Constantine A Stratakis

List of Publications by Year
in descending order

390 papers	18,265 citations	19608 61 h-index	18075 120 g-index
398 all docs	398 docs citations	398 times ranked	11591 citing authors

#	Article	IF	Citations
1	Scoping review of COVID-19-related systematic reviews and meta-analyses: can we really have confidence in their results?. Postgraduate Medical Journal, 2022, 98, 372-379.	0.9	5
2	Cushing syndrome and glucocorticoids: T-cell lymphopenia, apoptosis, and rescue by IL-21. Journal of Allergy and Clinical Immunology, 2022, 149, 302-314.	1.5	4
3	KDM1A inactivation causes hereditary food-dependent Cushing syndrome. Genetics in Medicine, 2022, 24, 374-383.	1.1	27
4	The regulation of PKA signaling in obesity and in the maintenance of metabolic health. , 2022, 237, 108113.		35
5	Inflammatory biomarkers in the evaluation of pediatric endogenous Cushing syndrome. European Journal of Endocrinology, 2022, 186, 503-510.	1.9	4
6	Copeptin Levels Before and After Transsphenoidal Surgery for Cushing Disease: A Potential Early Marker of Remission. Journal of the Endocrine Society, 2022, 6, bvac053.	0.1	1
7	Duplications disrupt chromatin architecture and rewire GPR101-enhancer communication in X-linked acrogigantism. American Journal of Human Genetics, 2022, 109, 553-570.	2.6	18
8	Neurofibromatosis Type 1 Has a Wide Spectrum of Growth Hormone Excess. Journal of Clinical Medicine, 2022, 11, 2168.	1.0	6
9	Genetic Alterations in Benign Adrenal Tumors. Biomedicines, 2022, 10, 1041.	1.4	6
10	Steroidogenic Factor-1 Lineage Origin of Skin Lesions in Carney Complex Syndrome. Journal of Investigative Dermatology, 2022, 142, 2949-2957.e9.	0.3	3
11	USP13 genetics and expression in a family with thyroid cancer. Endocrine, 2022, 77, 281-290.	1.1	5
12	Steroid hormone analysis of adolescents and young women with polycystic ovarian syndrome and adrenocortical dysfunction using UPC2-MS/MS. Pediatric Research, 2021, 89, 118-126.	1.1	11
13	Genetics, clinical features and outcomes of non-syndromic pituitary gigantism: experience of a single center from Sao Paulo, Brazil. Pituitary, 2021, 24, 252-261.	1.6	4
14	Recovery of hypothalamicâ€pituitaryâ€adrenal axis in paediatric Cushing disease. Clinical Endocrinology, 2021, 94, 40-47.	1.2	8
15	Pde8b haploinsufficiency in mice is associated with modest adrenal defects, impaired steroidogenesis, and male infertility, unaltered by concurrent PKA or Wnt activation. Molecular and Cellular Endocrinology, 2021, 522, 111117.	1.6	2
16	Volumetric Modeling of Adrenal Gland Size in Primary Bilateral Macronodular Adrenocortical Hyperplasia. Journal of the Endocrine Society, 2021, 5, bvaa162.	0.1	7
17	A phosphodiesterase 11 (Pde11a) knockout mouse expressed functional but reduced Pde11a: Phenotype and impact on adrenocortical function. Molecular and Cellular Endocrinology, 2021, 520, 111071.	1.6	6
18	Genomic and sequence variants of protein kinase A regulatory subunit type 1Î² (PRKAR1B) in patients with adrenocortical disease and Cushing syndrome. Genetics in Medicine, 2021, 23, 174-182.	1.1	8

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19	Predicting the risk of cardiac myxoma in Carney complex. Genetics in Medicine, 2021, 23, 80-85.	1.1	23
20	First Somatic PRKAR1A Defect Associated With Mosaicism for Another PRKAR1A Mutation in a Patient With Cushing Syndrome. Journal of the Endocrine Society, 2021, 5, bvab007.	0.1	3
21	The X-linked acrogigantism-associated gene gpr101 is a regulator of early embryonic development and growth in zebrafish. Molecular and Cellular Endocrinology, 2021, 520, 111091.	1.6	7
22	Phosphodiesterase 2A and 3B variants are associated with primary aldosteronism. Endocrine-Related Cancer, 2021, 28, 1-13.	1.6	17
