Maria Cbv Fragoso

List of Publications by Year
in descending order

72	2,991	185998 28	174990 52
papers	citations	h-index	g-index

77	77	77	3266
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. Cancer Cell, 2016, 29, 723-736.	7.7	482
2	An Inherited Mutation Outside the Highly Conserved DNA-Binding Domain of the p53 Tumor Suppressor Protein in Children and Adults with Sporadic Adrenocortical Tumors. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 4970-4973.	1.8	183
3	Cushingâ€™s Syndrome Secondary to Adrenocorticotropin-Independent Macronodular Adrenocortical Hyperplasia due to Activating Mutations ofGNAS1Gene. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 2147-2151.	1.8	174
4	Progression to Adrenocortical Tumorigenesis in Mice and Humans through Insulin-Like Growth Factor 2 and Î²-Catenin. American Journal of Pathology, 2012, 181, 1017-1033.	1.9	154
5	Expression of Insulin-Like Growth Factor-II and Its Receptor in Pediatric and Adult Adrenocortical Tumors. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 3524-3531.	1.8	149
6	The desmopressin stimulation test in the differential diagnosis of Cushing's syndrome. Clinical Endocrinology, 1993, 38, 463-472.	1.2	137
7	ARMC5Mutations Are a Frequent Cause of Primary Macronodular Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E1501-E1509.	1.8	120
8	Activating Mutation of the Stimulatory G Protein (gsp) as a Putative Cause of Ovarian and Testicular Human Stromal Leydig Cell Tumors. Journal of Clinical Endocrinology and Metabolism, 1998, 83, 2074-2078.	1.8	93
9	Combined expression of BUB1B, DLGAP5, and PINK1 as predictors of poor outcome in adrenocortical tumors: validation in a Brazilian cohort of adult and pediatric patients. European Journal of Endocrinology, 2012, 166, 61-67.	1.9	81
10	Steroidogenic Factor 1 Overexpression and Gene Amplification Are More Frequent in Adrenocortical Tumors from Children than from Adults. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1458-1462.	1.8	66
11	Primary bilateral macronodular adrenal hyperplasia. Current Opinion in Endocrinology, Diabetes and Obesity, 2014, 21, 177-184.	1.2	61
12	The role of desmopressin in bilateral and simultaneous inferior petrosal sinus sampling for differential diagnosis of ACTH-dependent Cushing's syndrome. Clinical Endocrinology, 2006, 66, 061120012318003-???.	1.2	55
13	ARMC5 mutations in a large French-Canadian family with cortisol-secreting Î²-adrenergic/vasopressin responsive bilateral macronodular adrenal hyperplasia. European Journal of Endocrinology, 2016, 174, 85-96.	1.9	55
14	Treatment of gonadotropin dependent precocious puberty due to hypothalamic hamartoma with gonadotropin releasing hormone agonist depot. Archives of Disease in Childhood, 1999, 80, 231-234.	1.0	45
15	High Penetrance of Pheochromocytoma Associated with the Novel C634Y/Y791F Double Germline Mutation in theRETProtooncogene. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1318-1327.	1.8	43
16	MANAGEMENT OF ENDOCRINE DISEASE: Management of pregnant patients with Cushing's syndrome. European Journal of Endocrinology, 2015, 173, R85-R91.	1.9	43
17	Genetics of primary macronodular adrenal hyperplasia. Journal of Endocrinology, 2015, 224, R31-R43.	1.2	41
18	No evidence of somatic activating mutations on gonadotropin receptor genes in sex cord stromal tumors. Fertility and Sterility, 2000, 74, 992-995.	0.5	34

