Constantine A Stratakis

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

63 361 14,139 109 h-index g-index citations papers 16,924 7.1 393 5.4 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
361	The regulation of PKA signaling in obesity and in the maintenance of metabolic health <i>Pharmacology & Therapeutics</i> , 2022 , 237, 108113	13.9	3
360	Copeptin Levels Before and After Transsphenoidal Surgery for Cushing Disease: A Potential Early Marker of Remission <i>Journal of the Endocrine Society</i> , 2022 , 6, bvac053	0.4	1
359	Duplications disrupt chromatin architecture and rewire GPR101-enhancer communication in X-linked acrogigantism <i>American Journal of Human Genetics</i> , 2022 ,	11	1
358	Genetic Alterations in Benign Adrenal Tumors. <i>Biomedicines</i> , 2022 , 10, 1041	4.8	0
357	CYP11B1 variants influence skeletal maturation via alternative splicing. <i>Communications Biology</i> , 2021 , 4, 1274	6.7	O
356	Lower hair cortisol among patients with sickle cell disease may indicate decreased adrenal reserves. American Journal of Blood Research, 2021 , 11, 140-148	1.6	
355	KDM1A inactivation causes hereditary food-dependent Cushing syndrome <i>Genetics in Medicine</i> , 2021 ,	8.1	2
354	Inherited Neuroendocrine Neoplasms 2021 , 409-459		5
353	A case of Carney triad complicated by renal cell carcinoma and a germline SDHA pathogenic variant. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2021 , 2021,	1.4	1
352	Corticotroph tumor progression after bilateral adrenalectomy (Nelson's syndrome): systematic review and expert consensus recommendations. <i>European Journal of Endocrinology</i> , 2021 , 184, P1-P16	6.5	6
351	Molecular Genetic and Genomic Alterations in Cushing's Syndrome and Primary Aldosteronism. <i>Frontiers in Endocrinology</i> , 2021 , 12, 632543	5.7	6
350	The PRKAR1B p.R115K Variant is Associated with Lipoprotein Profile in African American Youth with Metabolic Challenges. <i>Journal of the Endocrine Society</i> , 2021 , 5, bvab071	0.4	0
349	Is there a common cause for paediatric Cushing's disease?. <i>Endokrynologia Polska</i> , 2021 , 72, 104-107	1.1	1
348	Variants in PRKAR1B cause a neurodevelopmental disorder with autism spectrum disorder, apraxia, and insensitivity to pain. <i>Genetics in Medicine</i> , 2021 , 23, 1465-1473	8.1	1
347	Copeptin Levels Before and After Transsphenoidal Surgery for Cushing Disease: A Potential Marker of Remission. <i>Journal of the Endocrine Society</i> , 2021 , 5, A625-A625	0.4	
346	A Case of Carney Triad Complicated by Renal Cell Carcinoma and a Germline SDHA Pathogenic Variant. <i>Journal of the Endocrine Society</i> , 2021 , 5, A985-A985	0.4	78
345	Abnormal Pituitary Imaging and Associated Endocrine Dysfunctions in Erdheim-Chester Disease. <i>Journal of the Endocrine Society</i> , 2021 , 5, A622-A622	0.4	1

