

Kai Yang

List of Publications by Year in descending order

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29
papers

194
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1307366

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32
all docs

32
docs citations

32
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236
citing authors

#	ARTICLE	IF	CITATIONS
1	Novel nanostructured resin infiltrant containing calcium phosphate nanoparticles to prevent enamel white spot lesions. <i>Journal of the Mechanical Behavior of Biomedical Materials</i> , 2022, 126, 104990.	1.5	11
2	ROR2 Downregulation Activates the MSX2/NSUN2/p21 Regulatory Axis and Promotes Dental Pulp Stem Cell Senescence. <i>Stem Cells</i> , 2022, 40, 290-302.	1.4	7
3	Protective Effect of Maternal First-Trimester Low Body Mass Index Against Macrosomia: A 10-Year Cross-Sectional Study. <i>Frontiers in Endocrinology</i> , 2022, 13, 805636.	1.5	2
4	Whole exome sequencing identified a novel compound heterozygous variation in <i>COL7A1</i> gene causing dystrophic epidermolysis bullosa. <i>Molecular Genetics & Genomic Medicine</i> , 2022, , e1907.	0.6	1
5	Identification of novel heterozygous missense variant in the <i>COL11A1</i> causing fetal craniofacial anomalies. <i>International Journal of Transgender Health</i> , 2022, 15, 240-246.	1.1	0
6	Mechanisms of compensatory for cervical lordosis changes after laminectomy with fusion. <i>BMC Surgery</i> , 2022, 22, 129.	0.6	0
7	Relationship between TIA minus C0-7 angle and C2-7 SVA: analysis of 113 symptomatic patients. <i>BMC Musculoskeletal Disorders</i> , 2022, 23, 338.	0.8	3
8	Identification of novel variations in SLC6A8 and GAMT genes causing cerebral creatine deficiency syndrome. <i>Clinica Chimica Acta</i> , 2022, 532, 29-36.	0.5	1
9	Case Report: Exome Sequencing Identified a Novel Compound Heterozygous Variation in PLOD2 Causing Bruck Syndrome Type 2. <i>Frontiers in Genetics</i> , 2021, 12, 619948.	1.1	4
10	Comprehensive strategy improves the genetic diagnosis of different polycystic kidney diseases. <i>Journal of Cellular and Molecular Medicine</i> , 2021, 25, 6318.	1.6	6
11	Clinical research: low-level laser therapy in accelerating orthodontic tooth movement. <i>BMC Oral Health</i> , 2021, 21, 324.	0.8	26
12	Investigation of a Novel LRP6 Variant Causing Autosomal-Dominant Tooth Agenesis. <i>Frontiers in Genetics</i> , 2021, 12, 688241.	1.1	7
13	Downregulation of ROR2 promotes dental pulp stem cell senescence by inhibiting STK4-FOXO1/SMS1 axis in sphingomyelin biosynthesis. <i>Aging Cell</i> , 2021, 20, e13430.	3.0	7
14	Identification of novel variations in the <i>NTRK1</i> gene causing congenital insensitivity to pain with anhidrosis. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1839.	0.6	3
15	Four types of global spine sagittal alignment and compensation mechanism in adult patients with lumbar degenerative disease. <i>Journal of Orthopaedic Science</i> , 2021, , .	0.5	4
16	Metabolic and biophysical study of the MFN2 mutant causing Hereditary Motor and Sensory Neuropathy (HMSN). <i>American Journal of Translational Research (discontinued)</i> , 2021, 13, 11501-11512.	0.0	1
17	Three-dimensional changes in the upper airway and craniomaxillofacial morphology of patients with Angle Class III malocclusion treated with a Frankel III appliance. <i>BMC Oral Health</i> , 2021, 21, 634.	0.8	4
18	Investigation of a Novel NTRK1 Variation Causing Congenital Insensitivity to Pain With Anhidrosis. <i>Frontiers in Genetics</i> , 2021, 12, 763467.	1.1	4

#	ARTICLE	IF	CITATIONS
19	Whole-exome sequencing identified compound heterozygous variants in <i>ROR2</i> gene in a fetus with Robinow syndrome. <i>Journal of Clinical Laboratory Analysis</i> , 2020, 34, e23074.	0.9	9
20	Regulating Oral Biofilm from Cariogenic State to Non-Cariogenic State via Novel Combination of Bioactive Therapeutic Composite and Gene-Knockout. <i>Microorganisms</i> , 2020, 8, 1410.	1.6	3
21	Mean platelet volume in the first trimester as a predictor of gestational diabetes mellitus. <i>Chinese Medical Journal</i> , 2020, 133, 1364-1365.	0.9	0
22	A novel splicing mutation in the <i>PROS1</i> gene causes hereditary protein S deficiency in a Chinese family with thrombotic disease. <i>Thrombosis Research</i> , 2020, 189, 93-95.	0.8	2
23	A novel frameshift variant in <i>SON</i> causes Zhu-Tokita-Takenouchi-Kim Syndrome. <i>Journal of Clinical Laboratory Analysis</i> , 2020, 34, e23326.	0.9	11
24	Study on the correlation and predictive value of serum pregnancy-associated plasma protein A, triglyceride and serum 25-hydroxyvitamin D levels with gestational diabetes mellitus. <i>World Journal of Clinical Cases</i> , 2020, 8, 864-873.	0.3	8
25	A clinical and multi-omics study of Van der Woude syndrome in three generations of a Chinese family. <i>Molecular Medicine Reports</i> , 2020, 22, 2925-2931.	1.1	0
26	Transit amplifying cells coordinate mouse incisor mesenchymal stem cell activation. <i>Nature Communications</i> , 2019, 10, 3596.	5.8	31
27	Genetic Analysis in Fetal Skeletal Dysplasias by Trio Whole-Exome Sequencing. <i>BioMed Research International</i> , 2019, 2019, 1-8.	0.9	27
28	Comparative Study of 660 and 830nm Photobiomodulation in Promoting Orthodontic Tooth Movement. <i>Photobiomodulation, Photomedicine, and Laser Surgery</i> , 2019, 37, 349-355.	0.7	12
29	Research on picosecond laser refurbishing method for reconditioning ceramic orthodontic brackets. <i>Journal of Laser Applications</i> , 2014, 26, 022006.	0.8	0