

Yaser Alkhiary

List of Publications by Year in descending order

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Version: 2024-02-01

10
papers

641
citations

1163117

8
h-index

1372567

10
g-index

10
all docs

10
docs citations

10
times ranked

901
citing authors

#	ARTICLE	IF	CITATIONS
1	Enhancement of Experimental Fracture-Healing by Systemic Administration of Recombinant Human Parathyroid Hormone (PTH 1-34). <i>Journal of Bone and Joint Surgery - Series A</i> , 2005, 87, 731-741.	3.0	231
2	Three-dimensional Reconstruction of Fracture Callus Morphogenesis. <i>Journal of Histochemistry and Cytochemistry</i> , 2006, 54, 1215-1228.	2.5	164
3	Selective and Nonselective Cyclooxygenase-2 Inhibitors and Experimental Fracture-Healing. <i>Journal of Bone and Joint Surgery - Series A</i> , 2007, 89, 114-125.	3.0	106
4	Effects of the local mechanical environment on vertebrate tissue differentiation during repair: does repair recapitulate development?. <i>Journal of Experimental Biology</i> , 2003, 206, 2459-2471.	1.7	52
5	Effect of lip position and gingival display on smile and esthetics as perceived by college students with different educational backgrounds. <i>Clinical, Cosmetic and Investigational Dentistry</i> , 2013, 5, 77.	1.6	26
6	Effects of acid hydrolysis and mechanical polishing on surface residual stresses of low-fusing dental ceramics. <i>Journal of Prosthetic Dentistry</i> , 2003, 90, 133-142.	2.8	21
7	A novel homozygous PTH1R variant identified through whole-exome sequencing further expands the clinical spectrum of primary failure of tooth eruption in a consanguineous Saudi family. <i>Archives of Oral Biology</i> , 2016, 67, 28-33.	1.8	21
8	Effect of Clasp Design on Retention at Different Intervals Using Different Abutment Materials and in a Simulated Oral Condition. <i>Journal of Prosthodontics</i> , 2014, 23, 140-145.	3.7	11
9	Whole-exome sequencing reveals a recurrent mutation in the cathepsin C gene that causes Papillonâ€“Lefevre syndrome in a Saudi family. <i>Saudi Journal of Biological Sciences</i> , 2016, 23, 571-576.	3.8	6
10	Identification of Two Homozygous Sequence Variants in the <i>COL7A1</i> Gene Underlying Dystrophic Epidermolysis Bullosa by Wholeâ€“Exome Analysis in a Consanguineous Family. <i>Annals of Human Genetics</i> , 2015, 79, 350-356.	0.8	3