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344	Contralateral Suppression Index Does Not Predict Clinical Cure in Patients Undergoing Surgery for Primary Aldosteronism. <i>Annals of Surgical Oncology</i> , 2021 , 28, 7487-7495	3.1	2	
343	Safety and Efficacy of Pegvisomant in Pediatric Growth Hormone Excess. <i>Journal of the Endocrine Society</i> , 2021 , 5, A648-A648	0.4	78	
342	Selective Serotonin Reuptake Inhibitors Increase Urinary Free Cortisol in Patients with Carney Complex and Primary Pigmented Nodular Adrenocortical Disease. <i>Journal of the Endocrine Society</i> , 2021 , 5, A95-A95	0.4	78	
341	Potential Role for the RASD1 Glucocorticoid-Responsive Gene in Corticotroph Tumorigenesis. Journal of the Endocrine Society, 2021 , 5, A549-A549	0.4	78	
340	Health-Related Quality of Life in Cushing Disease: Discrepancy Between Parent and Child Reports. Journal of the Endocrine Society, 2021 , 5, A717-A718	0.4	78	
339	Inhibition of Aurora kinase A activity enhances the antitumor response of beta-catenin blockade in human adrenocortical cancer cells. <i>Molecular and Cellular Endocrinology</i> , 2021 , 528, 111243	4.4	3	
338	Carney Triad, Carney-Stratakis Syndrome, 3PAS and Other Tumors Due to SDH Deficiency. <i>Frontiers in Endocrinology</i> , 2021 , 12, 680609	5.7	1	
337	Whole-exome sequencing reveals insights into genetic susceptibility to Congenital Zika Syndrome. <i>PLoS Neglected Tropical Diseases</i> , 2021 , 15, e0009507	4.8	2	
336	Cushing syndrome and glucocorticoids: T-cell lymphopenia, apoptosis, and rescue by IL-21. <i>Journal of Allergy and Clinical Immunology</i> , 2021 ,	11.5	1	
335	Homozygous Variant () Linked to Gonadotropin-Independent Precocious Puberty in a Young Girl. Journal of the Endocrine Society, 2021 , 5, bvab125	0.4		
334	Steroid hormone analysis of adolescents and young women with polycystic ovarian syndrome and adrenocortical dysfunction using UPC-MS/MS. <i>Pediatric Research</i> , 2021 , 89, 118-126	3.2	5	
333	Genetics, clinical features and outcomes of non-syndromic pituitary gigantism: experience of a single center from Sao Paulo, Brazil. <i>Pituitary</i> , 2021 , 24, 252-261	4.3		
332	Recovery of hypothalamic-pituitary-adrenal axis in paediatric Cushing disease. <i>Clinical Endocrinology</i> , 2021 , 94, 40-47	3.4	5	
331	Pde8b haploinsufficiency in mice is associated with modest adrenal defects, impaired steroidogenesis, and male infertility, unaltered by concurrent PKA or Wnt activation. <i>Molecular and Cellular Endocrinology</i> , 2021 , 522, 111117	4.4	О	
330	Volumetric Modeling of Adrenal Gland Size in Primary Bilateral Macronodular Adrenocortical Hyperplasia. <i>Journal of the Endocrine Society</i> , 2021 , 5, bvaa162	0.4	2	
329	A phosphodiesterase 11 (Pde11a) knockout mouse expressed functional but reduced Pde11a: Phenotype and impact on adrenocortical function. <i>Molecular and Cellular Endocrinology</i> , 2021 , 520, 111	o 1 ·1	Ο	
328	Genomic and sequence variants of protein kinase A regulatory subunit type 1[[PRKAR1B] in patients with adrenocortical disease and Cushing syndrome. <i>Genetics in Medicine</i> , 2021 , 23, 174-182	8.1	5	
327	Predicting the risk of cardiac myxoma in Carney complex. <i>Genetics in Medicine</i> , 2021 , 23, 80-85	8.1	4	