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19	Deletion Mapping of Chromosome 17 in Benign and Malignant Adrenocortical Tumors Associated with the Arg337His Mutation of the p53 Tumor Suppressor Protein. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 2976-2981.	1.8	34
20	Metabolic reprogramming: a new relevant pathway in adult adrenocortical tumors. Oncotarget, 2015, 6, 44403-44421.	0.8	34
21	Isolated familial somatotropinoma: 11q13-loh and gene/protein expression analysis suggests a possible involvement of aip also in non-pituitary tumorigenesis. Clinics, 2010, 65, 407-415.	0.6	33
22	Cortisol and adrenocorticotropin response to desmopressin in women with Cushing's disease compared with depressive illness. Journal of Clinical Endocrinology and Metabolism, 1996, 81, 2233-2237.	1.8	32
23	¹⁸F-FDG-PET/CT Imaging of ACTH-Independent Macronodular Adrenocortical Hyperplasia (AIMAH) Demonstrating Increased¹⁸F-FDG Uptake. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 3300-3301.	1.8	31
24	The Use of Three-dimensional Printers for Partial Adrenalectomy: Estimating the Resection Limits. Urology, 2016, 90, 217-221.	0.5	31
25	Complete Resolution of Hypercortisolism with Sorafenib in a Patient with Advanced Medullary Thyroid Carcinoma and Ectopic ACTH (Adrenocorticotropic Hormone) Syndrome. Thyroid, 2014, 24, 1062-1066.	2.4	29
26	Cushingâ€™s disease due to somatic USP8 mutations: a systematic review and meta-analysis. Pituitary, 2019, 22, 435-442.	1.6	29
27	POD-1 binding to the E-box sequence inhibits SF-1 and StAR expression in human adrenocortical tumor cells. Molecular and Cellular Endocrinology, 2013, 371, 140-147.	1.6	28
28	Sonic Hedgehog Signaling Is Active in Human Adrenal Cortex Development and Deregulated in Adrenocortical Tumors. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E1209-E1216.	1.8	27
29	Association between the p27 rs2066827 variant and tumor multiplicity in patients harboring MEN1 germline mutations. European Journal of Endocrinology, 2014, 171, 335-342.	1.9	25
30	Expression of <scp>LIN</scp>28 and its regulatory micro<scp>RNA</scp>s in adult adrenocortical cancer. Clinical Endocrinology, 2015, 82, 481-488.	1.2	25
31	Transcriptome Analysis Showed a Differential Signature between Invasive and Non-invasive Corticotrophinomas. Frontiers in Endocrinology, 2017, 8, 55.	1.5	24
32	Radiographic Characteristics of Adrenal Masses Preceding the Diagnosis of Adrenocortical Cancer. Hormones and Cancer, 2015, 6, 176-181.	4.9	23
33	p27 variant and corticotropinoma susceptibility: a genetic and in vitro study. Endocrine-Related Cancer, 2014, 21, 395-404.	1.6	20
34	The role of fibroblast growth factor receptor 4 overexpression and gene amplification as prognostic markers in pediatric and adult adrenocortical tumors. Endocrine-Related Cancer, 2012, 19, L11-L13.	1.6	19
35	Altered expression of noncanonical Wnt pathway genes in paediatric and adult adrenocortical tumours. Clinical Endocrinology, 2014, 81, 503-510.	1.2	19
36	Pregnancy in Women Previously Treated for an Adrenocortical Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2015, 100, 4604-4611.	1.8	19

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37	Differential Expression of Stem Cell Markers in Human Adamantinomatous Craniopharyngioma and Pituitary Adenoma. Neuroendocrinology, 2017, 104, 183-193.	1.2	19
38	Mutation analysis of the follicle-stimulating hormone receptor gene in girls with gonadotropin-independent precocious puberty resulting from autonomous cystic ovaries. Fertility and Sterility, 2000, 73, 280-283.	0.5	18
39	Possible role of a radiation-induced p53 mutation in a Nelsonâ€™s syndrome patient with a fatal outcome. Pituitary, 2011, 14, 400-404.	1.6	18
40	Low DICER1 expression is associated with poor clinical outcome in adrenocortical carcinoma. Oncotarget, 2015, 6, 22724-22733.	0.8	18
41	Influence of the Fibroblast Growth Factor Receptor 4 Expression and the G388R Functional Polymorphism on Cushingâ€™s Disease Outcome. Journal of Clinical Endocrinology and Metabolism, 2010, 95, E271-E279.	1.8	17
42	Choroidal and Retinal Abnormalities by Optical Coherence Tomography in Endogenous Cushingâ€™s Syndrome. Frontiers in Endocrinology, 2016, 7, 154.	1.5	16
43	Negative correlation between tumour size and cortisol/ACTH ratios in patients with Cushingâ€™s disease harbouring microadenomas or macroadenomas. Journal of Endocrinological Investigation, 2016, 39, 1401-1409.	1.8	16
44	Guidelines for the management of neuroendocrine tumours by the Brazilian gastrointestinal tumour group. Ecancermedicalscience, 2017, 11, 716.	0.6	16
45	Recommendations of the Neuroendocrinology Department of the Brazilian Society of Endocrinology and Metabolism for the diagnosis of Cushingâ€™s disease in Brazil. Archives of Endocrinology and Metabolism, 2016, 60, 267-286.	0.3	14
46	Genetics of primary macronodular adrenal hyperplasia. Presse Medicale, 2018, 47, e139-e149.	0.8	14
47	A New Insight into the Surgical Treatment of Primary Macronodular Adrenal Hyperplasia. Journal of the Endocrine Society, 2020, 4, bvaa083.	0.1	14
48	An Inhibin B and Estrogen-Secreting Adrenocortical Carcinoma Leading to Selective FSH Suppression. Hormone Research in Paediatrics, 2007, 67, 7-11.	0.8	11
49	Amplification of theInsulin-Like Growth Factor 1 ReceptorGene Is a Rare Event in Adrenocortical Adenocarcinomas: Searching for Potential Mechanisms of Overexpression. BioMed Research International, 2014, 2014, 1-7.	0.9	11
50	An Overview of the Heterogeneous Causes of Cushing Syndrome Resulting From Primary Macronodular Adrenal Hyperplasia (PMAH). Journal of the Endocrine Society, 2022, 6, bvac041.	0.1	11
51	POD-1/TCF21Reduces SHP Expression, AffectingLRH-1Regulation and Cell Cycle Balance in Adrenocortical and Hepatocarcinoma Tumor Cells. BioMed Research International, 2015, 2015, 1-9.	0.9	10
52	Presentation and surgery outcomes in elderly with pheocromocytoma: a comparative analysis with young patients. International Braz J Urol: Official Journal of the Brazilian Society of Urology, 2016, 42, 671-677.	0.7	10
53	Low Protein Expression of both ATRX and ZNRF3 as Novel Negative Prognostic Markers of Adult Adrenocortical Carcinoma. International Journal of Molecular Sciences, 2021, 22, 1238.	1.8	10
54	DAX1 Overexpression in Pediatric Adrenocortical Tumors: A Synergic Role with SF1 in Tumorigenesis. Hormone and Metabolic Research, 2015, 47, 656-661.	0.7	9

