

# Maria Candida Barisson Villares Fragoso

## List of Publications by Year in descending order

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123  
papers

3,577  
citations

147801

31  
h-index

155660

55  
g-index

128  
all docs

128  
docs citations

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times ranked

3887  
citing authors

#	ARTICLE	IF	CITATIONS
1	Primary Adrenal Insufficiency Due to Bilateral Adrenal Infarction in COVID-19. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e394-e400.	3.6	24
2	Adrenal cysts of lymphatic origin: A clinical and pathological study of six cases and systematic literature review. <i>Annals of Diagnostic Pathology</i> , 2022, 57, 151888.	1.3	0
3	Response Letter to the Editor From de Ponthaud et al: "Cytoreductive Surgery of the Primary Tumor in Metastatic Adrenocortical Carcinoma: Impact on Patients' Survival". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, , .	3.6	0
4	An Overview of the Heterogeneous Causes of Cushing Syndrome Resulting From Primary Macronodular Adrenal Hyperplasia (PMAH). <i>Journal of the Endocrine Society</i> , 2022, 6, bvac041.	0.2	11
5	Low Protein Expression of both ATRX and ZNRF3 as Novel Negative Prognostic Markers of Adult Adrenocortical Carcinoma. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1238.	4.1	10
6	SDHB large deletions are associated with absence of MIBG uptake in metastatic lesions of malignant paragangliomas. <i>Endocrine</i> , 2021, 72, 586-590.	2.3	4
7	Phosphodiesterase 2A and 3B variants are associated with primary aldosteronism. <i>Endocrine-Related Cancer</i> , 2021, 28, 1-13.	3.1	17
8	Genetic and clinical aspects of paediatric pheochromocytomas and paragangliomas. <i>Clinical Endocrinology</i> , 2021, 95, 117-124.	2.4	10
9	Low Protein Expression of <i>ATRX</i> and <i>ZNRF3</i> as a Novel Prognostic Marker of Adult Adrenocortical Carcinoma. <i>Journal of the Endocrine Society</i> , 2021, 5, A87-A88.	0.2	0
10	Resolution of Cyclicity After Pasireotide LAR in a Patient With Cushing Disease. <i>AACE Clinical Case Reports</i> , 2021, 7, 277-281.	1.1	1
11	Role of the Mevalonate Pathway in Adrenocortical Tumorigenesis. <i>Hormone and Metabolic Research</i> , 2021, 53, 124-131.	1.5	1
12	Stathmin 1 is highly expressed and associated with survival outcome in malignant adrenocortical tumours. <i>Investigational New Drugs</i> , 2020, 38, 899-908.	2.6	9
13	Radiotherapy-induced malignancies in breast cancer patients with TP53 pathogenic germline variants (Li-Fraumeni syndrome). <i>Familial Cancer</i> , 2020, 19, 47-53.	1.9	16
14	A New Insight into the Surgical Treatment of Primary Macronodular Adrenal Hyperplasia. <i>Journal of the Endocrine Society</i> , 2020, 4, bvaa083.	0.2	14
15	High accuracy of bilateral and simultaneous petrosal sinus sampling with desmopressin for the differential diagnosis of pediatric ACTH-dependent Cushing's syndrome. <i>Pituitary</i> , 2020, 23, 507-514.	2.9	10
16	Pediatric adrenocortical tumor " review and management update. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2020, 27, 177-186.	2.3	16
17	High Prevalence of Alterations in DNA Mismatch Repair Genes of Lynch Syndrome in Pediatric Patients with Adrenocortical Tumors Carrying a Germline Mutation on TP53. <i>Cancers</i> , 2020, 12, 621.	3.7	4
18	XAF1 as a modifier of p53 function and cancer susceptibility. <i>Science Advances</i> , 2020, 6, eaba3231.	10.3	37