326	First Somatic Defect Associated With Mosaicism for Another Mutation in a Patient With Cushing Syndrome. <i>Journal of the Endocrine Society</i> , 2021 , 5, bvab007	0.4	1
325	The X-linked acrogigantism-associated gene gpr101 is a regulator of early embryonic development and growth in zebrafish. <i>Molecular and Cellular Endocrinology</i> , 2021 , 520, 111091	4.4	2
324	Phosphodiesterase 2A and 3B variants are associated with primary aldosteronism. Endocrine-Related Cancer, 2021 , 28, 1-13	5.7	7
323	Insulin-like growth factor 2 (IGF2) expression in adrenocortical disease due to PRKAR1A mutations compared to other benign adrenal tumors. <i>Endocrine</i> , 2021 , 72, 823-834	4	O
322	Insulin sensitivity and pancreatic Etell function in patients with primary aldosteronism. <i>Endocrine</i> , 2021 , 72, 96-103	4	3
321	Scoping review of COVID-19-related systematic reviews and meta-analyses: can we really have confidence in their results?. <i>Postgraduate Medical Journal</i> , 2021 ,	2	1
320	Cushing Syndrome in a Pediatric Patient With a KCNJ5 Variant and Successful Treatment With Low-dose Ketoconazole. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, 1606-1616	5.6	0
319	Loss of PKA regulatory subunit 1daggravates cardiomyocyte necrosis and myocardial ischemia/reperfusion injury. <i>Journal of Biological Chemistry</i> , 2021 , 297, 100850	5.4	5
318	Pituitary Imaging Abnormalities and Related Endocrine Disorders in Erdheim-Chester Disease. <i>Cancers</i> , 2021 , 13,	6.6	2
317	Paediatric patients with Cushing disease and negative pituitary MRI have a higher risk of nonremission after transsphenoidal surgery. <i>Clinical Endocrinology</i> , 2021 , 95, 856-862	3.4	1
316	Pediatric Cushing's syndrome: greater risk of being overweight or obese after long-term remission and its predictive factors. <i>European Journal of Endocrinology</i> , 2021 , 184, 179-187	6.5	0
315	Molecular Endocrinology, Endocrine Genetics, and Precision Medicine 2021 , 9-29		1
314	Germline CDKN1B Loss-of-Function Variants Cause Pediatric Cushing's Disease With or Without an MEN4 Phenotype. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	10
313	PRKAR1A deficiency impedes hypertrophy and reduces heart size. <i>Physiological Reports</i> , 2020 , 8, e1440	52.6	4
312	Adrenocortical tumorigenesis: Lessons from genetics. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2020 , 34, 101428	6.5	15
311	Endocrine Conditions and COVID-19. Hormone and Metabolic Research, 2020, 52, 471-484	3.1	20
310	A Gene-Based Classification of Primary Adrenocortical Hyperplasias. <i>Hormone and Metabolic Research</i> , 2020 , 52, 133-141	3.1	11
309	Germline Variants in Phosphodiesterase Genes and Genetic Predisposition to Pediatric Adrenocortical Tumors. <i>Cancers</i> , 2020 , 12,	6.6	12

(2020-2020)

