Guillaume Assié

List of Publications by Year in descending order

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CUULAUME ASSIÃO

#	Article	IF	CITATIONS
1	Integrated genomic characterization of adrenocortical carcinoma. Nature Genetics, 2014, 46, 607-612.	21.4	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.	3.7	559
3	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	16.8	482
4	<i>ARMC5</i> Mutations in Macronodular Adrenal Hyperplasia with Cushing's Syndrome. New England Journal of Medicine, 2013, 369, 2105-2114.	27.0	319
5	Corticotroph Tumor Progression after Adrenalectomy in Cushing's Disease: A Reappraisal of Nelson's Syndrome. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 172-179.	3.6	221
6	Pangenomic Classification of Pituitary Neuroendocrine Tumors. Cancer Cell, 2020, 37, 123-134.e5.	16.8	186
7	Aberrant DNA hypermethylation of SDHC: a novel mechanism of tumor development in Carney triad. Endocrine-Related Cancer, 2014, 21, 567-577.	3.1	161
8	Genetic Landscape of Sporadic Unilateral Adrenocortical Adenomas Without PRKACA p.Leu206Arg Mutation. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 3526-3538.	3.6	65
9	The pathophysiology, diagnosis and prognosis of adrenocortical tumors revisited by transcriptome analyses. Trends in Endocrinology and Metabolism, 2010, 21, 325-334.	7.1	42
10	The 'omics' of adrenocortical tumours for personalized medicine. Nature Reviews Endocrinology, 2014, 10, 215-228.	9.6	41
11	Somatic USP8 mutations are frequent events in corticotroph tumor progression causing Nelson's tumor. European Journal of Endocrinology, 2018, 178, 57-63.	3.7	37
12	Corticotroph tumor progression after bilateral adrenalectomy (Nelson's syndrome): systematic review and expert consensus recommendations. European Journal of Endocrinology, 2021, 184, P1-P16.	3.7	32
13	KDM1A inactivation causes hereditary food-dependent Cushing syndrome. Genetics in Medicine, 2022, 24, 374-383.	2.4	27
14	Biological and radiological exploration and management of non-functioning pituitary adenoma. Annales D'Endocrinologie, 2015, 76, 201-209.	1.4	25
15	Serum RARRES2 Is a Prognostic Marker in Patients With Adrenocortical Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 3345-3352.	3.6	21
16	Glucocorticoid Excess in Patients with Pheochromocytoma Compared with Paraganglioma and Other Forms of Hypertension. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e3374-e3383.	3.6	17
17	Genomics of benign adrenocortical tumors. Journal of Steroid Biochemistry and Molecular Biology, 2019, 193, 105414.	2.5	15
18	Calling Chromosome Alterations, DNA Methylation Statuses, and Mutations in Tumors by Simple Targeted Next-Generation Sequencing. Journal of Molecular Diagnostics, 2017, 19, 776-787.	2.8	7

#	Article	IF	CITATIONS
19	Somatotroph Tumors and the Epigenetic Status of the GNAS Locus. International Journal of Molecular Sciences, 2021, 22, 7570.	4.1	6
20	SUN-LB97 Targeted Metabolomics as a Screening Tool in the Diagnosis of Endocrine Hypertension. Journal of the Endocrine Society, 2020, 4, .	0.2	0