Kazuki Takeda

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9027925/publications.pdf

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22	721	759233 12	677142 22
papers	citations	h-index	g-index

25	25	25	876
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Risk factors for early-onset radiographical adjacent segment disease in patients with spondylolytic spondylolisthesis after single-level posterior lumbar interbody fusion. Spine Journal, 2022, 22, 1112-1118.	1.3	7
2	Polygenic Risk Score of Adolescent Idiopathic Scoliosis for Potential Clinical Use. Journal of Bone and Mineral Research, 2020, 36, 1481-1491.	2.8	5
3	Genome-wide association study identifies 14 previously unreported susceptibility loci for adolescent idiopathic scoliosis in Japanese. Nature Communications, 2019, 10, 3685.	12.8	47
4	Bi-allelic loss of function variants ofTBX6causes a spectrum of malformation of spine and rib including congenital scoliosis and spondylocostal dysostosis. Journal of Medical Genetics, 2019, 56, 622-628.	3.2	13
5	A multiethnic meta-analysis defined the association of rs12946942 with severe adolescent idiopathic scoliosis. Journal of Human Genetics, 2019, 64, 493-498.	2.3	11
6	TBX6-associated congenital scoliosis (TACS) as a clinically distinguishable subtype of congenital scoliosis: further evidence supporting the compound inheritance and TBX6 gene dosage model. Genetics in Medicine, 2019, 21, 1548-1558.	2.4	60
7	An international meta-analysis confirms the association of BNC2 with adolescent idiopathic scoliosis. Scientific Reports, 2018, 8, 4730.	3.3	20
8	Genome-wide meta-analysis and replication studies in multiple ethnicities identify novel adolescent idiopathic scoliosis susceptibility loci. Human Molecular Genetics, 2018, 27, 3986-3998.	2.9	34
9	Screening of known disease genes in congenital scoliosis. Molecular Genetics & Genomic Medicine, 2018, 6, 966-974.	1.2	20
10	A multi-ethnic meta-analysis confirms the association of rs6570507 with adolescent idiopathic scoliosis. Scientific Reports, 2018, 8, 11575.	3.3	33
11	Compound Heterozygosity for Null Mutations and a Common Hypomorphic Risk Haplotype inTBX6Causes Congenital Scoliosis. Human Mutation, 2017, 38, 317-323.	2.5	41
12	Response to Lefebvre et al. Clinical Genetics, 2017, 92, 563-564.	2.0	2
13	A Functional SNP in BNC2 Is Associated with Adolescent Idiopathic Scoliosis. American Journal of Human Genetics, 2015, 97, 337-342.	6.2	119
14	Preemptive Analgesic Effect of Fentanyl on Tourniquet Pain. Pain Medicine, 2007, 8, 618.2-618.	1.9	1
15	ATP Hydrolytic Activity of an Iron-Stimulated P-type ATPase of Mouse Liver Microsomes. Journal of UOEH, 2000, 22, 317-324.	0.6	2
16	Surgical Aspects and Management of Acute Necrotizing Pancreatitis. Pancreas, 1998, 16, 316-322.	1.1	50
17	Sudden death in chronic dialysis patients. Nephrology Dialysis Transplantation, 1997, 12, 952-955.	0.7	70
18	Continuous regional arterial infusion of protease inhibitor and antibiotics in acute necrotizing pancreatitis. American Journal of Surgery, 1996, 171, 394-398.	1.8	160

#	Article	IF	Citations
19	Lymphotoxin and TNF differ greatly in capacity to induce differentiation of human myeloblastic leukemia ML-1 cells. IUBMB Life, 1994, 32, 1109-19.	0.1	2
20	Roles of two tumor necrosis factor receptors in induction of differentiation of ML-1 cells. Anticancer Research, 1993, 13, 883-6.	1.1	2
21	Effect of fibroblast-derived differentiation inducing factor on the differentiation of human monocytoid and myeloid leukemia cell lines. Biochemical and Biophysical Research Communications, 1988, 155, 24-31.	2.1	21
22	Oxidation of NADH by Vanadate Plus Phenylmethanesulfonyl Fluoride (PMSF). Journal of UOEH, 1988, 10, 59-62.	0.6	0