Clinical characteristics and outcomes of SDHB-related pheochromocytoma and paraganglioma in children and adolescents. <i>Journal of Cancer Research and Clinical Oncology</i> , 2020 , 146, 1051-1063	4.9	14	
Medical Treatment of Pituitary Adenomas: A Celebration of Endocrinology (and Oncology)!. <i>Hormone and Metabolic Research</i> , 2020 , 52, 7	3.1	2	
Cushing syndrome: Old and new genes. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2020 , 34, 101418	6.5	4	
Loss of habenular Prkar2a reduces hedonic eating and increases exercise motivation. <i>JCI Insight</i> , 2020 , 5,	9.9	4	
The Association of ARMC5 with the Renin-Angiotensin-Aldosterone System, Blood Pressure, and Glycemia in African Americans. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	5	
Liver findings in patients with Carney complex, germline PRKAR1A pathogenic variants, and link to cardiac myxomas. <i>Endocrine-Related Cancer</i> , 2020 , 27, 355-360	5.7	1	
HEREDITARY ENDOCRINE TUMOURS: CURRENT STATE-OF-THE-ART AND RESEARCH OPPORTUNITIES: The roles of AIP and GPR101 in familial isolated pituitary adenomas (FIPA). <i>Endocrine-Related Cancer</i> , 2020 , 27, T77-T86	5.7	7	
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	5.7	7	
Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. <i>Endocrine-Related Cancer</i> , 2020 , 27, 403-413	5.7	4	
ARMC5 variants in PRKAR1A-mutated patients modify cortisol levels and Cushing's syndrome. <i>Endocrine-Related Cancer</i> , 2020 , 27, 509-517	5.7	2	
PRKACB variants in skeletal disease or adrenocortical hyperplasia: effects on protein kinase A. <i>Endocrine-Related Cancer</i> , 2020 , 27, 647-656	5.7	4	
PKA functions in metabolism and resistance to obesity: lessons from mouse and human studies. <i>Journal of Endocrinology</i> , 2020 , 246, R51-R64	4.7	11	
OR23-01 Intrapatient ACTH Variability in Cushing® Disease: Prognostic Significance. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78	
PRKAR1A deficiency delays postnatal heart growth. <i>FASEB Journal</i> , 2020 , 34, 1-1	0.9		
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. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78	
OR06-01 The Role of Germline Defects in Cushing Disease. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78	
MON-190 Telomere Length as a Novel Prognostic Marker of Cushing Complications. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78	
SAT-304 Pituitary Stem Cells May Drive Adenomas Causing Cushing Disease. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78	
	children and adolescents. Journal of Cancer Research and Clinical Oncology, 2020, 146, 1051-1063 Medical Treatment of Pituitary Adenomas: A Celebration of Endocrinology (and Oncology)!. Hormone and Metabolic Research, 2020, 52, 7 Cushing syndrome: Old and new genes. Best Practice and Research in Clinical Endocrinology and Metabolism, 2020, 34, 101418 Loss of habenular Prkar2a reduces hedonic eating and increases exercise motivation. JCI Insight, 2020, 5, 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, Liver findings in patients with Carney complex, germline PRKAR1A pathogenic variants, and link to cardiac myxomas. Endocrine-Related Cancer, 2020, 27, 355-360 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, T777-186 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, 187-197 Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. Endocrine-Related Cancer, 2020, 27, 509-517 PRKACB variants in PRKAR1A-mutated patients modify cortisol levels and Cushing's syndrome. Endocrine-Related Cancer, 2020, 27, 647-656 PKA functions in metabolism and resistance to obesity; lessons from mouse and human studies. Journal of Endocrine Society, 2020, 4, R51-R64 OR23-O1 Intrapatient ACTH Variability in Cushing® Disease: Prognostic Significance. Journal of the Endocrine Society, 2020, 4, PRKAR1A deficiency delays postnatal heart growth. FASEB Journal, 2020, 34, 1-1 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	hildren and adolescents. Journal of Cancer Research and Clinical Oncology, 2020, 146, 1051-1063 Medical Treatment of Pituitary Adenomas: A Celebration of Endocrinology (and Oncology)). Hormone and Metabolic Research, 2020, 52, 7 Cushing syndrome: Old and new genes. Best Practice and Research in Clinical Endocrinology and Metabolism, 2020, 34, 101418 Loss of habenular Prkar2a reduces hedonic eating and increases exercise motivation. JCI Insight, 2020, 5, 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, Liver findings in patients with Carney complex, germline PRKAR1A pathogenic variants, and link to cardiac myxomas. Endocrine-Related Cancer, 2020, 27, 355-360 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, 177-186 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, 187-197 Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. Endocrine-Related Cancer, 2020, 27, 403-413 ARMCS variants in PRKAR1A-mutated patients modify cortisol levels and Cushing's syndrome. Endocrine-Related Cancer, 2020, 27, 509-517 PRKACB variants in skeletal disease or adrenocortical hyperplasia: effects on protein kinase A. Endocrine-Related Cancer, 2020, 27, 647-656 PKA functions in metabolism and resistance to obesity: lessons from mouse and human studies. Journal of Endocrinology, 2020, 246, R51-R64 OR23-01 Intrapatient ACTH Variability in Cushing® Disease: Prognostic Significance. Journal of the Endocrine Society, 2020, 4, PRKAR1A deficiency delays postnatal heart growth. FASEB Journal, 2020, 34, 1-1 OGO-17 The Role of Germline Defects in Cushing® Disease. Jour	hildren and adolescents. Journal of Cancer Research and Clinical Oncology, 2020, 146, 1051-1063 49 14 Medical Treatment of Pituitary Adenomas: A Celebration of Endocrinology (and Oncology)!. Hormone and Metabolic Research, 2020, 52, 7 Cushing syndrome: Old and new genes. Best Practice and Research in Clinical Endocrinology and Metabolism, 2020, 34, 101418 Loss of habenular Prkar2a reduces hedonic eating and increases exercise motivation. JCl Insight, 2020, 34, 101418 Loss of habenular Prkar2a reduces hedonic eating and increases exercise motivation. JCl Insight, 2020, 34, 101418 The Association of ARMCS with the Renin-Angiotensin-Aldosterone System, Blood Pressure, and Clycemia in African Americans. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 56 5 Liver findings in patients with Carney complex, germline PRKAR1A pathogenic variants, and link to cardiac mysomas. Endocrine-Related Cancer, 2020, 27, 353-360 HEREDITARY ENDOCRINE TUMOURS: CURRENT STATE-OF-THE-ART AND RESEARCH OPPORTUNITIES. The roles of AlP and GPR101 in familial Isolated pituitary adenomas (FIPA). 57 7 HEREDITARY ENDOCRINE TUMOURS: CURRENT STATE-OF-THE-ART AND RESEARCH OPPORTUNITIES GPR 101, an orphan GPCR with roles in growth and pituitary tumorigenesis. 57 7 Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. Endocrine-Related Cancer, 2020, 27, 187-197 Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. Endocrine-Related Cancer, 2020, 27, 509-517 PRKACB variants in PRKAR1A-mutated patients modify cortisol levels and Cushing's syndrome. 57 4 PRAGet variants in metabolism and resistance to obesity; lessons from mouse and human studies. 57 4 PRAGet variants in metabolism and resistance to obesity; lessons from mouse and human studies. 57 4 PRKACB variants in metabolism and resistance to obesity; lessons from mouse and human studies. 57 4 PRAGet variants in metabolism and resistance to obesity; lessons from mouse and