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19	Crossed-Probes Cryoablation for the Treatment of a Sclerotic Vertebral Metastasis Abutting the Spinal Canal. <i>Journal of Vascular and Interventional Radiology</i> , 2020, 31, 284-285.	0.5	1
20	Allelic Variants of ARMC5 in Patients With Adrenal Incidentalomas and in Patients With Cushing's Syndrome Associated With Bilateral Adrenal Nodules. <i>Frontiers in Endocrinology</i> , 2020, 11, 36.	3.5	7
21	Sterol O-Acyl Transferase 1 as a Prognostic Marker of Adrenocortical Carcinoma. <i>Cancers</i> , 2020, 12, 247.	3.7	22
22	Internal validation and decision curve analysis of a preoperative nomogram predicting a postoperative complication in pheochromocytoma surgery: An international study. <i>International Journal of Urology</i> , 2020, 27, 463-468.	1.0	7
23	Expression of SOAT1 in Adrenocortical Carcinoma and Response to Mitotane Monotherapy: An ENSAT Multicenter Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, 2642-2653.	3.6	18
24	Cullin 3 targets the tumor suppressor gene ARMC5 for ubiquitination and degradation. <i>Endocrine-Related Cancer</i> , 2020, 27, 221-230.	3.1	15
25	ARMC5 mutations are associated with high levels of proliferating cell nuclear antigen and the presence of the serotonin receptor 5HT4R in PMAH nodules. <i>Archives of Endocrinology and Metabolism</i> , 2020, 64, 390-401.	0.6	3
26	SAT-560 Usefulness of Contralateral Suppression in Adrenal Venous Sampling to Define Lateralization in Primary Aldosteronism. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
27	SUN-178 Clinical and Anatomopathological Characteristics of Two Atypical Aldosterone-Producing Adenomas. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
28	MON-250 Late Diagnosis of ACTH-secreting Pulmonary Neuroendocrine Tumor by Repeated 68Ga Dotatate Pet/ct: Influence of Tumor Size in Abnormal Uptake?. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
29	MON-312 The Effects of Cabergoline in Pre-Surgical and Recurrence Periods of Cushing's Disease Patients. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
30	MON-206 Diagnosis of Non-Functional Masses in Adrenal Gland Topography - Experience of a Tertiary Health Center. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
31	SUN-172 A False Positive Result in Newborn Screening for Congenital Adrenal Hyperplasia (CAH) in a Girl with Beckwith Wiedemann Syndrome. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
32	Adrenocortical Tumors and gsp Mutations. , 2019, , 266-270.		0
33	Predictors of complication after adrenalectomy. <i>International Braz J Urol: Official Journal of the Brazilian Society of Urology</i> , 2019, 45, 514-522.	1.5	7
34	Cushing's disease due to somatic USP8 mutations: a systematic review and meta-analysis. <i>Pituitary</i> , 2019, 22, 435-442.	2.9	29
35	Molecular and cellular regulation of primary macronodular adrenal hyperplasia. <i>Current Opinion in Endocrine and Metabolic Research</i> , 2019, 8, 112-121.	1.4	0
36	KCNJ5 Somatic Mutation Is a Predictor of Hypertension Remission After Adrenalectomy for Unilateral Primary Aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 4695-4702.	3.6	42

