Mariko Eguchi

List of Publications by Year
in descending order

17	152	1478505 6	1199594 12
papers	citations	h-index	g-index

17	17	17	292
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Short stature in a child with a novel Aggrecan gene variant: A case report. Pediatrics International, 2022, 64, e15116.	0.5	0
2	Transvenous pacing approach for atrioventricular block in fontan - Possibility of transvenous approach by electrophysiological assessment -. Journal of Cardiology Cases, 2022, 25, 389-391.	0.5	3
3	Postchemotherapy immune status in infants with acute lymphoblastic leukemia: A report from the JPLSG MLLâ€10 trial. Pediatric Blood and Cancer, 2022, 69, .	1.5	2
4	Active aneurysm thrombosis after Kawasaki disease in an adult: Insight into anticoagulation therapy. Journal of Cardiology Cases, 2021, 23, 206-209.	0.5	1
5	Strayed guidewire into the epidural space during internal jugular vein puncture in a paediatric patient. European Heart Journal - Case Reports, 2021, 5, ytab347.	0.6	0
6	Activation of fibroblast growth factor-inducible 14 in the early phase of childhood IgA nephropathy. PLoS ONE, 2021, 16, e0258090.	2.5	1
7	A novel SOX10 variant in a Japanese girl with Waardenburg syndrome type 4C and Kallmann syndrome. Human Genome Variation, 2020, 7, 30.	0.7	2
8	Early detection of the PAX3â€FOXO1 fusion gene in circulating tumorâ€derived DNA in a case of alveolar rhabdomyosarcoma. Genes Chromosomes and Cancer, 2019, 58, 521-529.	2.8	20
9	Early Surgery Is Feasible for a Very Large Congenital Infantile Fibrosarcoma Associated With Life Threatening Coagulopathy: A Case Report and Literature Review. Frontiers in Pediatrics, 2019, 7, 529.	1.9	2
10	Manifestation of recessive combined Dâ€2â€, Lâ€2â€hydroxyglutaric aciduria in combination with 22q11.2 deletion syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 351-358.	1.2	7
11	Prolonged adrenal insufficiency after high-dose glucocorticoid in infants with leukemia. Pediatric Hematology and Oncology, 2018, 35, 355-361.	0.8	5
12	Lineageâ€dependent skewing of loss of heterozygosity (<scp>LOH</scp>) of <scp>KRAS</scp> gene in a case of juvenile myelomonocytic leukemia. European Journal of Haematology, 2015, 94, 177-181.	2.2	0
13	HMGA2 as a potential molecular target inKMT2A-AFF1-positive infant acute lymphoblastic leukaemia. British Journal of Haematology, 2015, 171, 818-829.	2.5	14
14	CLTCâ€ALK fusion as a primary event in congenital blastic plasmacytoid dendritic cell neoplasm. Genes Chromosomes and Cancer, 2014, 53, 78-89.	2.8	31
15	Successful treatment of very large congenital infantile fibrosarcoma. Pediatrics International, 2011, 53, 768-770.	0.5	6
16	Novel dominant-negative mutant of GATA3 in HDR syndrome. Journal of Molecular Medicine, 2011, 89, 43-50.	3.9	16
17	The small oligomerization domain of gephyrin converts MLL to an oncogene. Blood, 2004, 103, 3876-3882.	1.4	42