23	Insulin-like growth factor 2 (IGF2) expression in adrenocortical disease due to PRKAR1A mutations compared to other benign adrenal tumors. Endocrine, 2021, 72, 823-834.	1.1	1
24	Insulin sensitivity and pancreatic Î²-cell function in patients with primary aldosteronism. Endocrine, 2021, 72, 96-103.	1.1	8
25	Cushing Syndrome in a Pediatric Patient With a KCNJ5 Variant and Successful Treatment With Low-dose Ketoconazole. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 1606-1616.	1.8	4
26	A case of Carney triad complicated by renal cell carcinoma and a germline SDHA pathogenic variant. Endocrinology, Diabetes and Metabolism Case Reports, 2021, 2021, .	0.2	2
27	Corticotroph tumor progression after bilateral adrenalectomy (Nelsonâ€™s syndrome): systematic review and expert consensus recommendations. European Journal of Endocrinology, 2021, 184, P1-P16.	1.9	32
28	Molecular Genetic and Genomic Alterations in Cushingâ€™s Syndrome and Primary Aldosteronism. Frontiers in Endocrinology, 2021, 12, 632543.	1.5	19
29	The PRKAR1B p.R115K Variant is Associated with Lipoprotein Profile in African American Youth with Metabolic Challenges. Journal of the Endocrine Society, 2021, 5, bvab071.	0.1	3
30	Is there a common cause for paediatric Cushingâ€™s disease?. Endokrynologia Polska, 2021, 72, 104-107.	0.3	4
31	Variants in PRKAR1B cause a neurodevelopmental disorder with autism spectrum disorder, apraxia, and insensitivity to pain. Genetics in Medicine, 2021, 23, 1465-1473.	1.1	10
32	Copeptin Levels Before and After Transsphenoidal Surgery for Cushing Disease: A Potential Marker of Remission. Journal of the Endocrine Society, 2021, 5, A625-A625.	0.1	0
33	A Case of Carney Triad Complicated by Renal Cell Carcinoma and a Germline SDHA Pathogenic Variant. Journal of the Endocrine Society, 2021, 5, A985-A985.	0.1	0
34	Abnormal Pituitary Imaging and Associated Endocrine Dysfunctions in Erdheim-Chester Disease. Journal of the Endocrine Society, 2021, 5, A622-A622.	0.1	1
35	Contralateral Suppression Index Does Not Predict Clinical Cure in Patients Undergoing Surgery for Primary Aldosteronism. Annals of Surgical Oncology, 2021, 28, 7487-7495.	0.7	12
36	Family environment and development in children adopted from institutionalized care. Pediatric Research, 2021, , .	1.1	0

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37	Safety and Efficacy of Pegvisomant in Pediatric Growth Hormone Excess. Journal of the Endocrine Society, 2021, 5, A648-A648.	0.1	0
38	Selective Serotonin Reuptake Inhibitors Increase Urinary Free Cortisol in Patients with Carney Complex and Primary Pigmented Nodular Adrenocortical Disease. Journal of the Endocrine Society, 2021, 5, A95-A95.	0.1	0
39	Potential Role for the RASD1 Glucocorticoid-Responsive Gene in Corticotroph Tumorigenesis. Journal of the Endocrine Society, 2021, 5, A549-A549.	0.1	0
40	Health-Related Quality of Life in Cushing Disease: Discrepancy Between Parent and Child Reports. Journal of the Endocrine Society, 2021, 5, A717-A718.	0.1	0
41	Inhibition of Aurora kinase A activity enhances the antitumor response of beta-catenin blockade in human adrenocortical cancer cells. Molecular and Cellular Endocrinology, 2021, 528, 111243.	1.6	7
42	Carney Triad, Carney-Stratakis Syndrome, 3PAS and Other Tumors Due to SDH Deficiency. Frontiers in Endocrinology, 2021, 12, 680609.	1.5	11
43	Whole-exome sequencing reveals insights into genetic susceptibility to Congenital Zika Syndrome. PLoS Neglected Tropical Diseases, 2021, 15, e0009507.	1.3	5
44	Homozygous SHBG Variant (rs6258) Linked to Gonadotropin-Independent Precocious Puberty in a Young Girl. Journal of the Endocrine Society, 2021, 5, bvab125.	0.1	0
45	PRKAR1A and Thyroid Tumors. Cancers, 2021, 13, 3834.	1.7	11
46	Loss of PKA regulatory subunit 1Î± aggravates cardiomyocyte necrosis and myocardial ischemia/reperfusion injury. Journal of Biological Chemistry, 2021, 297, 100850.	1.6	11
