

Exome sequencing identifies MAX mutations as a cause

Nature Genetics

43, 663-667

DOI: 10.1038/ng.861

Citation Report

#	ARTICLE	IF	CITATIONS
5	Genetics and clinical characteristics of hereditary pheochromocytomas and paragangliomas. <i>Endocrine-Related Cancer</i> , 2011, 18, R253-R276.	3.1	299
6	Exome Sequencing Identifies Rare Deleterious Mutations in DNA Repair Genes FANCC and BLM as Potential Breast Cancer Susceptibility Alleles. <i>PLoS Genetics</i> , 2012, 8, e1002894.	3.5	186
7	Epithelial to Mesenchymal Transition Is Activated in Metastatic Pheochromocytomas and Paragangliomas Caused by SDHB Gene Mutations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012, 97, E954-E962.	3.6	87
8	Mitochondrial function and content in pheochromocytoma/paraganglioma of succinate dehydrogenase mutation carriers. <i>Endocrine-Related Cancer</i> , 2012, 19, 261-269.	3.1	23
9	False-negative 123I-MIBG SPECT is most commonly found in SDHB-related pheochromocytoma or paraganglioma with high frequency to develop metastatic disease. <i>Endocrine-Related Cancer</i> , 2012, 19, 83-93.	3.1	106
10	Combined blockade of signalling pathways shows marked anti-tumour potential in phaeochromocytoma cell lines. <i>Journal of Molecular Endocrinology</i> , 2012, 49, 79-96.	2.5	44
11	Head and neck paragangliomas: genetic spectrum and clinical variability in 79 consecutive patients. <i>Endocrine-Related Cancer</i> , 2012, 19, 149-155.	3.1	71
12	ENDOCRINE DISORDERS IN PREGNANCY: Pheochromocytoma and pregnancy: a deceptive connection. <i>European Journal of Endocrinology</i> , 2012, 166, 143-150.	3.7	122
13	GENETICS IN ENDOCRINOLOGY: The genetics of phaeochromocytoma: using clinical features to guide genetic testing. <i>European Journal of Endocrinology</i> , 2012, 166, 151-158.	3.7	98
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17	Somatic NF1 inactivation is a frequent event in sporadic pheochromocytoma. <i>Human Molecular Genetics</i> , 2012, 21, 5397-5405.	2.9	126
18	Trp53 inactivation leads to earlier phaeochromocytoma formation in pten knockout mice. <i>Endocrine-Related Cancer</i> , 2012, 19, 731-740.	3.1	3
19	Pheochromocytoma and Paraganglioma: Recent Progress and New Vistas for Improved Patient Care. <i>Hormone and Metabolic Research</i> , 2012, 44, 325-327.	1.5	8
20	Current and Future Anatomical and Functional Imaging Approaches to Pheochromocytoma and Paraganglioma. <i>Hormone and Metabolic Research</i> , 2012, 44, 367-372.	1.5	71
21	A Decade (2001-2010) of Genetic Testing for Pheochromocytoma and Paraganglioma. <i>Hormone and Metabolic Research</i> , 2012, 44, 359-366.	1.5	103
22	Peptide Receptor Radionuclide Therapy (PRRT) with 177Lu-DOTATATE in Individuals with Neck or Mediastinal Paraganglioma (PGL). <i>Hormone and Metabolic Research</i> , 2012, 44, 411-414.	1.5	71

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64	An update on the genetics of pheochromocytoma. <i>Journal of Human Hypertension</i> , 2013, 27, 141-147.	2.2	54
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74	Translational Research in Endocrine Surgery. <i>Surgical Oncology Clinics of North America</i> , 2013, 22, 857-884.	1.5	11
75	Gestational diastolic hypertension with gene mutation-related pheochromocytoma positive at 18F-DOPA PET/CT: Diagnostic and therapeutic implications. <i>Revista Espanola De Medicina Nuclear E Imagen Molecular</i> , 2013, 32, 111-112.	0.0	1
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80	10 rare tumors that warrant a genetics referral. <i>Familial Cancer</i> , 2013, 12, 1-18.	1.9	24
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