

Kosei Hasegawa

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

407
citations

933447

10
h-index

794594

19
g-index

37
all docs

37
docs citations

37
times ranked

677
citing authors

#	ARTICLE	IF	CITATIONS
1	Heterozygous Orthodontic Homeobox 2 Mutations Are Associated with Variable Pituitary Phenotype. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 756-764.	3.6	98
2	Hypophosphatemic osteomalacia and bone sclerosis caused by a novel homozygous mutation of the FAM20C gene in an elderly man with a mild variant of Raine syndrome. <i>Bone</i> , 2014, 67, 56-62.	2.9	59
3	Clinical Practice Guidelines for Hypophosphatasia*. <i>Clinical Pediatric Endocrinology</i> , 2020, 29, 9-24.	0.8	28
4	Mutation spectrum of <i>COL1A1</i> and <i>COL1A2</i> genes in Indian patients with osteogenesis imperfecta. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 1482-1489.	1.2	24
5	A novel mutation in the COL2A1 gene in a patient with Stickler syndrome type 1: a case report and review of the literature. <i>Journal of Medical Case Reports</i> , 2017, 11, 237.	0.8	22
6	Mutations in type I collagen genes in Japanese osteogenesis imperfecta patients. <i>Pediatrics International</i> , 2007, 49, 564-569.	0.5	21
7	Clinical Practice Guidelines for Achondroplasia*. <i>Clinical Pediatric Endocrinology</i> , 2020, 29, 25-42.	0.8	19
8	Novel heterozygous mutation in <i>TBX1</i> in an infant with hypocalcemic seizures. <i>Clinical Pediatric Endocrinology</i> , 2018, 27, 159-164.	0.8	13
9	Incidence rate and characteristics of symptomatic vitamin D deficiency in children: a nationwide survey in Japan. <i>Endocrine Journal</i> , 2018, 65, 593-599.	1.6	12
10	Growth of infants with osteogenesis imperfecta treated with bisphosphonate. <i>Pediatrics International</i> , 2009, 51, 54-58.	0.5	11
11	A novel mutation p.Ser348Cys in FGFR3 causes achondroplasia. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 1370-1372.	1.2	10
12	HDR syndrome in a Japanese girl with biliary atresia: a case report. <i>BMC Pediatrics</i> , 2016, 16, 14.	1.7	8
13	Three cases of Japanese acromicric/geleophysic dysplasia with FBN1 mutations: a comparison of clinical and radiological features. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2017, 30, 117-121.	0.9	8
14	Novel and recurrent PORCN gene mutations in almost unilateral and typical focal dermal hypoplasia patients. <i>European Journal of Dermatology</i> , 2013, 23, 64-67.	0.6	7
15	Children with short-limbed short stature in pediatric endocrinological services in Japan. <i>Pediatrics International</i> , 2014, 56, 809-812.	0.5	7
16	Acanthosis nigricans in a Japanese boy with hypochondroplasia due to a K650T mutation in <i>FGFR3</i> . <i>Clinical Pediatric Endocrinology</i> , 2017, 26, 223-228.	0.8	7
17	Urinary N-telopeptides of type I collagen in healthy children. <i>Pediatrics International</i> , 2010, 52, 398-401.	0.5	6
18	Pyridoxal in the Cerebrospinal Fluid May Be a Better Indicator of Vitamin B6-dependent Epilepsy Than Pyridoxal 5-Phosphate. <i>Pediatric Neurology</i> , 2020, 113, 33-41.	2.1	6

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19	Impaired pyridinoline cross-link formation in patients with osteogenesis imperfecta. <i>Journal of Bone and Mineral Metabolism</i> , 2008, 26, 394-399.	2.7	5
20	Genetic analysis in Japanese patients with osteogenesis imperfecta: Genotype and phenotype spectra in 96 probands. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1675.	1.2	5
21	Residual endogenous insulin secretion in Japanese children with type 1A diabetes. <i>Clinical Pediatric Endocrinology</i> , 2021, 30, 27-33.	0.8	5
22	Japanese familial case with metaphyseal dysplasia, Schmid Type caused by the p.T555P mutation in the <i>COL10A1</i> gene. <i>Clinical Pediatric Endocrinology</i> , 2015, 24, 33-36.	0.8	4
23	Changes in facial appearance from neonate to adult in 3-M syndrome patient with novel CUL7 gene mutations. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2016, 29, 241-6.	0.9	4
24	A case of pyridoxine-dependent epilepsy with novel ALDH7A1 mutations. <i>Oxford Medical Case Reports</i> , 2020, 2020, omaa008.	0.4	4
25	Thioredoxin interacting protein protects mice from fasting induced liver steatosis by activating ER stress and its downstream signaling pathways. <i>Scientific Reports</i> , 2022, 12, 4819.	3.3	4
26	The clinical course of Rathke's cleft cysts in pediatric patients: impact on growth and pubertal development. <i>Clinical Pediatric Endocrinology</i> , 2022, 31, 38-43.	0.8	3
27	A novel pathogenic variant p.<sc>Asp797Val</sc> in <sc><i>IFIH1</i></sc> in a Japanese boy with overlapping <sc>Singletonâ€Merten</sc> syndrome and <sc>Aicardiâ€GoutiÃˆres</sc> syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2022, 188, 249-252.	1.2	2
28	A Case of Nager Syndrome Diagnosed Before Birth. <i>Acta Medica Okayama</i> , 2019, 73, 273-277.	0.2	2
29	Neonatal-Onset Hereditary Coproporphyrria: A New Variant of Hereditary Coproporphyrria. <i>JIMD Reports</i> , 2017, 37, 99-106.	1.5	1
30	Laboratory changes during adrenocorticotrophic hormone therapy associated with renal calcified lesions. <i>Pediatrics International</i> , 2020, 62, 587-592.	0.5	1
31	Achondroplasia. , 2019, , 145-154.		0
32	Novel <i>AVPR2</i> variant in a male infant with nephrogenic diabetes insipidus who showed delayed head control. <i>Clinical Pediatric Endocrinology</i> , 2019, 28, 155-158.	0.8	0
33	Preclinical diagnosis and identification of the chimeric CYP11B1/CYP11B2 gene in two pediatric cases of a Japanese family with glucocorticoid-remediable aldosteronism. <i>Hypertension Research</i> , 2021, 44, 891-893.	2.7	0
34	A pediatric case of extrapontine myelinolysis due to rapid fluctuation of sodium concentration after craniopharyngioma surgery. <i>Journal of the Japanese Society of Intensive Care Medicine</i> , 2021, 28, 227-229.	0.0	0
35	Transient central diabetes insipidus after cranioplasty for craniosynostosis in an infant with septo-optic dysplasia. <i>Clinical Pediatric Endocrinology</i> , 2022, 31, 50-53.	0.8	0