric Endocrinology and Metabolism, 2020, 25, 282-286.	2.3	1
14	Predictive value of an early amplitude-integrated electroencephalogram for short-term neurologic outcomes in preterm infants. Turkish Journal of Pediatrics, 2020, 62, 367.	0.6	0
15	Utility of Magnetic Resonance Imaging (MRI) in Children With Strabismus. Journal of Child Neurology, 2019, 34, 574-581.	1.4	2
16	Characteristics of Bilirubin According to the Results of the Direct Antiglobulin Test and Its Impact in Hemolytic Disease of the Newborn. Laboratory Medicine, 2019, 50, 138-144.	1.2	4
17	Clinical characterization of anti-GQ1b antibody syndrome in Korean children. Journal of Neuroimmunology, 2019, 330, 170-173.	2.3	11
18	Bilateral Acute Retinal Necrosis and Encephalomalacia Due to Herpes Simplex Virus Infection in a Premature Infant. Neonatal Medicine, 2019, 26, 63-66.	0.2	1

#	Article	IF	Citations
19	The clinical characteristics and neurodevelopmental outcome of preterm infants with persistent periventricular echogenicity. Pediatrics and Neonatology, 2018, 59, 606-610.	0.9	4
20	Comparison of conservative therapy and steroid therapy for Bellâ€™s palsy in children. Korean Journal of Pediatrics, 2018, 61, 332-337.	1.9	8
21	Chylous Ascites in an Infant with Thanatophoric Dysplasia Type I with FGFR3 Mutation Surviving Five Months. Fetal and Pediatric Pathology, 2018, 37, 363-371.	0.7	1
22	Extended Red Blood Cell Genotyping to Investigate Immunohematology Problems. Annals of Laboratory Medicine, 2018, 38, 387-388.	2.5	3
23	Vitamin D deficiency in children aged 6 to 12 years: single center's experience in Busan. Annals of Pediatric Endocrinology and Metabolism, 2016, 21, 149.	2.3	21
24	Venous angioma may be associated with epilepsy in children. Korean Journal of Pediatrics, 2016, 59, 341.	1.9	5
25	Novel and Recurrent ACADS Mutations and Clinical Manifestations Observed in Korean Patients with Short-chain Acyl-coenzyme a Dehydrogenase Deficiency. Annals of Clinical and Laboratory Science, 2016, 46, 360-6.	0.2	5
26	Neonatal Iliopsoas Abscess: The First Korean Case. Journal of Korean Medical Science, 2015, 30, 1203.	2.5	8
27	Optic Neuritis in Korean Children: Low Risk of Subsequent Multiple Sclerosis. Pediatric Neurology, 2015, 53, 221-225.	2.1	17
28	Cornelia de Lange Syndrome with NIPBL Gene Mutation: A Case Report. Journal of Korean Medical Science, 2010, 25, 1821.	2.5	9