

Exome sequencing identifies MAX mutations as a cause

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Citation Report

#	ARTICLE	IF	CITATIONS
5	Genetics and clinical characteristics of hereditary pheochromocytomas and paragangliomas. <i>Endocrine-Related Cancer</i> , 2011, 18, R253-R276.	1.6	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.	1.5	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.	1.8	87
8	Mitochondrial function and content in pheochromocytoma/paraganglioma of succinate dehydrogenase mutation carriers. <i>Endocrine-Related Cancer</i> , 2012, 19, 261-269.	1.6	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.	1.6	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.	1.1	44
11	Head and neck paragangliomas: genetic spectrum and clinical variability in 79 consecutive patients. <i>Endocrine-Related Cancer</i> , 2012, 19, 149-155.	1.6	71
12	ENDOCRINE DISORDERS IN PREGNANCY: Pheochromocytoma and pregnancy: a deceptive connection. <i>European Journal of Endocrinology</i> , 2012, 166, 143-150.	1.9	122
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17	Somatic NF1 inactivation is a frequent event in sporadic pheochromocytoma. <i>Human Molecular Genetics</i> , 2012, 21, 5397-5405.	1.4	126
18	Trp53 inactivation leads to earlier phaeochromocytoma formation in pten knockout mice. <i>Endocrine-Related Cancer</i> , 2012, 19, 731-740.	1.6	3
19	Pheochromocytoma and Paraganglioma: Recent Progress and New Vistas for Improved Patient Care. <i>Hormone and Metabolic Research</i> , 2012, 44, 325-327.	0.7	8
20	Current and Future Anatomical and Functional Imaging Approaches to Pheochromocytoma and Paraganglioma. <i>Hormone and Metabolic Research</i> , 2012, 44, 367-372.	0.7	71
21	A Decade (2001-2010) of Genetic Testing for Pheochromocytoma and Paraganglioma. <i>Hormone and Metabolic Research</i> , 2012, 44, 359-366.	0.7	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.	0.7	71

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33	Identical germline mutations in the TMEM127 gene in two unrelated Japanese patients with bilateral pheochromocytoma. <i>Clinical Endocrinology</i> , 2012, 77, 707-714.	1.2	14
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45	Improving Indel Detection Specificity of the Ion Torrent PGM Benchtop Sequencer. <i>PLoS ONE</i> , 2012, 7, e45798.	1.1	48
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