Raffaella Morotti

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9980379/publications.pdf

Version: 2024-02-01

22 papers

911 citations

687363 13 h-index 713466 21 g-index

23 all docs 23 docs citations

23 times ranked

1447 citing authors

#	Article	IF	CITATIONS
1	Neuropathology of Chiari Malformation II with Chromosome X Alterations: An Autopsy Study in a 17-Month-Old and Review of Literature. Journal of Neuropathology and Experimental Neurology, 2022, 81, 296-298.	1.7	1
2	Placental Tissue Destruction and Insufficiency From COVID-19 Causes Stillbirth and Neonatal Death From Hypoxic-Ischemic Injury. Archives of Pathology and Laboratory Medicine, 2022, 146, 660-676.	2.5	127
3	<i>DLG5</i> variants are associated with multiple congenital anomalies including ciliopathy phenotypes. Journal of Medical Genetics, 2021, 58, 453-464.	3.2	10
4	Maternal respiratory SARS-CoV-2 infection in pregnancy is associated with a robust inflammatory response at the maternal-fetal interface. Med, 2021, 2, 591-610.e10.	4.4	122
5	Complete Hydatidiform Mole and Coexisting Fetus With Gastroschisis: A Case Report Highlighting the Importance of Diagnostic Genotyping. Pediatric and Developmental Pathology, 2021, 24, 575-580.	1.0	2
6	Human autoinflammatory disease reveals ELF4 as a transcriptional regulator of inflammation. Nature Immunology, 2021, 22, 1118-1126.	14.5	30
7	Hofbauer Cells and COVID-19 in Pregnancy. Archives of Pathology and Laboratory Medicine, 2021, 145, 1328-1340.	2.5	40
8	Chronic Histiocytic Intervillositis With Trophoblast Necrosis Is a Risk Factor Associated With Placental Infection From Coronavirus Disease 2019 (COVID-19) and Intrauterine Maternal-Fetal Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Transmission in Live-Born and Stillborn Infants. Archives of Pathology and Laboratory Medicine, 2021, 145, 517-528.	2.5	125
9	Outcomes of neonates with listeriosis supported with extracorporeal membrane oxygenation from 1991 to 2017. Journal of Perinatology, 2020, 40, 105-111.	2.0	2
10	Recurrence and Complications in Pediatric and Adolescent Papillary Thyroid Cancer in a High-Volume Practice. Journal of Surgical Research, 2020, 249, 58-66.	1.6	24
11	Progressive Splenomegaly and Hypersplenism: An Unusual Case of Splenic Vein Stenosis with Histologic Findings of Hepatoportal Sclerosis. Journal of Pediatrics, 2020, 218, 222-227.e1.	1.8	1
12	Autopsy Services and Emergency Preparedness of a Tertiary Academic Hospital Mortuary for the COVID-19 Public Health Emergency: The Yale Plan. Advances in Anatomic Pathology, 2020, 27, 355-362.	4.3	6
13	Human Small Intestine Transplantation: Segmental Susceptibility to Ischemia Using Different Preservation Solutions and Conditions. Transplantation Proceedings, 2020, 52, 2934-2940.	0.6	12
14	Cytomorphologic features of thyroid disease in patients with <i>DICER1</i> mutations: A report of cytologyâ€"histopathology correlation in 7 patients. Cancer Cytopathology, 2020, 128, 746-756.	2.4	16
15	Lymph node ratio predicts recurrence in pediatric papillary thyroid cancer. Journal of Pediatric Surgery, 2019, 54, 129-132.	1.6	25
16	Pediatric Nonalcoholic Fatty Liver Disease in New York City: An Autopsy Study. Journal of Pediatrics, 2018, 200, 174-180.	1.8	32
17	Congenital Extrahepatic Portosystemic Shunt (Abernethy Malformation Type Ib) With Associated Hepatocellular Carcinoma. Pediatric and Developmental Pathology, 2017, 20, 354-362.	1.0	39
18	Use of Fluorescein Isothiocyanate-Inulin as a Marker for Intestinal Ischemic Injury. Journal of the American College of Surgeons, 2017, 224, 1066-1073.	0.5	7

#	Article	IF	CITATION
19	Modulation of Intestinal Microbiome Prevents Intestinal Ischemic Injury. Frontiers in Physiology, 2017, 8, 1064.	2.8	21
20	<i>NTRK</i> fusion oncogenes in pediatric papillary thyroid carcinoma in northeast United States. Cancer, 2016, 122, 1097-1107.	4.1	195
21	A hydrogel-endothelial cell implant mimics infantile hemangioma: modulation by survivin and the Hippo pathway. Laboratory Investigation, 2015, 95, 765-780.	3.7	7
22	Paediatric hepatocellular carcinoma due to somatic CTNNB1 and NFE2L2 mutations in the setting of inherited bi-allelic ABCB11 mutations. Journal of Hepatology, 2014, 61, 1178-1183.	3.7	48