

Preventive medicine of von Hippelâ€“Lindau disease-associated  
tumors

Endocrine-Related Cancer

25, 783-793

DOI: 10.1530/erc-18-0100

Citation Report

#	ARTICLE	IF	CITATIONS
1	Nutrition and neuroendocrine tumors: An update of the literature. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 159-167.	5.7	38
2	Curative and palliative surgery in patients with neuroendocrine tumors of the gastro-entero-pancreatic (GEP) tract. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 169-178.	5.7	12
3	New Insights Into Pheochromocytoma Surveillance of Young Patients With VHL Missense Mutations. Journal of the Endocrine Society, 2019, 3, 1682-1692.	0.2	15
4	Prognostic and predictive factors on overall survival and surgical outcomes in pancreatic neuroendocrine tumors: recent advances and controversies. Expert Review of Anticancer Therapy, 2019, 19, 1029-1050.	2.4	53
5	Concomitant pancreatic neuroendocrine tumors in hereditary tumor syndromes: who, when and how to operate?. Journal of Pancreatology, 2019, 2, 48-53.	0.9	3
6	von Hippel-Lindau Disease: an Update. Current Genetic Medicine Reports, 2019, 7, 227-235.	1.9	8
7	Neuroendocrine Neoplasms of the Small Bowel and Pancreas. Neuroendocrinology, 2020, 110, 444-476.	2.5	70
8	The North American Neuroendocrine Tumor Society Consensus Paper on the Surgical Management of Pancreatic Neuroendocrine Tumors. Pancreas, 2020, 49, 1-33.	1.1	226
9	Genetic and epigenetic alterations in pancreatic neuroendocrine tumors. Journal of Gastrointestinal Oncology, 2020, 11, 567-577.	1.4	14
10	Inherited syndromes involving pancreatic neuroendocrine tumors. Journal of Gastrointestinal Oncology, 2020, 11, 559-566.	1.4	19
11	A Case of Von Hippel-Lindau Disease with Bilateral Pheochromocytoma and Ectopic Hypersecretion of Intact Parathyroid Hormone in an Adolescent Girl. Case Reports in Endocrinology, 2020, 2020, 1-5.	0.4	1
12	Hemangioblastoma and von Hippel-Lindau disease: genetic background, spectrum of disease, and neurosurgical treatment. Child's Nervous System, 2020, 36, 2537-2552.	1.1	23
13	&lt;p&gt;Von Hippel-Lindau Disease: Current Challenges and Future Prospects&lt;/p&gt;. OncoTargets and Therapy, 2020, Volume 13, 5669-5690.	2.0	66
14	Maternal and fetal outcomes in phaeochromocytoma and pregnancy: a multicentre retrospective cohort study and systematic review of literature. Lancet Diabetes and Endocrinology,the, 2021, 9, 13-21.	11.4	37
15	Prenatal Diagnosis and Preimplantation Genetic Diagnosis. , 2021, , 769-800.		0
16	Neuroendocrine Neoplasms with Peculiar Biology and Features: MEN1, MEN2A, MEN2B, MEN4, VHL, NF1. , 2021, , 233-267.		0
17	Pheochromocytoma/Paraganglioma, Medullary Thyroid Carcinoma, and Hereditary Endocrine Neoplasia Syndromes. , 2021, , 491-527.		1
18	Diagnostic and management strategies for pNETs in Von Hippel-Lindau: a systematic review. Endocrine-Related Cancer, 2021, 28, 151-160.	3.1	5

