

Mutations in PAX1 may be associated with Klippel-Feil syndrome

European Journal of Human Genetics

11, 468-474

DOI: [10.1038/sj.ejhg.5200987](https://doi.org/10.1038/sj.ejhg.5200987)

Citation Report

#	ARTICLE	IF	CITATIONS
1	A 38-year-old female with chronic cervicgia: case discussion. Clinical Chiropractic, 2004, 7, 100-100.	0.1	0
2	A 38-year-old female with chronic cervicgia: case discussion. Clinical Chiropractic, 2004, 7, 100-104.	0.1	0
3	An analysis of <i>PAX1</i> in the development of vertebral malformations. Clinical Genetics, 2005, 68, 448-453.	2.0	64
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6	DLL3 as a candidate gene for vertebral malformations. American Journal of Medical Genetics, Part A, 2006, 140A, 2447-2453.	1.2	36
7	Association of methylenetetrahydrofolate reductase genetic polymorphisms with atlantoaxial dislocation. Journal of Neurosurgery: Spine, 2007, 7, 623-630.	1.7	8
8	Ruptured Aneurysm of the Sinus of Valsalva With Wildervanck Syndrome (Cervico-Oculo-Acoustic) Tj ETQq1 1 0.784314 rgBTg /Overlo	1.6	8
9	Lack of evidence of WNT3A as a candidate gene for congenital vertebral malformations. Scoliosis, 2007, 2, 13.	0.4	19
10	PAX genes: Roles in development, pathophysiology, and cancer. Biochemical Pharmacology, 2007, 73, 1-14.	4.4	239
11	Mutations in GDF6 are associated with vertebral segmentation defects in Klippel-Feil syndrome. Human Mutation, 2008, 29, 1017-1027.	2.5	170
12	The basal chordate amphioxus as a simple model for elucidating developmental mechanisms in vertebrates. Birth Defects Research Part C: Embryo Today Reviews, 2008, 84, 175-187.	3.6	34
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17	Progress in the Understanding of the Genetic Etiology of Vertebral Segmentation Disorders in Humans. Annals of the New York Academy of Sciences, 2009, 1151, 38-67.	3.8	70
18	Quantitative Analysis of Methylation Status of the PAX1 Gene for Detection of Cervical Cancer. International Journal of Gynecological Cancer, 2010, 20, 513-519.	2.5	42

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19	Characterization of the human nucleus pulposus cell phenotype and evaluation of novel marker gene expression to define adult stem cell differentiation. <i>Arthritis and Rheumatism</i> , 2010, 62, 3695-3705.	6.7	194
20	Molecular basis of androgenetic alopecia: From androgen to paracrine mediators through dermal papilla. <i>Journal of Dermatological Science</i> , 2011, 61, 1-6.	1.9	147
21	Can mutations in the ribosomal protein S26 (RPS26) gene lead to Klippel-Feil syndrome in Diamond-Blackfan anemia patients? An update from the Czech Diamond-Blackfan Anemia registry. <i>Blood Cells, Molecules, and Diseases</i> , 2011, 46, 300-301.	1.4	10
22	Influence of Porcine Intervertebral Disc Matrix on Stem Cell Differentiation. <i>Journal of Functional Biomaterials</i> , 2011, 2, 155-172.	4.4	10
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58	Expanding the neurodevelopmental phenotypes of individuals with de novo KMT2A variants. <i>Npj Genomic Medicine</i> , 2019, 4, 9.	3.8	29
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