47	Pituitary Imaging Abnormalities and Related Endocrine Disorders in Erdheimâ€“Chester Disease. Cancers, 2021, 13, 4126.	1.7	4
48	Paediatric patients with Cushing disease and negative pituitary MRI have a higher risk of nonremission after transsphenoidal surgery. Clinical Endocrinology, 2021, 95, 856-862.	1.2	5
49	Pediatric Cushingâ€™s syndrome: greater risk of being overweight or obese after long-term remission and its predictive factors. European Journal of Endocrinology, 2021, 184, 179-187.	1.9	3
50	Molecular Endocrinology, Endocrine Genetics, and Precision Medicine. , 2021, , 9-29.		1
51	Inherited Neuroendocrine Neoplasms. , 2021, , 409-459.		7
52	Younger age and early puberty are associated with cognitive function decline in children with Cushing disease. Clinical Endocrinology, 2021, , .	1.2	3
53	CYP11B1 variants influence skeletal maturation via alternative splicing. Communications Biology, 2021, 4, 1274.	2.0	3
54	Lower hair cortisol among patients with sickle cell disease may indicate decreased adrenal reserves. American Journal of Blood Research, 2021, 11, 140-148.	0.6	0

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55	Association between Maternal Non-Coding Interferon-Î» Polymorphisms and Congenital Zika Syndrome in a Cohort from Brazilian Northeast. Viruses, 2021, 13, 2253.	1.5	1
56	Protein kinase A drives paracrine crisis and WNT4-dependent testis tumor in Carney complex. Journal of Clinical Investigation, 2021, 131, .	3.9	2
57	Chaperones, somatotroph tumors and the cyclic AMP (cAMP)-dependent protein kinase (PKA) pathway. Molecular and Cellular Endocrinology, 2020, 499, 110607.	1.6	8
58	Mosaicism for KCNJ5 Causing Early-Onset Primary Aldosteronism due to Bilateral Adrenocortical Hyperplasia. American Journal of Hypertension, 2020, 33, 124-130.	1.0	20
59	Aggressive pituitary tumors in the young and elderly. Reviews in Endocrine and Metabolic Disorders, 2020, 21, 213-223.	2.6	22
60	Preventing disease in the twenty-first century: â€œLife is short, the Art long, opportunity fleetingâ€¦â€• Pediatric Research, 2020, 87, 181-182.	1.1	0
61	Computerized Analysis of Brain MRI Parameter Dynamics in Young Patients With Cushing Syndromeâ€”A Case-Control Study. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2069-e2077.	1.8	7
62	The Genetics of Pituitary Adenomas. Journal of Clinical Medicine, 2020, 9, 30.	1.0	37
63	Carney complex syndrome manifesting as cardioembolic stroke: a case report and review of the literature. International Journal of Neuroscience, 2020, , 1-7.	0.8	5
64	Kisspeptin deficiency leads to abnormal adrenal glands and excess steroid hormone secretion. Human Molecular Genetics, 2020, 29, 3443-3450.	1.4	3
65	Prevalence of Diabetes and Hypertension and Their Associated Risks for Poor Outcomes in Covid-19 Patients. Journal of the Endocrine Society, 2020, 4, bvaa102.	0.1	56
66	Rare Germline DICER1 Variants in Pediatric Patients With Cushing's Disease: What Is Their Role?. Frontiers in Endocrinology, 2020, 11, 433.	1.5	7
67	Hemodynamics of Prefrontal Cortex in Ornithine Transcarbamylase Deficiency: A Twin Case Study. Frontiers in Neurology, 2020, 11, 809.	1.1	3
68	Prkar1a haploinsufficiency ameliorates the growth hormone excess phenotype in Aip-deficient mice. Human Molecular Genetics, 2020, 29, 2951-2961.	1.4	2
69	A Century After the Description of â€œHormonesâ€: Our Golden Jubilee Celebration Goes on with What is New in Endocrine Oncology: And a Lot is New!. Hormone and Metabolic Research, 2020, 52, 551-552.	0.7	0
70	ARMC5 Alterations in Patients With Sporadic Neuroendocrine Tumors and Multiple Endocrine Neoplasia Type 1 (MEN1). Journal of Clinical Endocrinology and Metabolism, 2020, 105, e4531-e4542.	1.8	5