#	ARTICLE	IF	CITATIONS
19	A Direct Comparison of Patients With Hereditary and Sporadic Pancreatic Neuroendocrine Tumors: Evaluation of Clinical Course, Prognostic Factors and Genotypeâ€“Phenotype Correlations. <i>Frontiers in Endocrinology</i> , 2021, 12, 681013.	3.5	7
20	Spectral CT in clinical routine imaging of neuroendocrine neoplasms. <i>Clinical Radiology</i> , 2021, 76, 348-357.	1.1	4
21	Clinical presentation, genotypeâ€“phenotype correlations, and outcome of pancreatic neuroendocrine tumors in Von Hippelâ€“Lindau syndrome. <i>Endocrine</i> , 2021, 74, 180-187.	2.3	5
22	Phakomatoses and Endocrine Gland Tumors: Noteworthy and (Not so) Rare Associations. <i>Frontiers in Endocrinology</i> , 2021, 12, 678869.	3.5	3
23	Hereditary syndromes associated with neuroendocrine tumors. <i>Current Opinion in Endocrine and Metabolic Research</i> , 2021, 18, 230-235.	1.4	0
24	Pancreas in Hereditary Syndromes: Cross-sectional Imaging Spectrum. <i>Radiographics</i> , 2021, 41, 200164.	3.3	2
25	Unsuspected Von Hippel-Lindau syndrome in acute-onset resistant hypertension. <i>BMJ Case Reports</i> , 2018, 2018, bcr-2018-225162.	0.5	1
26	VON HIPPEL-LINDAU SYNDROME: ASPECTS OF TREATMENT AND MANAGEMENT. CASE REPORT AND LITERATURE REVIEW. <i>Russian Archives of Internal Medicine</i> , 2019, 9, 165-171.	0.2	1
27	Clinical diagnosis, treatment and screening of the VHL gene in three von Hippelâ€“Lindau disease pedigrees. <i>Experimental and Therapeutic Medicine</i> , 2020, 20, 1237-1244.	1.8	4
28	Management recommendations for pancreatic manifestations of von Hippelâ€“Lindau disease. <i>Cancer</i> , 2022, 128, 435-446.	4.1	14
29	Pancreatic neuroendocrine neoplasms: Updates on genomic changes in inherited tumour syndromes and sporadic tumours based on WHO classification. <i>Critical Reviews in Oncology/Hematology</i> , 2022, 172, 103648.	4.4	8
30	Identification of a VHL gene mutation in atypical Von Hippel-Lindau syndrome: genotypeâ€“phenotype correlation and gene therapy perspective. <i>Cancer Cell International</i> , 2021, 21, 685.	4.1	0
31	Is There a Role for Biomarkers in Surveillance of Pancreatic Neuroendocrine Neoplasms in Von Hippel-Lindau Disease?. <i>Journal of the Endocrine Society</i> , 2022, 6, bvab191.	0.2	0
32	Metastatic Pheochromocytoma Diagnosed with 131I-MIBG SPECT/CT Imaging in a Patient with Pathogenic VHL Mutation. <i>World Journal of Nuclear Medicine</i> , 2022, 21, 073-075.	0.5	0
33	Large scale genotypeâ€“and phenotypeâ€“driven machine learning in Von Hippelâ€“Lindau disease. <i>Human Mutation</i> , 2022, 43, 1268-1285.	2.5	6
34	Neuroendocrine neoplasms in the context of inherited tumor syndromes: a reappraisal focused on targeted therapies. <i>Journal of Endocrinological Investigation</i> , 2023, 46, 213-234.	3.3	13
35	Genetics of Pancreatic Neuroendocrine Tumors. <i>Hematology/Oncology Clinics of North America</i> , 2022, 36, 1033-1051.	2.2	3
36	Multidisciplinary management of patients diagnosed with von Hippel-Lindau disease: A practical review of the literature for clinicians. <i>Asian Journal of Urology</i> , 2022, 9, 430-442.	1.2	2

#	ARTICLE	IF	CITATIONS
37	PANCREATODUODENECTOMY IN PATIENT WITH VON HIPPEL-LINDAU DISEASE: A LITERATURE REVIEW. Arquivos Brasileiros De Cirurgia Digestiva: ABCD = Brazilian Archives of Digestive Surgery, 0, 35, .	0.5	1
38	Synchronous or metachronous presentation of pancreatic neuroendocrine tumor versus secondary lesion to pancreas in patients affected by renal cell carcinoma. Systematic review. Seminars in Oncology, 2022, 49, 476-481.	2.2	0
39	Unique Characteristics of Patients with Von Hippel-Lindau Disease Defined by Various Diagnostic Criteria. Cancers, 2023, 15, 1657.	3.7	1
40	Non-Interventional Management of Advanced Pancreatic Neuroendocrine Neoplasms in Patients with von Hippel-Lindau Disease. Cancers, 2023, 15, 1739.	3.7	2
41	Guidelines for surveillance of patients with von Hippel-Lindau disease: Consensus statement of the International VHL Surveillance Guidelines Consortium and VHL Alliance. Cancer, 2023, 129, 2927-2940.	4.1	2