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37	Prognostic Relevance of Steroid Sulfation in Adrenocortical Carcinoma Revealed by Molecular Phenotyping Using High-Resolution Mass Spectrometry Imaging. <i>Clinical Chemistry</i> , 2019, 65, 1276-1286.	3.2	19
38	New Insights Into Pheochromocytoma Surveillance of Young Patients With VHL Missense Mutations. <i>Journal of the Endocrine Society</i> , 2019, 3, 1682-1692.	0.2	15
39	Clinical spectrum of Li-Fraumeni syndrome/Li-Fraumeni-like syndrome in Brazilian individuals with the TP53 p.R3337H mutation. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 190, 250-255.	2.5	23
40	Filamin A and DRD2 expression in corticotrophinomas. <i>Pituitary</i> , 2019, 22, 163-169.	2.9	5
41	Targeted Assessment of GOS2 Methylation Identifies a Rapidly Recurrent, Routinely Fatal Molecular Subtype of Adrenocortical Carcinoma. <i>Clinical Cancer Research</i> , 2019, 25, 3276-3288.	7.0	51
42	SAT-064 Validation of Furosemide Upright Test in Primary Aldosteronism Diagnosis Using Direct Renin Assay. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.2	0
43	The role of ARMC5 in human cell cultures from nodules of primary macronodular adrenocortical hyperplasia (PMAH). <i>Molecular and Cellular Endocrinology</i> , 2018, 460, 36-46.	3.2	38
44	Primary malignant tumors of the adrenal glands. <i>Clinics</i> , 2018, 73, e756s.	1.5	27
45	Genetics of primary macronodular adrenal hyperplasia. <i>Presse Medicale</i> , 2018, 47, e139-e149.	1.9	14
46	A review of Cushing's disease treatment by the Department of Neuroendocrinology of the Brazilian Society of Endocrinology and Metabolism. <i>Archives of Endocrinology and Metabolism</i> , 2018, 62, 87-105.	0.6	3
47	TCF21/POD-1, a Transcriptional Regulator of SF-1/NR5A1, as a Potential Prognosis Marker in Adult and Pediatric Adrenocortical Tumors. <i>Frontiers in Endocrinology</i> , 2018, 9, 38.	3.5	9
48	Preventive medicine of von Hippel-Lindau disease-associated pancreatic neuroendocrine tumors. <i>Endocrine-Related Cancer</i> , 2018, 25, 783-793.	3.1	42
49	Somatic USP8 mutations are frequent events in corticotroph tumor progression causing Nelson's tumor. <i>European Journal of Endocrinology</i> , 2018, 178, 57-63.	3.7	37
50	Triple A Syndrome: Preliminary Response to the Antioxidant N-Acetylcysteine Treatment in a Child. <i>Hormone Research in Paediatrics</i> , 2017, 88, 167-171.	1.8	4
51	Differential Expression of Stem Cell Markers in Human Adamantinomatous Craniopharyngioma and Pituitary Adenoma. <i>Neuroendocrinology</i> , 2017, 104, 183-193.	2.5	19
52	New evidences on the regulation of SF-1 expression by POD1/TCF21 in adrenocortical tumor cells. <i>Clinics</i> , 2017, 72, 391-394.	1.5	6
53	Transcriptome Analysis Showed a Differential Signature between Invasive and Non-invasive Corticotrophinomas. <i>Frontiers in Endocrinology</i> , 2017, 8, 55.	3.5	24
54	Adjuvant radiotherapy for the primary treatment of adrenocortical carcinoma: are we offering the best?. <i>International Braz J Urol: Official Journal of the Brazilian Society of Urology</i> , 2017, 43, 841-848.	1.5	17