71	Letter to the Editor from Berthon: â€œCardiac Myxoma Caused by Fumarate Hydratase Gene Deletion in Patient With Cortisol-Secreting Adrenocortical Adenomaâ€• Journal of Clinical Endocrinology and Metabolism, 2020, 105, e4183-e4184.	1.8	1
72	Germline CDKN1B Loss-of-Function Variants Cause Pediatric Cushingâ€™s Disease With or Without an MEN4 Phenotype. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1983-2005.	1.8	31

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73	Epidemics will always come (and go): The need to prepare for the next one, research on COVID-19, and the role of molecular and cellular endocrinology. Molecular and Cellular Endocrinology, 2020, 511, 110863.	1.6	3
74	PRKAR1A deficiency impedes hypertrophy and reduces heart size. Physiological Reports, 2020, 8, e14405.	0.7	8
75	Adrenocortical tumorigenesis: Lessons from genetics. Best Practice and Research in Clinical Endocrinology and Metabolism, 2020, 34, 101428.	2.2	36
76	Endocrine Conditions and COVID-19. Hormone and Metabolic Research, 2020, 52, 471-484.	0.7	34
77	A Gene-Based Classification of Primary Adrenocortical Hyperplasias. Hormone and Metabolic Research, 2020, 52, 133-141.	0.7	15
78	Germline Variants in Phosphodiesterase Genes and Genetic Predisposition to Pediatric Adrenocortical Tumors. Cancers, 2020, 12, 506.	1.7	17
79	Clinical characteristics and outcomes of SDHB-related pheochromocytoma and paraganglioma in children and adolescents. Journal of Cancer Research and Clinical Oncology, 2020, 146, 1051-1063.	1.2	30
80	Medical Treatment of Pituitary Adenomas: A Celebration of Endocrinology (and Oncology)!. Hormone and Metabolic Research, 2020, 52, 7-7.	0.7	2
81	Editorial: Congenital Adrenal Hyperplasia, Unresolved Issues and Implications on Clinical Management. Frontiers in Endocrinology, 2020, 11, 170.	1.5	2
82	Cushing syndrome: Old and new genes. Best Practice and Research in Clinical Endocrinology and Metabolism, 2020, 34, 101418.	2.2	13
83	Letter to the Editor: â€œIGSF1 Deficiency Results in Human and Murine Somatotrope Neurosecretory Hyperfunctionâ€: Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2310-e2310.	1.8	0
84	Loss of habenular Prkar2a reduces hedonic eating and increases exercise motivation. JCI Insight, 2020, 5, .	2.3	8
85	The Association of ARMC5 with the Renin-Angiotensin-Aldosterone System, Blood Pressure, and Glycemia in African Americans. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 2625-2633.	1.8	9
86	Liver findings in patients with Carney complex, germline PRKAR1A pathogenic variants, and link to cardiac myxomas. Endocrine-Related Cancer, 2020, 27, 355-360.	1.6	2
87	HEREDITARY ENDOCRINE TUMOURS: CURRENT STATE-OF-THE-ART AND RESEARCH OPPORTUNITIES: The roles of AIP and GPR101 in familial isolated pituitary adenomas (FIPA). Endocrine-Related Cancer, 2020, 27, T77-T86.	1.6	11
88	HEREDITARY ENDOCRINE TUMOURS: CURRENT STATE-OF-THE-ART AND RESEARCH OPPORTUNITIES: GPR101, an orphan GPCR with roles in growth and pituitary tumorigenesis. Endocrine-Related Cancer, 2020, 27, T87-T97.	1.6	12
89	Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. Endocrine-Related Cancer, 2020, 27, 403-413.	1.6	13
90	ARMC5 variants in PRKAR1A-mutated patients modify cortisol levels and Cushingâ€™s syndrome. Endocrine-Related Cancer, 2020, 27, 509-517.	1.6	7

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91	PRKACB variants in skeletal disease or adrenocortical hyperplasia: effects on protein kinase A. Endocrine-Related Cancer, 2020, 27, 647-656.	1.6	7
92	PKA functions in metabolism and resistance to obesity: lessons from mouse and human studies. Journal of Endocrinology, 2020, 246, R51-R64.	1.2	50
93	Acute Statin Administration Reduces Levels of Steroid Hormone Precursors. Hormone and Metabolic Research, 2020, 52, 742-746.	0.7	3
