Patricia L M Dahia

List of Publications by Year
in descending order

51 papers	7,420 citations	172457 29 h-index	197818 49 g-index
69 all docs	69 docs citations	69 times ranked	6671 citing authors

#	Article	IF	Citations
1	Endocrine and Neuroendocrine Tumors Special Issueâ€"Checkpoint Inhibitors for Adrenocortical Carcinoma and Metastatic Pheochromocytoma and Paraganglioma: Do They Work?. Cancers, 2022, 14, 467.	3.7	18
2	Thyroid nodules of indeterminate cytology in Hispanic/Latinx patients. Head and Neck, 2022, , .	2.0	2
3	Genotype-Phenotype Features of Germline Variants of the TMEM127 Pheochromocytoma Susceptibility Gene: A 10-Year Update. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e350-e364.	3.6	8
4	The North American Neuroendocrine Tumor Society Consensus Guidelines for Surveillance and Management of Metastatic and/or Unresectable Pheochromocytoma and Paraganglioma. Pancreas, 2021, 50, 469-493.	1.1	55
5	International consensus on initial screening and follow-up of asymptomatic SDHx mutation carriers. Nature Reviews Endocrinology, 2021, 17, 435-444.	9.6	80
6	Insights into Mechanisms of Pheochromocytomas and Paragangliomas Driven by Known or New Genetic Drivers. Cancers, 2021, 13, 4602.	3.7	11
7	Functional Characterization of TMEM127 Variants Reveals Novel Insights into Its Membrane Topology and Trafficking. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e3142-e3156.	3.6	8
8	MYC Regulation of D2HGDH and L2HGDH Influences the Epigenome and Epitranscriptome. Cell Chemical Biology, 2020, 27, 538-550.e7.	5.2	14
9	Recognizing hypoxia in phaeochromocytomas and paragangliomas. Nature Reviews Endocrinology, 2020, 16, 191-192.	9.6	9
10	HEREDITARY ENDOCRINE TUMOURS: CURRENT STATE-OF-THE-ART AND RESEARCH OPPORTUNITIES: Metastatic pheochromocytomas and paragangliomas: proceedings of the MEN2019 workshop. Endocrine-Related Cancer, 2020, 27, T41-T52.	3.1	33
11	Genetic predisposition to neural crest-derived tumors: revisiting the role of KIF1B. Endocrine Connections, 2020, 9, 1042-1050.	1.9	1
12	Genetic predisposition to neural crest-derived tumors: revisiting the role of KIF1B. Endocrine Connections, 2020, 9, 1042-1050.	1.9	5
13	Synonymous but Not Silent: A Synonymous VHL Variant in Exon 2 Confers Susceptibility to Familial Pheochromocytoma and von Hippel-Lindau Disease. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 3826-3834.	3.6	17
14	The tumor suppressor TMEM127 regulates insulin sensitivity in a tissue-specific manner. Nature Communications, 2019, 10, 4720.	12.8	14
15	A MYC Driven and Metabolite Executed Axis Controls the Subcellular Localization and Activity of TET DNA Hydroxylases and RNA Demethylases in B Cell Lymphomas. Blood, 2019, 134, 114-114.	1.4	0
16	EPAS1 Mutations and Paragangliomas in Cyanotic Congenital Heart Disease. New England Journal of Medicine, 2018, 378, 1259-1261.	27.0	41
17	The TMEM127 human tumor suppressor is a component of the mTORC1 lysosomal nutrient-sensing complex. Human Molecular Genetics, 2018, 27, 1794-1808.	2.9	18
18	A Unique Case of Metastatic, Functional, Hereditary Paraganglioma Associated With anSDHC Germline Mutation. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 2802-2806.	3.6	6

