

# A Mutation in the Fibroblast Growth Factor 14 Gene Is A Dominant Cerebral Ataxia

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Dominant ataxias. , 2010, , 242-283.		0
2	Dominantly inherited ataxias. <i>Seminars in Pediatric Neurology</i> , 2003, 10, 210-222.	1.0	10
3	Paediatric and adult ataxias (update). <i>European Journal of Paediatric Neurology</i> , 2003, 7, 231-233.	0.7	1
4	Do CTG expansions at the SCA8 locus cause ataxia?. <i>Annals of Neurology</i> , 2003, 54, 110-115.	2.8	50
5	Discoveries in sphingolipid metabolism, spinocerebellar ataxia and autoimmune disease. <i>Clinical Genetics</i> , 2003, 64, 1-3.	1.0	0
6	A new mechanism for spinocerebellar ataxia involving mutations in protein kinase C $\delta$ . <i>Clinical Genetics</i> , 2003, 64, 3-4.	1.0	1
7	Can we throw the master-switch in autoimmune disease?. <i>Clinical Genetics</i> , 2003, 64, 5-6.	1.0	0
8	Fibroblast Growth Factor (FGF) Homologous Factors Share Structural but Not Functional Homology with FGFs. <i>Journal of Biological Chemistry</i> , 2003, 278, 34226-34236.	1.6	221
9	Missense Mutations in the Regulatory Domain of PKC $\delta$ : A New Mechanism for Dominant Nonepisodic Cerebellar Ataxia. <i>American Journal of Human Genetics</i> , 2003, 72, 839-849.	2.6	236
10	Identification of a novel SCA14 mutation in a Dutch autosomal dominant cerebellar ataxia family. <i>Neurology</i> , 2003, 61, 1760-1765.	1.5	97
11	Dominant spinocerebellar ataxias: a molecular approach to classification, diagnosis, pathogenesis and the future. <i>Expert Review of Molecular Diagnostics</i> , 2003, 3, 715-732.	1.5	12
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13	Title is missing!. <i>Current Opinion in Neurology</i> , 2003, 16, 451-458.	1.8	4
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15	Molecular Genetics of Hereditary Spinocerebellar Ataxia. <i>Archives of Neurology</i> , 2004, 61, 727.	4.9	130
16	Mapping of the SCA23 locus involved in autosomal dominant cerebellar ataxia to chromosome region 20p13-12.3. <i>Brain</i> , 2004, 127, 2551-2557.	3.7	79
17	Protein kinase C gamma mutations in spinocerebellar ataxia 14 increase kinase activity and alter membrane targeting. <i>Brain</i> , 2004, 128, 436-442.	3.7	70
18	A novel neurodegenerative disease characterised by posterior column ataxia and pyramidal tract involvement maps to chromosome 8p12-8q12.1. <i>Journal of Medical Genetics</i> , 2004, 41, 634-639.	1.5	13

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20	Mutations in the <i>FGF14</i> gene are not a major cause of spinocerebellar ataxia in Caucasians. <i>Neurology</i> , 2004, 63, 936-936.	1.5	12
21	Autosomal dominant cerebellar ataxias: clinical features, genetics, and pathogenesis. <i>Lancet Neurology</i> , The, 2004, 3, 291-304.	4.9	963
22	Fine mapping of 16q-linked autosomal dominant cerebellar ataxia type III in Japanese families. <i>Neurogenetics</i> , 2004, 5, 215-221.	0.7	14
23	Spinocerebellar ataxia with sensory neuropathy (SCA25) maps to chromosome 2p. <i>Annals of Neurology</i> , 2004, 55, 97-104.	2.8	78
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26	Hereditary ataxias. <i>Handbook of Clinical Neurophysiology</i> , 2004, 4, 655-673.	0.0	1
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43	Fibroblast growth factor homologous factors: Evolution, structure, and function. <i>Cytokine and Growth Factor Reviews</i> , 2005, 16, 215-220.	3.2	184
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53	Identification of a new family of spinocerebellar ataxia type 14 in the japanese spinocerebellar ataxia population by the screening of PRKCG exon 4. <i>Movement Disorders</i> , 2006, 21, 1355-1360.	2.2	29
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156	Genetics and Clinical Features of Inherited Ataxias. , 2015, , 939-978.		1
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