94	SUN-235 Deficient Fear Extinction in PRKAR1A-Defective Mice. Journal of the Endocrine Society, 2020, 4, .	0.1	0
95	SAT-543 Human Hair Aldosterone Measurements for Evaluation of Primary Aldosteronism. Journal of the Endocrine Society, 2020, 4, .	0.1	0
96	OR24-06 USP8 Genetic Variants May Contribute to the Development of Bilateral Adrenal Hyperplasia and ACTH-Independent Cushing Syndrome. Journal of the Endocrine Society, 2020, 4, .	0.1	0
97	SUN-713 Prevalence of Renal Cysts in Patients with Carney Complex. Journal of the Endocrine Society, 2020, 4, .	0.1	0
98	Telomere Length Changes in Children With Cushing Disease: A Pilot Study. Journal of the Endocrine Society, 2020, 4, bvaa067.	0.1	1
99	Curative resection of an aldosteronoma causing primary aldosteronism in the second trimester of pregnancy. Endocrinology, Diabetes and Metabolism Case Reports, 2020, 2020, .	0.2	2
100	A SOX5 gene variant as a possible contributor to short stature. Endocrinology, Diabetes and Metabolism Case Reports, 2020, 2020, .	0.2	2
101	OR23-01 Intrapatient ACTH Variability in Cushingâ€™s Disease: Prognostic Significance. Journal of the Endocrine Society, 2020, 4, .	0.1	0
102	Inhibin A as a tumor marker for primary bilateral macronodular adrenal hyperplasia. Endocrinology, Diabetes and Metabolism Case Reports, 2020, 2020, .	0.2	0
103	PRKAR1A deficiency delays postnatal heart growth. FASEB Journal, 2020, 34, 1-1.	0.2	0
104	SUN-917 Aggressive De Novo MEN1 Variant in a Child with Metastatic Pancreatic Acth and Crh Co-Secreting Neuroendocrine Tumor: Diagnosis and 10-Year Follow Up. Journal of the Endocrine Society, 2020, 4, .	0.1	0
105	OR06-01 The Role of Germline Defects in Cushingâ€™s Disease. Journal of the Endocrine Society, 2020, 4, .	0.1	0
106	OR22-07 Novel Variants in Protein Kinase a Signaling-Related Genes Identified in Obese Children with and Without NAFLD. Journal of the Endocrine Society, 2020, 4, .	0.1	1
107	MON-190 Telomere Length as a Novel Prognostic Marker of Cushing Complications. Journal of the Endocrine Society, 2020, 4, .	0.1	0
108	SAT-304 Pituitary Stem Cells May Drive Adenomas Causing Cushingâ€™s Disease. Journal of the Endocrine Society, 2020, 4, .	0.1	0

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109	Abstract MP166: PRKAR1A Deficiency Abrogates Cardiac Hypertrophy Through Inhibition of Mitochondrial Fission. Circulation Research, 2020, 127, .	2.0	0
110	c-KIT oncogene expression in PRKAR1A-mutant adrenal cortex. Endocrine-Related Cancer, 2020, 27, 591-599.	1.6	0
111	c-KIT oncogene expression in PRKAR1A-mutant adrenal cortex. Endocrine-Related Cancer, 2020, 27, 591-599.	1.6	1
112	Molecular mechanisms of ARMC5 mutations in adrenal pathophysiology. Current Opinion in Endocrine and Metabolic Research, 2019, 8, 104-111.	0.6	16
113	Pheochromocytomas: Fabulous, Fascinating, and First (in everything)!. Hormone and Metabolic Research, 2019, 51, 401-402.	0.7	2
114	Somatic PRKAR1A Gene Mutation in a Nonsyndromic Metastatic Large Cell Calcifying Sertoli Cell Tumor. Journal of the Endocrine Society, 2019, 3, 1375-1382.	0.1	16
115	Cushing disease in a patient with nonbullous congenital ichthyosiform erythroderma: lessons in avoiding glucocorticoids in ichthyosis. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 911-914.	0.4	0
116	Multiple Endocrine Neoplasia Type 1 (MEN1): An Update and the Significance of Early Genetic and Clinical Diagnosis. Frontiers in Endocrinology, 2019, 10, 339.	1.5	118
117	ARMC5 Variants and Risk of Hypertension in Blacks: MHâ€GRID Study. Journal of the American Heart Association, 2019, 8, e012508.	1.6	8
118	Resistant Hypertension. Endocrinology and Metabolism Clinics of North America, 2019, 48, 811-828.	1.2	5
