

# Mutations in LRP5 or FZD4 Underlie the Common Familial Exostosis Locus on Chromosome 11q

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Citation Report

#	ARTICLE	IF	CITATIONS
1	WNT and $\beta$ -catenin signalling: diseases and therapies. <i>Nature Reviews Genetics</i> , 2004, 5, 691-701.	7.7	1,675
2	Wnt signaling in osteoblasts and bone diseases. <i>Gene</i> , 2004, 341, 19-39.	1.0	724
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4	Autosomal Recessive Familial Exudative Vitreoretinopathy Is Associated with Mutations in LRP5. <i>American Journal of Human Genetics</i> , 2004, 75, 878-884.	2.6	174
5	Pathogenic mutations and polymorphisms in the lipoprotein receptor-related protein 5 reveal a new biological pathway for the control of bone mass. <i>Current Opinion in Lipidology</i> , 2005, 16, 207-214.	1.2	47
7	The Wnt Signaling Pathway in Retinal Degenerations. <i>IUBMB Life</i> , 2005, 57, 381-388.	1.5	51
8	Genetic variants of frizzled-4 gene in familial exudative vitreoretinopathy and advanced retinopathy of prematurity. <i>Clinical Genetics</i> , 2005, 67, 363-366.	1.0	49
9	The role of the cysteine-rich domain of Frizzled in Wingless-Armadillo signaling. <i>EMBO Journal</i> , 2005, 24, 3493-3503.	3.5	52
10	LRP5 mutations in osteoporosis-pseudoglioma syndrome and high-bone-mass disorders. <i>Joint Bone Spine</i> , 2005, 72, 207-214.	0.8	80
11	Cell-surface co-receptors: emerging roles in signaling and human disease. <i>Trends in Biochemical Sciences</i> , 2005, 30, 611-621.	3.7	68
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19	Role of the Norrie Disease Pseudoglioma Gene in Sprouting Angiogenesis during Development of the Retinal Vasculature. , 2005, 46, 3372.		129

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