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55	Guidelines for the management of neuroendocrine tumours by the Brazilian gastrointestinal tumour group. <i>Ecancelmedicalsecience</i> , 2017, 11, 716.	1.1	16
56	GLUT1 expression in pediatric adrenocortical tumors: a promising candidate to predict clinical behavior. <i>Oncotarget</i> , 2017, 8, 63835-63845.	1.8	8
57	Presentation and surgery outcomes in elderly with pheochromocytoma: a comparative analysis with young patients. <i>International Braz J Urol: Official Journal of the Brazilian Society of Urology</i> , 2016, 42, 671-677.	1.5	10
58	Long-term Results after CT-Guided Percutaneous Ethanol Ablation for the Treatment of Hyperfunctioning Adrenal Disorders. <i>Clinics</i> , 2016, 71, 600-605.	1.5	6
59	The Role of gsp Mutations on the Development of Adrenocortical Tumors and Adrenal Hyperplasia. <i>Frontiers in Endocrinology</i> , 2016, 7, 104.	3.5	8
60	Choroidal and Retinal Abnormalities by Optical Coherence Tomography in Endogenous Cushing's Syndrome. <i>Frontiers in Endocrinology</i> , 2016, 7, 154.	3.5	16
61	Recommendations of the Neuroendocrinology Department of the Brazilian Society of Endocrinology and Metabolism for the diagnosis of Cushing's disease in Brazil. <i>Archives of Endocrinology and Metabolism</i> , 2016, 60, 267-286.	0.6	14
62	Negative correlation between tumour size and cortisol/ACTH ratios in patients with Cushing's disease harbouring microadenomas or macroadenomas. <i>Journal of Endocrinological Investigation</i> , 2016, 39, 1401-1409.	3.3	16
63	ARMC5 mutations in a large French-Canadian family with cortisol-secreting $\beta$ -adrenergic/vasopressin responsive bilateral macronodular adrenal hyperplasia. <i>European Journal of Endocrinology</i> , 2016, 174, 85-96.	3.7	55
64	Comprehensive Pan-Genomic Characterization of Adrenocortical Carcinoma. <i>Cancer Cell</i> , 2016, 29, 723-736.	16.8	482
65	The Use of Three-dimensional Printers for Partial Adrenalectomy: Estimating the Resection Limits. <i>Urology</i> , 2016, 90, 217-221.	1.0	31
66	High 18F-FDG uptake in PMAH correlated with normal expression of Glut1, HK1, HK2, and HK3. <i>Acta Radiologica</i> , 2016, 57, 370-377.	1.1	9
67	Fatal factitious Cushing syndrome (MÃ¼nchhausen's syndrome) in a patient with macroprolactinoma and silent corticotrophinoma: case report and literature review. <i>Clinical Diabetes and Endocrinology</i> , 2015, 1, 3.	2.7	5
68	POD-1/TCF21 Reduces SHP Expression, Affecting LRH-1 Regulation and Cell Cycle Balance in Adrenocortical and Hepatocarcinoma Tumor Cells. <i>BioMed Research International</i> , 2015, 2015, 1-9.	1.9	10
69	Low DICER1 expression is associated with poor clinical outcome in adrenocortical carcinoma. <i>Oncotarget</i> , 2015, 6, 22724-22733.	1.8	18
70	Pregnancy in Women Previously Treated for an Adrenocortical Carcinoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 4604-4611.	3.6	19
71	Expression of $\text{LIN}28$ and its regulatory microRNAs in adult adrenocortical cancer. <i>Clinical Endocrinology</i> , 2015, 82, 481-488.	2.4	25
72	MANAGEMENT OF ENDOCRINE DISEASE: Management of pregnant patients with Cushing's syndrome. <i>European Journal of Endocrinology</i> , 2015, 173, R85-R91.	3.7	43

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73	Genetics of primary macronodular adrenal hyperplasia. <i>Journal of Endocrinology</i> , 2015, 224, R31-R43.	2.6	41
74	Radiographic Characteristics of Adrenal Masses Preceding the Diagnosis of Adrenocortical Cancer. <i>Hormones and Cancer</i> , 2015, 6, 176-181.	4.9	23
75	DAX1 Overexpression in Pediatric Adrenocortical Tumors: A Synergic Role with SF1 in Tumorigenesis. <i>Hormone and Metabolic Research</i> , 2015, 47, 656-661.	1.5	9
76	Metabolic reprogramming: a new relevant pathway in adult adrenocortical tumors. <i>Oncotarget</i> , 2015, 6, 44403-44421.	1.8	34
77	Abstract 3464: Prognostic value of DICER1 expression in adrenocortical cancer patients. , 2015, , .		0
78	p27 variant and corticotropinoma susceptibility: a genetic and in vitro study. <i>Endocrine-Related Cancer</i> , 2014, 21, 395-404.	3.1	20
79	Association between the p27 rs2066827 variant and tumor multiplicity in patients harboring MEN1 germline mutations. <i>European Journal of Endocrinology</i> , 2014, 171, 335-342.	3.7	25
80	Amplification of the <i>Insulin-Like Growth Factor 1 Receptor</i> Gene Is a Rare Event in Adrenocortical Adenocarcinomas: Searching for Potential Mechanisms of Overexpression. <i>BioMed Research International</i> , 2014, 2014, 1-7.	1.9	11
81	Altered expression of noncanonical Wnt pathway genes in paediatric and adult adrenocortical tumours. <i>Clinical Endocrinology</i> , 2014, 81, 503-510.	2.4	19
82	Primary bilateral macronodular adrenal hyperplasia. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2014, 21, 177-184.	2.3	61
83	Complete Resolution of Hypercortisolism with Sorafenib in a Patient with Advanced Medullary Thyroid Carcinoma and Ectopic ACTH (Adrenocorticotrophic Hormone) Syndrome. <i>Thyroid</i> , 2014, 24, 1062-1066.	4.5	29
84	Sonic Hedgehog Signaling Is Active in Human Adrenal Cortex Development and Deregulated in Adrenocortical Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, E1209-E1216.	3.6	27
85	<i>ARMC5</i> Mutations Are a Frequent Cause of Primary Macronodular Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, E1501-E1509.	3.6	120
86	POD-1 binding to the E-box sequence inhibits SF-1 and StAR expression in human adrenocortical tumor cells. <i>Molecular and Cellular Endocrinology</i> , 2013, 371, 140-147.	3.2	28
87	Modulatory effect of Bcl GR gene polymorphisms on the obesity phenotype in Brazilian patients with Cushing's disease. <i>Clinics</i> , 2013, 68, 579-585.	1.5	8
88	PROP1 overexpression in corticotrophinomas: evidence for the role of PROP1 in the maintenance of cells committed to corticotrophic differentiation. <i>Clinics</i> , 2013, 68, 887-891.	1.5	9
89	The role of fibroblast growth factor receptor 4 overexpression and gene amplification as prognostic markers in pediatric and adult adrenocortical tumors. <i>Endocrine-Related Cancer</i> , 2012, 19, L11-L13.	3.1	19
90	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. <i>European Journal of Endocrinology</i> , 2012, 166, 61-67.	3.7	81