c-KIT oncogene expression in PRKAR1A-mutant adrenal cortex. *Endocrine-Related Cancer*, **2020**, 27, 591-599

289	c-KIT oncogene expression in PRKAR1A-mutant adrenal cortex. <i>Endocrine-Related Cancer</i> , 2020 , 27, 59	I- §9/9	
288	Acute Statin Administration Reduces Levels of Steroid Hormone Precursors. <i>Hormone and Metabolic Research</i> , 2020 , 52, 742-746	3.1	
287	SUN-235 Deficient Fear Extinction in PRKAR1A-Defective Mice. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78
286	SAT-543 Human Hair Aldosterone Measurements for Evaluation of Primary Aldosteronism. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78
285	OR24-06 USP8 Genetic Variants May Contribute to the Development of Bilateral Adrenal Hyperplasia and ACTH-Independent Cushing Syndrome. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78
284	SUN-713 Prevalence of Renal Cysts in Patients with Carney Complex. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78
283	Telomere Length Changes in Children With Cushing Disease: A Pilot Study. <i>Journal of the Endocrine Society</i> , 2020 , 4, bvaa067	0.4	O
282	Aggressive pituitary tumors in the young and elderly. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2020 , 21, 213-223	10.5	10
281	Computerized Analysis of Brain MRI Parameter Dynamics in Young Patients With Cushing Syndrome-A Case-Control Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	4
280	Carney complex syndrome manifesting as cardioembolic stroke: a case report and review of the literature. <i>International Journal of Neuroscience</i> , 2020 , 1-7	2	1
279	Kisspeptin deficiency leads to abnormal adrenal glands and excess steroid hormone secretion. <i>Human Molecular Genetics</i> , 2020 , 29, 3443-3450	5.6	1
278	Prevalence of Diabetes and Hypertension and Their Associated Risks for Poor Outcomes in Covid-19 Patients. <i>Journal of the Endocrine Society</i> , 2020 , 4, bvaa102	0.4	32
277	Rare Germline Variants in Pediatric Patients With Cushing's Disease: What Is Their Role?. <i>Frontiers in Endocrinology</i> , 2020 , 11, 433	5.7	4
276	Hemodynamics of Prefrontal Cortex in Ornithine Transcarbamylase Deficiency: A Twin Case Study. <i>Frontiers in Neurology</i> , 2020 , 11, 809	4.1	2
275	Prkar1a haploinsufficiency ameliorates the growth hormone excess phenotype in Aip-deficient mice. <i>Human Molecular Genetics</i> , 2020 , 29, 2951-2961	5.6	1
274	A Century After the Description of "Hormones", Our Golden Jubilee Celebration Goes on with What is New in Endocrine Oncology: And a Lot is New!. <i>Hormone and Metabolic Research</i> , 2020 , 52, 551-552	3.1	
273	ARMC5 Alterations in Patients With Sporadic Neuroendocrine Tumors and Multiple Endocrine Neoplasia Type 1 (MEN1). <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	2