119	Illicit Upregulation of Serotonin Signaling Pathway in Adrenals of Patients With High Plasma or Intra-Adrenal ACTH Levels. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4967-4980.	1.8	15
120	Germline USP8 Mutation Associated With Pediatric Cushing Disease and Other Clinical Features: A New Syndrome. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 4676-4682.	1.8	45
121	Hormone and Metabolic Research: 50 Years of Research. Hormone and Metabolic Research, 2019, 51, 8-10.	0.7	1
122	Subspecialty training in adult inherited metabolic diseases: a call to action for unmet needs. Lancet Diabetes and Endocrinology,the, 2019, 7, 82-84.	5.5	7
123	The Catalytic Subunit Î² of PKA Affects Energy Balance and Catecholaminergic Activity. Journal of the Endocrine Society, 2019, 3, 1062-1078.	0.1	9
124	Called and Uncalled for Functions of the Main Catalytic Subunit of Protein Kinase A: One Enzyme, Many Faces. Endocrinology, 2019, 160, 1674-1676.	1.4	2
125	CRH stimulation improves 18F-FDG-PET detection of pituitary adenomas in Cushingâ€™s disease. Endocrine, 2019, 65, 155-165.	1.1	25
126	Inflammation and Metabolism in Cancer Cellâ€"Mitochondria Key Player. Frontiers in Oncology, 2019, 9, 348.	1.3	115

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127	Large Genomic Aberrations in Corticotropinomas Are Associated With Greater Aggressiveness. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 1792-1801.	1.8	20
128	High expression of adrenal P450 aromatase (CYP19A1) in association with ARMC5-primary bilateral macronodular adrenocortical hyperplasia. Journal of Steroid Biochemistry and Molecular Biology, 2019, 191, 105316.	1.2	13
129	Clinical, Diagnostic, and Treatment Characteristics of SDHA-Related Metastatic Pheochromocytoma and Paraganglioma. Frontiers in Oncology, 2019, 9, 53.	1.3	39
130	SGPL1 Deficiency: A Rare Cause of Primary Adrenal Insufficiency. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 1484-1490.	1.8	27
131	Genetics of Hypertension in African Americans and Others of African Descent. International Journal of Molecular Sciences, 2019, 20, 1081.	1.8	43
132	CD40LG duplications in patients with X-LAG syndrome commonly undergo random X-chromosome inactivation. Journal of Allergy and Clinical Immunology, 2019, 143, 1659.	1.5	4
133	Genetic Characteristics of Aldosterone-Producing Adenomas in Blacks. Hypertension, 2019, 73, 885-892.	1.3	121
134	A novel mutation in the glucocorticoid receptor gene as a cause of severe glucocorticoid resistance complicated by hypertensive encephalopathy. Journal of Hypertension, 2019, 37, 1475-1481.	0.3	12
135	Variations in maternal adenylate cyclase genes are associated with congenital Zika syndrome in a cohort from Northeast, Brazil. Journal of Internal Medicine, 2019, 285, 215-222.	2.7	18
136	Carney Complex. Experimental and Clinical Endocrinology and Diabetes, 2019, 127, 156-164.	0.6	84
137	Optical Imaging Technology: A Useful Tool to Identify Remission in Cushing Disease After Surgery. Hormone and Metabolic Research, 2019, 51, 120-126.	0.7	1
138	Growth hormone excess in neurofibromatosis 1. Genetics in Medicine, 2019, 21, 1254-1255.	1.1	13
139	The 3PAs: An Update on the Association of Pheochromocytomas, Paragangliomas, and Pituitary Tumors. Hormone and Metabolic Research, 2019, 51, 419-436.	0.7	22
140	Genetic Tumor Syndromes with Endocrine Involvement: A Compendium and an Update. Pediatric Endocrinology Reviews, 2019, 16, 311-334.	1.2	0
141	Pediatric Cushing Syndrome; an Overview. Pediatric Endocrinology Reviews, 2019, 17, 100-109.	1.2	6
142	Incidence of Autoimmune and Related Disorders After Resolution of Endogenous Cushing Syndrome in Children. Hormone and Metabolic Research, 2018, 50, 290-295.	0.7	9
143	Anxiety-like behavior and other consequences of early life stress in mice with increased protein kinase A activity. Behavioural Brain Research, 2018, 348, 22-30.	1.2	3