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91	Progression to Adrenocortical Tumorigenesis in Mice and Humans through Insulin-Like Growth Factor 2 and $\beta$ -Catenin. American Journal of Pathology, 2012, 181, 1017-1033.	3.8	154
92	Genotype analysis of the human endostatin variant p.D104N in benign and malignant adrenocortical tumors. Clinics, 2012, 67, 95-98.	1.5	6
93	ACTH-Independent Macronodular Adrenal Hyperplasia. Endocrinology and Metabolism, 2011, 26, 1.	3.0	4
94	Possible role of a radiation-induced p53 mutation in a Nelson's syndrome patient with a fatal outcome. Pituitary, 2011, 14, 400-404.	2.9	18
95	<sup>18</sup> F-FDG-PET/CT Imaging of ACTH-Independent Macronodular Adrenocortical Hyperplasia (AIMAH) Demonstrating Increased <sup>18</sup> F-FDG Uptake. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 3300-3301.	3.6	31
96	Glucose-dependent insulinotropic peptide receptor overexpression in adrenocortical hyperplasia in MEN1 syndrome without loss of heterozygosity at the 11q13 locus. Clinics, 2011, 66, 529-33.	1.5	6
97	Isolated familial somatotropinoma: 11q13-loh and gene/protein expression analysis suggests a possible involvement of <i>aipl1</i> also in non-pituitary tumorigenesis. Clinics, 2010, 65, 407-415.	1.5	33
98	Influence of the Fibroblast Growth Factor Receptor 4 Expression and the G388R Functional Polymorphism on Cushing's Disease Outcome. Journal of Clinical Endocrinology and Metabolism, 2010, 95, E271-E279.	3.6	17
99	High Penetrance of Pheochromocytoma Associated with the Novel C634Y/Y791F Double Germline Mutation in the <i>RET</i> Protooncogene. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1318-1327.	3.6	43
100	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.	3.6	66
101	A missense TCF1 mutation in a patient with <i>mody-3</i> and liver adenomatosis. Clinics, 2010, 65, 1059-1060.	1.5	7
102	ACTH-Independent Cushing's Syndrome: Adrenocortical Tumors. , 2010, , 189-208.		0
103	Stem Cell Markers Gene Expression in Corticotroph Pituitary Adenomas. , 2010, , P1-76-P1-76.		0
104	A Sellar Mass in a Patient with Systemic Non Hodgkin Lymphoma: Is There a Role for PET Scan in Differential Diagnosis?.. , 2010, , P1-297-P1-297.		0
105	Correlations between the Expression of BUBB1, DLG7 and PINK1 Genes and Outcomes in a Brazilian Cohort of Adrenocortical Tumors of Adult and Pediatric Patients.. , 2010, , P3-75-P3-75.		0
106	Clinical and Subclinical ACTH-Independent Macronodular Adrenal Hyperplasia (AIMAH) Affecting Members of a Large Brazilian Kindred.. , 2010, , P3-631-P3-631.		0
107	Expression profiles of the glucose-dependent insulinotropic peptide receptor and LHCGR in sporadic adrenocortical tumors. Journal of Endocrinology, 2009, 200, 167-175.	2.6	5
108	ADRENOCORTICAL CARCINOMA: A 30-YEAR EXPERIENCE AT A SINGLE INSTITUTION. Journal of Urology, 2009, 181, 8-9.	0.4	1