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272	Letter to the Editor from Berthon: "Cardiac Myxoma Caused by Fumarate Hydratase Gene Deletion in Patient With Cortisol-Secreting Adrenocortical Adenoma". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	1	
271	Chaperones, somatotroph tumors and the cyclic AMP (cAMP)-dependent protein kinase (PKA) pathway. <i>Molecular and Cellular Endocrinology</i> , 2020 , 499, 110607	4.4	5	
270	Mosaicism for KCNJ5 Causing Early-Onset Primary Aldosteronism due to Bilateral Adrenocortical Hyperplasia. <i>American Journal of Hypertension</i> , 2020 , 33, 124-130	2.3	12	
269	Letter to the Editor: "IGSF1 Deficiency Results in Human and Murine Somatotrope Neurosecretory Hyperfunction". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6		
268	Germline USP8 Mutation Associated With Pediatric Cushing Disease and Other Clinical Features: A New Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 4676-4682	5.6	28	
267	Subspecialty training in adult inherited metabolic diseases: a call to action for unmet needs. <i>Lancet Diabetes and Endocrinology,the</i> , 2019 , 7, 82-84	18.1	2	
266	The Catalytic Subunit of PKA Affects Energy Balance and Catecholaminergic Activity. <i>Journal of the Endocrine Society</i> , 2019 , 3, 1062-1078	0.4	7	
265	Called and Uncalled for Functions of the Main Catalytic Subunit of Protein Kinase A: One Enzyme, Many Faces. <i>Endocrinology</i> , 2019 , 160, 1674-1676	4.8	2	
264	CRH stimulation improves F-FDG-PET detection of pituitary adenomas in Cushing's disease. <i>Endocrine</i> , 2019 , 65, 155-165	4	13	
263	Inflammation and Metabolism in Cancer Cell-Mitochondria Key Player. <i>Frontiers in Oncology</i> , 2019 , 9, 348	5.3	60	
262	Large Genomic Aberrations in Corticotropinomas Are Associated With Greater Aggressiveness. Journal of Clinical Endocrinology and Metabolism, 2019 , 104, 1792-1801	5.6	10	
261	High expression of adrenal P450 aromatase (CYP19A1) in association with ARMC5-primary bilateral macronodular adrenocortical hyperplasia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019 , 191, 105316	5.1	5	
260	Clinical, Diagnostic, and Treatment Characteristics of -Related Metastatic Pheochromocytoma and Paraganglioma. <i>Frontiers in Oncology</i> , 2019 , 9, 53	5.3	24	
259	SGPL1 Deficiency: A Rare Cause of Primary Adrenal Insufficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 1484-1490	5.6	13	
258	Genetics of Hypertension in African Americans and Others of African Descent. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	23	
257	CD40LG duplications in patients with X-LAG syndrome commonly undergo random X-chromosome inactivation. <i>Journal of Allergy and Clinical Immunology</i> , 2019 , 143, 1659	11.5	3	
256	Molecular mechanisms of mutations in adrenal pathophysiology. <i>Current Opinion in Endocrine and Metabolic Research</i> , 2019 , 8, 104-111	1.7	8	
255	Pheochromocytomas: Fabulous, Fascinating, and First (in everything)!. <i>Hormone and Metabolic Research</i> , 2019 , 51, 401-402	3.1	2	

