

# The Mutational Spectrum of the Sonic Hedgehog Gene in Mutations Cause a Significant Proportion of Autosomal

Human Molecular Genetics

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Hemophilia B in a 46,XX female probably caused by nonrandom X inactivation. <i>Clinical Genetics</i> , 1993, 43, 1-4.	1.0	17
2	Molecular Mechanisms of Holoprosencephaly. <i>Molecular Genetics and Metabolism</i> , 1999, 68, 126-138.	0.5	121
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6	The structure and function of genes causing human holoprosencephaly. <i>Gene Function &amp; Disease</i> , 2000, 1, 7-20.	0.3	3
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8	Mutations in TGIF cause holoprosencephaly and link NODAL signalling to human neural axis determination. <i>Nature Genetics</i> , 2000, 25, 205-208.	9.4	368
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18	Molecular Genetics of Holoprosencephaly. <i>Fetal and Pediatric Pathology</i> , 2000, 19, 1-19.	0.3	0
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21	Prenatal diagnosis of partial monosomy 18p(18p11.2?pter) and trisomy 21q(21q22.3?qter) with alobar holoprosencephaly and premaxillary agenesis. <i>Prenatal Diagnosis</i> , 2001, 21, 346-350.	1.1	28
22	Expression of sonic hedgehog downstream genes is modified in rat embryos exposed in utero to a distal inhibitor of cholesterol biosynthesis. <i>Developmental Dynamics</i> , 2001, 220, 99-111.	0.8	25
23	SHH mutation is associated with solitary median maxillary central incisor: A study of 13 patients and review of the literature. <i>American Journal of Medical Genetics Part A</i> , 2001, 102, 1-10.	2.4	157
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42	Mosaic trisomy 9 and lobar holoprosencephaly. American Journal of Medical Genetics Part A, 2002, 111, 295-300.	2.4	16
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