144	An update on Cushing syndrome in pediatrics. Annales D'Endocrinologie, 2018, 79, 125-131.	0.6	34

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145	Neonatal Cushing Syndrome. Clinics in Perinatology, 2018, 45, 103-118.	0.8	22
146	Succinate dehydrogenase (SDH) deficiency, Carney triad and the epigenome. Molecular and Cellular Endocrinology, 2018, 469, 107-111.	1.6	45
147	An update on the genetics of benign pituitary adenomas in children and adolescents. Current Opinion in Endocrine and Metabolic Research, 2018, 1, 19-24.	0.6	14
148	An orphan G-protein-coupled receptor causes human gigantism and/or acromegaly: Molecular biology and clinical correlations. Best Practice and Research in Clinical Endocrinology and Metabolism, 2018, 32, 125-140.	2.2	26
149	Cyclic 3â€²,5â€²-adenosine monophosphate (cAMP) signaling in theÂanterior pituitary gland in health and disease. Molecular and Cellular Endocrinology, 2018, 463, 72-86.	1.6	27
150	The Spectrum of Thyroid Gland Pathology in Carney Complex. American Journal of Surgical Pathology, 2018, 42, 587-594.	2.1	35
151	Successful Treatment of Estrogen Excess in Primary Bilateral Macronodular Adrenocortical Hyperplasia with Leuprolide Acetate. Hormone and Metabolic Research, 2018, 50, 124-132.	0.7	7
152	Decreased lymphocytes and increased risk for infection are common in endogenous pediatric Cushing syndrome. Pediatric Research, 2018, 83, 431-437.	1.1	24
153	Cyclic AMPâ€dependent protein kinase catalytic subunit A (PRKACA): the expected, the unexpected, and what might be next. Journal of Pathology, 2018, 244, 257-259.	2.1	16
154	Superiority of 68Ga-DOTATATE over 18F-FDG and anatomic imaging in the detection of succinate dehydrogenase mutation (SDHx)-related pheochromocytoma and paraganglioma in the pediatric population. European Journal of Nuclear Medicine and Molecular Imaging, 2018, 45, 787-797.	3.3	64
155	Failure to Thrive in the Context of Carney Complex. Hormone Research in Paediatrics, 2018, 89, 38-46.	0.8	3
156	Xq26.3 Duplication in a Boy With Motor Delay and Low Muscle Tone Refines the X-Linked Acrogigantism Genetic Locus. Journal of the Endocrine Society, 2018, 2, 1100-1108.	0.1	7
157	Medullary thyroid cancer, leukemia, mesothelioma and meningioma associated with germline APC and RASAL1 variants: a new syndrome?. Hormones, 2018, 16, 423-428.	0.9	3
158	An update on adrenal endocrinology: significant discoveries in the last 10Âyears and where the field is heading in the next decade. Hormones, 2018, 17, 479-490.	0.9	5
159	Primary hypophysitis and other autoimmune disorders of the sellar and suprasellar regions. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 335-347.	2.6	34
160	Mini-review of hair cortisol concentration for evaluation of Cushing syndrome. Expert Review of Endocrinology and Metabolism, 2018, 13, 225-231.	1.2	24
161	An overview of inborn errors of metabolism manifesting with primary adrenal insufficiency. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 53-67.	2.6	13
162	Children with <scp>MEN</scp>1 gene mutations may present first (and at a young age) with Cushing disease. Clinical Endocrinology, 2018, 89, 437-443.	1.2	19

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163	Genetics of Cushingâ€™s Syndrome. Endocrinology and Metabolism Clinics of North America, 2018, 47, 275-297.	1.2	33
164	Cushingâ€™s Syndrome in Pediatrics. Endocrinology and Metabolism Clinics of North America, 2018, 47, 451-462.	1.2	60
165	Interaction of AIP with protein kinase A (cAMP-dependent protein kinase). Human Molecular Genetics, 2018, 27, 2604-2613.	1.4	25
166	Genetics of micronodular adrenal hyperplasia and Carney complex. Presse Medicale, 2018, 47, e127-e137.	0.8	17
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