

Mutations in the TGF- β 2 repressor SKI cause Shprintzen aneurysm

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Connective Tissue Disorders. , 0, , 537-546.		0
2	The Cardiac Society of Australia and New Zealand. Circulation Journal, 2012, 76, 1051-1053.	0.7	1
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5	TGF- β^2 and Cardiovascular Disorders. , 2013, , 297-322.		1
6	Molecular mechanisms of thoracic aortic dissection. Journal of Surgical Research, 2013, 184, 907-924.	0.8	182
7	Severe congenital lipodystrophy and a progeroid appearance: Mutation in the penultimate exon of <i>FBN1</i> causing a recognizable phenotype. American Journal of Medical Genetics, Part A, 2013, 161, 3057-3062.	0.7	56
8	Fine Mapping of the 1p36 Deletion Syndrome Identifies Mutation of PRDM16 as a Cause of Cardiomyopathy. American Journal of Human Genetics, 2013, 93, 67-77.	2.6	164
9	Genetic biomarkers in aortopathy. Biomarkers in Medicine, 2013, 7, 547-563.	0.6	6
10	576kb deletion in 1p36.33-36.32 containing SKI is associated with limb malformation, congenital heart disease and epilepsy. Gene, 2013, 528, 352-355.	1.0	27
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