Young-Mi Han

List of Publications by Year
in descending order

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Version: 2024-02-01

23 papers	54 citations	1937685 4 h-index	1872680 6 g-index
24 all docs	24 docs citations	24 times ranked	60 citing authors

#	Article	IF	Citations
1	Comparison of effectiveness of growth hormone therapy according to disease-causing genes in children with Noonan syndrome. Korean Journal of Pediatrics, 2019, 62, 274-280.	1.9	11
2	Neonatal Iliopsoas Abscess: The First Korean Case. Journal of Korean Medical Science, 2015, 30, 1203.	2.5	8
3	Effect of Prophylactic Palivizumab on Admission Due to Respiratory Syncytial Virus Infection in Former Very Low Birth Weight Infants with Bronchopulmonary Dysplasia. Journal of Korean Medical Science, 2015, 30, 924.	2.5	6
4	Meconium peritonitis resulting from different etiologies in siblings: a case report. BMC Pediatrics, 2020, 20, 106.	1.7	6
5	Bartâ€™s Syndrome with Novel Frameshift Mutations in theCOL7A1Gene. Fetal and Pediatric Pathology, 2019, 38, 72-79.	0.7	5
6	The clinical characteristics and neurodevelopmental outcome of preterm infants with persistent periventricular echogenicity. Pediatrics and Neonatology, 2018, 59, 606-610.	0.9	4
7	Association between vitamin D deficiency at one month of age and bronchopulmonary dysplasia. Medicine (United States), 2021, 100, e27966.	1.0	3
8	Neonatal Lupus Erythematosus as a Rare Cause of Fever in Young Infants. Journal of Clinical Medicine, 2021, 10, 3195.	2.4	2
9	Establishing reference values for amplitude-integrated electroencephalography in preterms below 35 weeks of gestational age: A prospective observational cohort study. Turkish Journal of Pediatrics, 2016, 58, 592-601.	0.6	2
10	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
11	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
12	Fatal outcome of autosomal recessive polycystic kidney disease in neonates with recessive PKHD1 mutations. Medicine (United States), 2020, 99, e20113.	1.0	1
13	Poland Syndrome in One Dizygotic Twin: A Case Report. Korean Journal of Perinatology, 2015, 26, 352.	0.1	1
14	Risk Factors for Delayed Hyperthyrotropinemia in Late Preterm Infants. Neonatal Medicine, 2019, 26, 204-212.	0.2	1
15	Scrotal pyocele secondary to gastrointestinal perforation in infants: a case series. Yeungnam University Journal of Medicine, 2023, 40, 86-90.	1.4	1
16	Congenital Syphilis in Neonate: A Single Center Study for 10 Years. Journal of the Korean Society of Maternal and Child Health, 2021, 25, 204-210.	0.6	0
17	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
18	Clinical Features of Cricopharyngeal Incoordination in Newborns and Infants. Korean Journal of Pediatric Gastroenterology and Nutrition, 2008, 11, 116.	0.2	0

#	Article	IF	Citations
19	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
20	Lung Volume Reduction Surgery in Preterm Infant with Giant Bullae due to Bronchopulmonary Dysplasia: A Case Report. Perinatology, 2021, 32, 204.	0.1	0
21	Implementing the Golden Hour Protocol to Improve the Clinical Outcomes in Preterm Infants. Neonatal Medicine, 2022, 29, 10-17.	0.2	0
22	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
23	Perinatal Prognostic Factors for Congenital Diaphragmatic Hernia: A Korean Single-Center Study. Neonatal Medicine, 2022, 29, 76-83.	0.2	0