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55	High 18F-FDG uptake in PMAH correlated with normal expression of Glut1, HK1, HK2, and HK3. Acta Radiologica, 2016, 57, 370-377.	0.5	9
56	TCF21/POD-1, a Transcritional Regulator of SF-1/NR5A1, as a Potential Prognosis Marker in Adult and Pediatric Adrenocortical Tumors. Frontiers in Endocrinology, 2018, 9, 38.	1.5	9
57	Stathmin 1 is highly expressed and associated with survival outcome in malignant adrenocortical tumours. Investigational New Drugs, 2020, 38, 899-908.	1.2	9
58	PROP1 overexpression in corticotrophinomas: evidence for the role of PROP1 in the maintenance of cells committed to corticotrophic differentiation. Clinics, 2013, 68, 887-891.	0.6	9
59	The Role of gsp Mutations on the Development of Adrenocortical Tumors and Adrenal Hyperplasia. Frontiers in Endocrinology, 2016, 7, 104.	1.5	8
60	GLUT1 expression in pediatric adrenocortical tumors: a promising candidate to predict clinical behavior. Oncotarget, 2017, 8, 63835-63845.	0.8	8
61	Allelic Variants of ARMC5 in Patients With Adrenal Incidentalomas and in Patients With Cushing's Syndrome Associated With Bilateral Adrenal Nodules. Frontiers in Endocrinology, 2020, 11, 36.	1.5	7
62	A missense TCF1 mutation in a patient with mody-3 and liver adenomatosis. Clinics, 2010, 65, 1059-1060.	0.6	7
63	Long-term Results after CT-Guided Percutaneous Ethanol Ablation for the Treatment of Hyperfunctioning Adrenal Disorders. Clinics, 2016, 71, 600-605.	0.6	6
64	Genotype analysis of the human endostatin variant p.D104N in benign and malignant adrenocortical tumors. Clinics, 2012, 67, 95-98.	0.6	6
65	Expression profiles of the glucose-dependent insulinotropic peptide receptor and LHCGR in sporadic adrenocortical tumors. Journal of Endocrinology, 2009, 200, 167-175.	1.2	5
66	Analysis of glucose-dependent insulinotropic peptide receptor (GIPR) and luteinizing hormone receptor (LHCGR) expression in human adrenocortical hyperplasia. Arquivos Brasileiros De Endocrinologia E Metabologia, 2009, 53, 326-331.	1.3	5
67	High Prevalence of Alterations in DNA Mismatch Repair Genes of Lynch Syndrome in Pediatric Patients with Adrenocortical Tumors Carrying a Germline Mutation on TP53. Cancers, 2020, 12, 621.	1.7	4
68	Assessing the emerging oncogene protein kinase C epsilon as a candidate gene in families with Carney complexâ€2. Clinical Endocrinology, 2012, 76, 147-148.	1.2	2
69	Adrenocortical Tumors and gsp Mutations. , 2019, , 266-270.		0
70	Molecular and cellular regulation of primary macronodular adrenal hyperplasia. Current Opinion in Endocrine and Metabolic Research, 2019, 8, 112-121.	0.6	0
71	ACTH-Independent Cushingâ€™s Syndrome: Adrenocortical Tumors. , 2010, , 189-208.		0
72	Abstract 3464: Prognostic value of DICER1 expression in adrenocortical cancer patients. , 2015, , .		0