254	Somatic Gene Mutation in a Nonsyndromic Metastatic Large Cell Calcifying Sertoli Cell Tumor. Journal of the Endocrine Society, 2019 , 3, 1375-1382	0.4	2
253	Cushing disease in a patient with nonbullous congenital ichthyosiform erythroderma: lessons in avoiding glucocorticoids in ichthyosis. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2019 , 32, 911-	9 ⁴ 14	
252	Multiple Endocrine Neoplasia Type 1 (MEN1): An Update and the Significance of Early Genetic and Clinical Diagnosis. <i>Frontiers in Endocrinology</i> , 2019 , 10, 339	5.7	61
251	ARMC 5 Variants and Risk of Hypertension in Blacks: MH- GRID Study. <i>Journal of the American Heart Association</i> , 2019 , 8, e012508	6	5
250	Resistant Hypertension: A Clinical Perspective. <i>Endocrinology and Metabolism Clinics of North America</i> , 2019 , 48, 811-828	5.5	2
249	Illicit Upregulation of Serotonin Signaling Pathway in Adrenals of Patients With High Plasma or Intra-Adrenal ACTH Levels. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 4967-4980	5.6	9
248	The Genetics of Pituitary Adenomas. <i>Journal of Clinical Medicine</i> , 2019 , 9,	5.1	19
247	Genetic Characteristics of Aldosterone-Producing Adenomas in Blacks. <i>Hypertension</i> , 2019 , 73, 885-892	8.5	78
246	A novel mutation in the glucocorticoid receptor gene as a cause of severe glucocorticoid resistance complicated by hypertensive encephalopathy. <i>Journal of Hypertension</i> , 2019 , 37, 1475-1481	1.9	4
245	Variations in maternal adenylate cyclase genes are associated with congenital Zika syndrome in a cohort from Northeast, Brazil. <i>Journal of Internal Medicine</i> , 2019 , 285, 215-222	10.8	12
244	Carney Complex. Experimental and Clinical Endocrinology and Diabetes, 2019, 127, 156-164	2.3	51
243	Optical Imaging Technology: A Useful Tool to Identify Remission in Cushing Disease After Surgery. <i>Hormone and Metabolic Research</i> , 2019 , 51, 120-126	3.1	1
242	Growth hormone excess in neurofibromatosis 1. <i>Genetics in Medicine</i> , 2019 , 21, 1254-1255	8.1	8
241	The 3PAs: An Update on the Association of Pheochromocytomas, Paragangliomas, and Pituitary Tumors. <i>Hormone and Metabolic Research</i> , 2019 , 51, 419-436	3.1	14
240	Genetic Tumor Syndromes with Endocrine Involvement: A Compendium and an Update. <i>Pediatric Endocrinology Reviews</i> , 2019 , 16, 311-334	1.1	
239	Pediatric Cushing Syndrome; an Overview. <i>Pediatric Endocrinology Reviews</i> , 2019 , 17, 100-109	1.1	4
238	Incidence of Autoimmune and Related Disorders After Resolution of Endogenous Cushing Syndrome in Children. <i>Hormone and Metabolic Research</i> , 2018 , 50, 290-295	3.1	7
237	Anxiety-like behavior and other consequences of early life stress in mice with increased protein kinase A activity. <i>Behavioural Brain Research</i> , 2018 , 348, 22-30	3.4	2

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Mutational Analysis of the PRL Receptor Gene in Human Breast Tumors with Differential PRL Receptor Protein Expression

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Genetics of Carney Complex and Related Familial Lentiginoses, and other Multiple Tumor Syndromes

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