Ronald De Krijger

List of Publications by Year in descending order

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687220 940416 1,957 16 13 16 citations h-index g-index papers 16 16 16 2430 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Integrated genomic characterization of adrenocortical carcinoma. Nature Genetics, 2014, 46, 607-612.	9.4	560
2	European Society of Endocrinology Clinical Practice Guidelines on the management of adrenocortical carcinoma in adults, in collaboration with the European Network for the Study of Adrenal Tumors. European Journal of Endocrinology, 2018, 179, G1-G46.	1.9	559
3	Major Prognostic Role of Ki67 in Localized Adrenocortical Carcinoma After Complete Resection. Journal of Clinical Endocrinology and Metabolism, 2015, 100, 841-849.	1.8	274
4	Observer Variation in the Application of the Pheochromocytoma of the Adrenal Gland Scaled Score. American Journal of Surgical Pathology, 2009, 33, 599-608.	2.1	152
5	Reactivity of Germ Cell Maturation Stage-Specific Markers in Spermatocytic Seminoma: Diagnostic and Etiological Implications. Laboratory Investigation, 2001, 81, 919-928.	1.7	106
6	Value of Molecular Classification for Prognostic Assessment of Adrenocortical Carcinoma. JAMA Oncology, 2019, 5, 1440.	3.4	57
7	The window period of NEUROGENIN3 during human gestation. Islets, 2014, 6, e954436.	0.9	47
8	DNA methylation is an independent prognostic marker of survival in adrenocortical cancer. Journal of Clinical Endocrinology and Metabolism, 2016, 102, jc.2016-3205.	1.8	44
9	Premature differentiation of vascular smooth muscle cells in human congenital diaphragmatic hernia. Experimental and Molecular Pathology, 2013, 94, 195-202.	0.9	43
10	Genotype and Tumor Locus Determine Expression Profile of Pseudohypoxic Pheochromocytomas and Paragangliomas. Neoplasia, 2013, 15, 435-IN22.	2.3	33
11	Altered Phenotype of \hat{l}^2 -Cells and Other Pancreatic Cell Lineages in Patients With Diffuse Congenital Hyperinsulinism in Infancy Caused by Mutations in the ATP-Sensitive K-Channel. Diabetes, 2015, 64, 3182-3188.	0.3	20
12	Usage of TCRAV and TCRBV gene families in human fetal and adult TCR rearrangements. Immunogenetics, 1994, 39, 343-350.	1.2	18
13	Hypoxia-Inducible Factor 2α Mutation-Related Paragangliomas Classify as Discrete Pseudohypoxic Subcluster. Neoplasia, 2016, 18, 567-576.	2.3	16
14	Vascular Pattern Analysis for the Prediction of Clinical Behaviour in Pheochromocytomas and Paragangliomas. PLoS ONE, 2015, 10, e0121361.	1.1	14
15	Analysis of early fetal T-cell receptor δchain in humans. Immunogenetics, 1990, 32, 331-336.	1.2	9
16	Hypoxia inducible factor 2α (HIF2α/EPAS1) is associated with development of pulmonary hypertension in severe congenital diaphragmatic hernia patients. Pulmonary Circulation, 2018, 8, 1-4.	0.8	5