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109	Analysis of glucose-dependent insulinotropic peptide receptor (GIPR) and luteinizing hormone receptor (LHCGR) expression in human adrenocortical hyperplasia. <i>Arquivos Brasileiros De Endocrinologia E Metabologia</i> , 2009, 53, 326-331.	1.3	5
110	Expression of Insulin-Like Growth Factor-II and Its Receptor in Pediatric and Adult Adrenocortical Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008, 93, 3524-3531.	3.6	149
111	An Inhibin B and Estrogen-Secreting Adrenocortical Carcinoma Leading to Selective FSH Suppression. <i>Hormone Research in Paediatrics</i> , 2007, 67, 7-11.	1.8	11
112	The role of desmopressin in bilateral and simultaneous inferior petrosal sinus sampling for differential diagnosis of ACTH-dependent Cushing's syndrome. <i>Clinical Endocrinology</i> , 2006, 66, 061120012318003-???	2.4	55
113	Ectopic ACTH syndrome: our experience with 25 cases. <i>European Journal of Endocrinology</i> , 2006, 155, 725-733.	3.7	121
114	Deletion Mapping of Chromosome 17 in Benign and Malignant Adrenocortical Tumors Associated with the Arg337His Mutation of the p53 Tumor Suppressor Protein. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2005, 90, 2976-2981.	3.6	34
115	Cushing's Syndrome Secondary to Adrenocorticotropin-Independent Macronodular Adrenocortical Hyperplasia due to Activating Mutations of <i>GNAS1</i> Gene. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 2147-2151.	3.6	174
116	An Inherited Mutation Outside the Highly Conserved DNA-Binding Domain of the p53 Tumor Suppressor Protein in Children and Adults with Sporadic Adrenocortical Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 4970-4973.	3.6	183
117	An Inherited Mutation Outside the Highly Conserved DNA-Binding Domain of the p53 Tumor Suppressor Protein in Children and Adults with Sporadic Adrenocortical Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 4970-4973.	3.6	36
118	No evidence of somatic activating mutations on gonadotropin receptor genes in sex cord stromal tumors. <i>Fertility and Sterility</i> , 2000, 74, 992-995.	1.0	34
119	Mutation analysis of the follicle-stimulating hormone receptor gene in girls with gonadotropin-independent precocious puberty resulting from autonomous cystic ovaries. <i>Fertility and Sterility</i> , 2000, 73, 280-283.	1.0	18
120	Treatment of gonadotropin dependent precocious puberty due to hypothalamic hamartoma with gonadotropin releasing hormone agonist depot. <i>Archives of Disease in Childhood</i> , 1999, 80, 231-234.	1.9	45
121	Activating Mutation of the Stimulatory G Protein ( <i>gsp</i> ) as a Putative Cause of Ovarian and Testicular Human Stromal Leydig Cell Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1998, 83, 2074-2078.	3.6	93
122	Cortisol and adrenocorticotropin response to desmopressin in women with Cushing's disease compared with depressive illness. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1996, 81, 2233-2237.	3.6	32
123	The desmopressin stimulation test in the differential diagnosis of Cushing's syndrome. <i>Clinical Endocrinology</i> , 1993, 38, 463-472.	2.4	137