

Tricia Bhatti

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

339
citations

1163117

8
h-index

888059

17
g-index

26
all docs

26
docs citations

26
times ranked

699
citing authors

#	ARTICLE	IF	CITATIONS
1	Expanding the phenotypic spectrum of ARCN1-related syndrome. <i>Genetics in Medicine</i> , 2022, 24, 1227-1237.	2.4	5
2	A Novel TP53 Tandem Duplication in a Child with Li-Fraumeni Syndrome. <i>Journal of Physical Education and Sports Management</i> , 2022, , mcs.a006181.	1.2	2
3	Ingested Foreign Bodies Can Cause Appendicitis and Perforation: A Multi-Institutional Case Series. <i>Pediatric and Developmental Pathology</i> , 2022, , 109352662210831.	1.0	5
4	Localized islet nuclear enlargement hyperinsulinism (LINE-HI) due to <i>ABCC8</i> and <i>GCK</i> mosaic mutations. <i>European Journal of Endocrinology</i> , 2022, 187, 301-313.	3.7	4
5	Case Report: Two Distinct Focal Congenital Hyperinsulinism Lesions Resulting From Separate Genetic Events. <i>Frontiers in Pediatrics</i> , 2021, 9, 699129.	1.9	3
6	Utility of Fine-Needle Aspirations to Diagnose Pediatric Thyroid Nodules. <i>Hormone Research in Paediatrics</i> , 2021, 94, 263-274.	1.8	8
7	Excision and Reconstruction of Alveolar Rhabdomyosarcoma Involving the Achilles Tendon in a Pediatric Patient. <i>JBJS Case Connector</i> , 2021, 11, .	0.3	0
8	CAMKV Is a Candidate Immunotherapeutic Target in MYCN Amplified Neuroblastoma. <i>Frontiers in Oncology</i> , 2020, 10, 302.	2.8	13
9	Skull Base Nodular Fasciitis in and Adolescent Male. <i>Journal of Neurological Surgery, Part B: Skull Base</i> , 2020, 81, .	0.8	0
10	Androgenetic chimerism as an etiology for Beckwithâ€“Wiedemann syndrome: diagnosis and management. <i>Genetics in Medicine</i> , 2019, 21, 2644-2649.	2.4	15
11	Characteristics of Follicular Variant Papillary Thyroid Carcinoma in a Pediatric Cohort. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 1639-1648.	3.6	19
12	Multiple Hybrid Cysts in a Child: Challenge. <i>American Journal of Dermatopathology</i> , 2018, 40, e7-e8.	0.6	0
13	Multiple Hybrid Cysts in a Child: Answer. <i>American Journal of Dermatopathology</i> , 2018, 40, 70-71.	0.6	0
14	Tender Nodules on the Lower Legs. <i>JAMA Dermatology</i> , 2018, 154, 471.	4.1	0
15	Erythematous Nodule on the Face of a Child: Challenge. <i>American Journal of Dermatopathology</i> , 2018, 40, e119-e120.	0.6	0
16	Erythematous Nodule on the Face of a Child: Answer. <i>American Journal of Dermatopathology</i> , 2018, 40, 699-700.	0.6	0
17	Congenital syphilis as a clinical and histopathologic mimic of neonatal lupus. <i>Journal of Cutaneous Pathology</i> , 2018, 45, 791-793.	1.3	5
18	Cytotoxic T-Lymphocyte-Associated Protein 4 Haploinsufficiency-Associated Inflammation Can Occur Independently of T-Cell Hyperproliferation. <i>Frontiers in Immunology</i> , 2018, 9, 1715.	4.8	13

#	ARTICLE	IF	CITATIONS
19	Identification of GPC2 as an Oncoprotein and Candidate Immunotherapeutic Target in High-Risk Neuroblastoma. <i>Cancer Cell</i> , 2017, 32, 295-309.e12.	16.8	148
20	Degos disease mimicking primary vasculitis of the CNS. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2016, 3, e206.	6.0	9
21	Post-streptococcal glomerulonephritis associated with atypical hemolytic uremic syndrome: to treat or not to treat with eculizumab?. <i>CKJ: Clinical Kidney Journal</i> , 2016, 9, 90-96.	2.9	10
22	Severe human parechovirus type 3 myocarditis and encephalitis in an adolescent with hypogammaglobulinemia. <i>International Journal of Infectious Diseases</i> , 2015, 36, 6-8.	3.3	19
23	Rapid progression to end-stage renal disease in a child with a sporadic ACTN4 mutation. <i>Clinical Nephrology Case Studies</i> , 2015, 3, 14-18.	0.7	2
24	Stable long-term mixed chimerism achieved in a canine model of allogeneic in utero hematopoietic cell transplantation. <i>Blood</i> , 2014, 124, 1987-1995.	1.4	59