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19	Pheochromocytomas and Paragangliomas, Genetically Diverse and Minimalist, All at Once!. Cancer Cell, 2017, 31, 159-161.	16.8	30
20	IDH Mutation, Competitive Inhibition of FTO, and RNA Methylation. Cancer Cell, 2017, 31, 619-620.	16.8	65
21	Molecular and phenotypic evaluation of a novel germline TMEM127 mutation with an uncommon clinical presentation. Endocrine-Related Cancer, 2017, 24, L79-L82.	3.1	8
22	Consensus Statement on next-generation-sequencing-based diagnostic testing of hereditary phaeochromocytomas and paragangliomas. Nature Reviews Endocrinology, 2017, 13, 233-247.	9.6	198
23	Predictors of outcome in phaeochromocytomas and paragangliomas. F1000Research, 2017, 6, 2160.	1.6	3
24	Recurrent Mutations of Chromatin-Remodeling Genes and Kinase Receptors in Pheochromocytomas and Paragangliomas. Clinical Cancer Research, 2016, 22, 2301-2310.	7.0	136
25	Penetrance and Clinical Features of Pheochromocytoma in a Six-Generation Family Carrying a Germline TMEM127 Mutation. Journal of Clinical Endocrinology and Metabolism, 2015, 100, E308-E318.	3.6	44
26	D2HGDH regulates alpha-ketoglutarate levels and dioxygenase function by modulating IDH2. Nature Communications, 2015, 6, 7768.	12.8	64
27	Pheochromocytoma and paraganglioma pathogenesis: learning from genetic heterogeneity. Nature Reviews Cancer, 2014, 14, 108-119.	28.4	442
28	Nextâ€generation sequencing for the genetic screening of phaeochromcytomas and paragangliomas: riding the new wave, but with caution. Clinical Endocrinology, 2014, 80, 23-24.	2.4	6
29	A Novel Germline Mutation in BAP1 Predisposes to Familial Clear-Cell Renal Cell Carcinoma. Molecular Cancer Research, 2013, 11, 1061-1071.	3.4	135
30	Novel Hereditary Forms of Pheochromocytomas and Paragangliomas. Frontiers of Hormone Research, 2013, 41, 79-91.	1.0	13
31	In vivo and in vitro oncogenic effects of HIF2A mutations in pheochromocytomas and paragangliomas. Endocrine-Related Cancer, 2013, 20, 349-359.	3.1	110
32	MAX Mutations Cause Hereditary and Sporadic Pheochromocytoma and Paraganglioma. Clinical Cancer Research, 2012, 18, 2828-2837.	7.0	277
33	Defects in succinate dehydrogenase in gastrointestinal stromal tumors lacking KIT and PDGFRA mutations. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 314-318.	7.1	574
34	Minireview: The Busy Road to Pheochromocytomas and Paragangliomas Has a New Member, TMEM127. Endocrinology, 2011, 152, 2133-2140.	2.8	42
35	Germline mutations in TMEM127 confer susceptibility to pheochromocytoma. Nature Genetics, 2010, 42, 229-233.	21.4	364
36	Mutations of the Metabolic GenesIDH1,IDH2, andSDHAF2Are Not Major Determinants of the Pseudohypoxic Phenotype of Sporadic Pheochromocytomas and Paragangliomas. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1469-1472.	3.6	25

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37	VHL Disease. Best Practice and Research in Clinical Endocrinology and Metabolism, 2010, 24, 401-413.	4.7	53
38	Spectrum and Prevalence of FP/TMEM127 Gene Mutations in Pheochromocytomas and Paragangliomas. JAMA - Journal of the American Medical Association, 2010, 304, 2611.	7.4	174
39	Molecular Biology of Pheochromocytomas and Paragangliomas. , 2010, , 297-305.		2
40	A germline mutation of the KIF1BÎ² gene on 1p36 in a family with neural and nonneural tumors. Human Genetics, 2008, 124, 279-285.	3.8	139
41	Transcription Association of VHL and SDH Mutations Link Hypoxia and Oxidoreductase Signals in Pheochromocytomas. Annals of the New York Academy of Sciences, 2006, 1073, 208-220.	3.8	42
42	Evolving concepts in pheochromocytoma and paraganglioma. Current Opinion in Oncology, 2006, 18, 1-8.	2.4	64
43	Clinical Presentation and Penetrance of Pheochromocytoma/Paraganglioma Syndromes. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 827-836.	3.6	560
44	A HIF1Î± Regulatory Loop Links Hypoxia and Mitochondrial Signals in Pheochromocytomas. PLoS Genetics, 2005, 1, e8.	3.5	394
45	Novel Pheochromocytoma Susceptibility Loci Identified by Integrative Genomics. Cancer Research, 2005, 65, 9651-9658.	0.9	88
46	RASSF1A promoter region CpG island hypermethylation in phaeochromocytomas and neuroblastoma tumours. Oncogene, 2001, 20, 7573-7577.	5.9	127
47	Allelic imbalance, including deletion ofPTEN/MMAC1, at the Cowden disease locus on 10q22-23, in hamartomas from patients with cowden syndrome and germlinePTEN mutation. , 1998, 21, 61-69.		85
48	Germline Dinucleotide Mutation in Codon 883 of theRETProto-Oncogene in Multiple Endocrine Neoplasia Type 2B Without Codon 918 Mutation. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 3902-3904.	3.6	216
49	Germline mutations of the PTEN gene in Cowden disease, an inherited breast and thyroid cancer syndrome. Nature Genetics, 1997, 16, 64-67.	21.4	1,902
50	Germline mutations in PTEN are present in Bannayan-Zonana syndrome. Nature Genetics, 1997, 16, 333-334.	21.4	622
51	Germline Dinucleotide Mutation in Codon 883 of the RETProto-Oncogene in Multiple Endocrine Neoplasia Type 2B Without Codon 918 Mutation. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 3902-3904.	3.6	73