Kazuki Yamazawa

List of Publications by Year
in descending order

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30	1,354	687363 13	526287 27
papers	citations	h-index	g-index

30	30	30	2366
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	In utero undernourishment perturbs the adult sperm methylome and intergenerational metabolism. Science, 2014, 345, 1255903.	12.6	535
2	Uniparental disomy and human disease: An overview. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2010, 154C, 329-334.	1.6	151
3	Allele-specific binding of ZFP57 in the epigenetic regulation of imprinted and non-imprinted monoallelic expression. Genome Biology, 2015, 16, 112.	8.8	150
4	Silver-Russell syndrome in a girl born after in vitro fertilization: partial hypermethylation at the differentially methylated region of PEG1/MEST. Journal of Assisted Reproduction and Genetics, 2007, 24, 131-136.	2.5	121
5	Molecular and Clinical Studies in 138 Japanese Patients with Silver-Russell Syndrome. PLoS ONE, 2013, 8, e60105.	2.5	55
6	Molecular and clinical findings and their correlations in Silver-Russell syndrome: implications for a positive role of IGF2 in growth determination and differential imprinting regulation of the IGF2â€“H19 domain in bodies and placentas. Journal of Molecular Medicine, 2008, 86, 1171-1181.	3.9	54
7	Monozygotic female twins discordant for Silverâ€“Russell syndrome and hypomethylation of the H19-DMR. Journal of Human Genetics, 2008, 53, 950-955.	2.3	44
8	H1foo Has a Pivotal Role in Qualifying Induced Pluripotent Stem Cells. Stem Cell Reports, 2016, 6, 825-833.	4.8	40
9	Contribution of gene mutations to Silver-Russell syndrome phenotype: multigene sequencing analysis in 92 etiology-unknown patients. Clinical Epigenetics, 2020, 12, 86.	4.1	29
10	Androgenetic/biparental mosaicism in a girl with Beckwithâ€“Wiedemann syndrome-like and upd(14)pat-like phenotypes. Journal of Human Genetics, 2011, 56, 91-93.	2.3	25
11	Molecular and clinical analyses of two patients with UPD(16)mat detected by screening 94 patients with Silver-Russell syndrome phenotype of unknown aetiology. Journal of Medical Genetics, 2019, 56, 413-418.	3.2	23
12	Mutation and Polymorphism Analyses of <i>INSL3</i> and <i>LGR8</i>/<i>GREAT</i> in 62 Japanese Patients with Cryptorchidism. Hormone Research in Paediatrics, 2007, 67, 73-76.	1.8	20
13	Role of Imprinting Disorders in Short Children Born SGA and Silver-Russell Syndrome Spectrum. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 802-813.	3.6	16
14	Genetic heterogeneity of patients with suspected Silver-Russell syndrome: genome-wide copy number analysis in 82 patients without imprinting defects. Clinical Epigenetics, 2017, 9, 52.	4.1	15
15	Hyponatremia, Hypophosphatemia, and Hypouricemia in a Girl With Macrophage Activation Syndrome. Pediatrics, 2006, 118, 2557-2560.	2.1	14
16	Placental hypoplasia in maternal uniparental disomy for chromosome 7. American Journal of Medical Genetics, Part A, 2008, 146A, 514-516.	1.2	11
17	Mosaic upd(7)mat in a patient with Silverâ€“Russell syndrome. American Journal of Medical Genetics, Part A, 2012, 158A, 465-468.	1.2	7
18	Genome-wide methylation analysis in Silverâ€“Russell syndrome, Temple syndrome, and Praderâ€“Willi syndrome. Clinical Epigenetics, 2020, 12, 159.	4.1	7

#	Article	IF	Citations
19	Pathogenic Copy Number and Sequence Variants in Children Born SGA With Short Stature Without Imprinting Disorders. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e3121-e3133.	3.6	7
20	High-level heteroplasmy for the m.7445A>G mitochondrial DNA mutation can cause progressive sensorineural hearing loss in infancy. International Journal of Pediatric Otorhinolaryngology, 2018, 108, 125-131.	1.0	6
21	Loss of imprinting of the human-specific imprinted gene ZNF597 causes prenatal growth retardation and dysmorphic features: implications for phenotypic overlap with Silver-Russell syndrome. Journal of Medical Genetics, 2021, 58, 427-432.	3.2	6
22	A Pediatric Case of Antibiotic-Associated Hemorrhagic Colitis Caused by Klebsiella Oxytoca. Global Pediatric Health, 2014, 1, 2333794X1455052.	0.7	4
23	Spontaneous intramural duodenal hematoma as the manifestation of Noonan syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 496-498.	1.2	4
24	Neonatal intestinal volvulus and preduodenal portal vein associated with situs ambiguus: Report of a case. Surgery Today, 2011, 41, 726-729.	1.5	3
25	Compound heterozygous variants in the ABCG8 gene in a Japanese girl with sitosterolemia. Human Genome Variation, 2020, 7, 25.	0.7	3
26	A novel pathogenic variant of the FH gene in a family with hereditary leiomyomatosis and renal cell carcinoma. Human Genome Variation, 2022, 9, 3.	0.7	3
27	A Familial Case of a Whole Germline <i>CDC73</i> Deletion Discordant for Primary Hyperparathyroidism. Hormone Research in Paediatrics, 2019, 92, 56-63.	1.8	1
28	Metacarpophalangeal pattern profile analysis for a 3â€•monthâ€•old infant with Feingold syndrome 2. American Journal of Medical Genetics, Part A, 2021, 185, 952-954.	1.2	0
29	A Japanese boy with double diagnoses of 2p15p16.1 microdeletion syndrome and RP2-associated retinal disorder. Human Genome Variation, 2021, 8, 46.	0.7	0
30	Frequency and clinical characteristics of distinct etiologies in patients with Silver-Russell syndrome diagnosed based on the Netchine-Harbison clinical scoring system. Journal of Human Genetics, 0, , .	2.3	0