

Germline loss-of-function mutations in LZTR1 predispose to multiple schwannomas

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Citation Report

#	ARTICLE	IF	CITATIONS
1	A germline missense mutation in COQ6 is associated with susceptibility to familial schwannomatosis. <i>Genetics in Medicine</i> , 2014, 16, 787-792.	1.1	20
2	Schwannomas and Their Pathogenesis. <i>Brain Pathology</i> , 2014, 24, 205-220.	2.1	151
3	Premature termination of SMARCB1 translation may be followed by reinitiation in schwannomatosis-associated schwannomas, but results in absence of SMARCB1 expression in rhabdoid tumors. <i>Acta Neuropathologica</i> , 2014, 128, 439-448.	3.9	23
4	Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained after excluding SMARCB1 and LZTR1 germline variants. <i>Acta Neuropathologica</i> , 2014, 128, 449-452.	3.9	36
5	SMARCB1 mutations in schwannomatosis and genotype correlations with rhabdoid tumors. <i>Cancer Genetics</i> , 2014, 207, 373-378.	0.2	71
6	Concurrent DNA Copy-Number Alterations and Mutations in Genes Related to Maintenance of Genome Stability in Uninvolved Mammary Glandular Tissue from Breast Cancer Patients. <i>Human Mutation</i> , 2015, 36, 1088-1099.	1.1	11
7	Neurofibromatosis type 2. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2015, 132, 87-96.	1.0	45
8	Immortalized Human Schwann Cell Lines Derived From Tumors of Schwannomatosis Patients. <i>PLoS ONE</i> , 2015, 10, e0144620.	1.1	10
9	Mutations in <i>LZTR1</i> add to the complex heterogeneity of schwannomatosis. <i>Neurology</i> , 2015, 84, 141-147.	1.5	90
10	Synchronous Hodgkin's lymphoma and schwannomas mimicking refractory Hodgkin's lymphoma. <i>Annals of Hematology</i> , 2015, 94, 1593-1595.	0.8	0
11	Diagnosis, Management, and New Therapeutic Options in Childhood Neurofibromatosis Type 2 and Related Forms. <i>Seminars in Pediatric Neurology</i> , 2015, 22, 240-258.	1.0	68
12	Is there a link between COQ6 and schwannomatosis?. <i>Genetics in Medicine</i> , 2015, 17, 312-313.	1.1	7
13	Response to Trevisson et al.. <i>Genetics in Medicine</i> , 2015, 17, 313-314.	1.1	1
14	Mosaic Neurocutaneous Disorders and Their Causes. <i>Seminars in Pediatric Neurology</i> , 2015, 22, 207-233.	1.0	81
15	Hereditary Predisposition to Primary CNS Tumors. <i>Molecular Pathology Library</i> , 2015, , 1-22.	0.1	0
16	Cancer genomics: why rare is valuable. <i>Journal of Molecular Medicine</i> , 2015, 93, 369-381.	1.7	8
17	Familial schwannomatosis with a germline mutation of SMARCB1 in Japan. <i>Brain Tumor Pathology</i> , 2015, 32, 216-220.	1.1	11
18	Segmental neurofibromatosis type 2: discriminating two hit from four hit in a patient presenting multiple schwannomas confined to one limb. <i>BMC Medical Genomics</i> , 2015, 8, 2.	0.7	24

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19	Broadening the spectrum of SMARCB1-associated malignant tumors: a case of uterine leiomyosarcoma in a patient with schwannomatosis. <i>Human Pathology</i> , 2015, 46, 1226-1231.	1.1	10
20	Pediatric Peripheral Nerve Tumors. , 2015, , 839-846.		4
21	Expanding schwannomatosis phenotype. <i>Journal of Neuro-Oncology</i> , 2015, 122, 607-609.	1.4	5
22	Rare variants in <i>SOS2</i> and <i>LZTR1</i> are associated with Noonan syndrome. <i>Journal of Medical Genetics</i> , 2015, 52, 413-421.	1.5	187
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