

Puppala Venkat Ramchander

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8246440/publications.pdf>

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papers

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docs citations

9
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89
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic Association of rs1021188 and DNA Methylation Signatures of TNFSF11 in the Risk of Conductive Hearing Loss. <i>Frontiers in Medicine</i> , 2022, 9, 870244.	1.2	0
2	The risks of RELN polymorphisms and its expression in the development of otosclerosis. <i>PLoS ONE</i> , 2022, 17, e0269558.	1.1	3
3	Association of <i>ISL1</i> polymorphisms and eosinophilic levels among otitis media patients. <i>Journal of Clinical Laboratory Analysis</i> , 2021, 35, e23702.	0.9	2
4	Evaluation of the Genetic Association and mRNA Expression of the <i>COL1A1</i> , <i>BMP2</i> , and <i>BMP4</i> Genes in the Development of Otosclerosis. <i>Genetic Testing and Molecular Biomarkers</i> , 2020, 24, 343-351.	0.3	5
5	Osteoprotegerin gene polymorphisms and otosclerosis: an additional genetic association study, multilocus interaction and meta-analysis. <i>BMC Medical Genetics</i> , 2020, 21, 122.	2.1	8
6	Otosclerosis Associated with a De Novo Mutation $\hat{\sim}832G\hat{\sim}$ in the <i>TGFB1</i> Gene Promoter Causing a Decreased Expression Level. <i>Scientific Reports</i> , 2016, 6, 29572.	1.6	9
7	Genetic Association and Altered Gene Expression of Osteoprotegerin in Otosclerosis Patients. <i>Annals of Human Genetics</i> , 2015, 79, 225-237.	0.3	13
8	Genetic Association and Gene Expression Profiles of <i>TGFB1</i> and the Contribution of <i>TGFB1</i> to Otosclerosis Susceptibility. <i>Journal of Bone and Mineral Research</i> , 2013, 28, 2490-2497.	3.1	16
9	Mutations in the Connexin 29 Gene Are Not a Major Cause of Nonsyndromic Hearing Impairment in India. <i>Genetic Testing and Molecular Biomarkers</i> , 2010, 14, 539-541.	0.3	3