Philippe Chanson

List of Publications by Year
in descending order

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280 papers	19,683 citations	6592 79 h-index	14702 127 g-index
298 all docs	298 docs citations	298 times ranked	11844 citing authors

#	Article	IF	Citations
1	A Consensus on Criteria for Cure of Acromegaly. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3141-3148.	1.8	697
2	A Family with Hypogonadotropic Hypogonadism and Mutations in the Gonadotropin-Releasing Hormone Receptor. New England Journal of Medicine, 1997, 337, 1597-1603.	13.9	473
3	A Consensus Statement on acromegaly therapeutic outcomes. Nature Reviews Endocrinology, 2018, 14, 552-561.	4.3	382
4	GNAS-activating mutations define a rare subgroup of inflammatory liver tumors characterized by STAT3 activation. Journal of Hepatology, 2012, 56, 184-191.	1.8	354
5	Clinical Characteristics and Therapeutic Responses in Patients with Germ-LineAIPMutations and Pituitary Adenomas: An International Collaborative Study. Journal of Clinical Endocrinology and Metabolism, 2010, 95, E373-E383.	1.8	323
6	The European Registry on Cushing's syndrome: 2-year experience. Baseline demographic and clinical characteristics. European Journal of Endocrinology, 2011, 165, 383-392.	1.9	322
7	Impact of Growth Hormone (GH) Treatment on Cardiovascular Risk Factors in GH-Deficient Adults: A Metaanalysis of Blinded, Randomized, Placebo-Controlled Trials. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 2192-2199.	1.8	321
8	Consensus on diagnosis and management of Cushing's disease: a guideline update. Lancet Diabetes and Endocrinology,the, 2021, 9, 847-875.	5.5	315
9	Expert consensus document: A consensus on the medical treatment of acromegaly. Nature Reviews Endocrinology, 2014, 10, 243-248.	4.3	306
10	Gigantism and Acromegaly Due to Xq26 Microduplications and GPR101 Mutation. New England Journal of Medicine, 2014, 371, 2363-2374.	13.9	292
11	MAX Mutations Cause Hereditary and Sporadic Pheochromocytoma and Paraganglioma. Clinical Cancer Research, 2012, 18, 2828-2837.	3.2	277
12	Pituitary Apoplexy. Endocrine Reviews, 2015, 36, 622-645.	8.9	270
13	Place of Cabergoline in Acromegaly: A Meta-Analysis. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 1327-1335.	1.8	255
14	Isolated Familial Hypogonadotropic Hypogonadism and aGNRH1Mutation. New England Journal of Medicine, 2009, 360, 2742-2748.	13.9	247
15	Acromegaly. Nature Reviews Disease Primers, 2019, 5, 20.	18.1	247
16	Ketoconazole in Cushing's Disease: Is It Worth a Try?. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1623-1630.	1.8	231
17	TAC3andTACR3Defects Cause Hypothalamic Congenital Hypogonadotropic Hypogonadism in Humans. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 2287-2295.	1.8	214
18	Outcome of Gamma Knife Radiosurgery in 82 Patients with Acromegaly: Correlation with Initial Hypersecretion. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4483-4488.	1.8	209

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19	Endocrine Aspects of Obstructive Sleep Apnea. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 483-495.	1.8	202
20	Temozolomide Treatment in Aggressive Pituitary Tumors and Pituitary Carcinomas: A French Multicenter Experience. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 4592-4599.	1.8	202
21	Germ-Line Mutation Analysis in Patients with Multiple Endocrine Neoplasia Type 1 and Related Disorders. American Journal of Human Genetics, 1998, 63, 455-467.	2.6	197
22	Treatment of aggressive pituitary tumours and carcinomas: results of a European Society of Endocrinology (ESE) survey 2016. European Journal of Endocrinology, 2018, 178, 265-276.	1.9	196
23	A Critical Analysis of Pituitary Tumor Shrinkage during Primary Medical Therapy in Acromegaly. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4405-4410.	1.8	193
24	Effects of Somatostatin Analogs on Glucose Homeostasis: A Metaanalysis of Acromegaly Studies. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 1500-1508.	1.8	191
25	Octreotide Therapy for Thyroid-Stimulating Hormone-Secreting Pituitary Adenomas: A Follow-up of 52 Patients. Annals of Internal Medicine, 1993, 119, 236.	2.0	189
26	Acromegaly. Orphanet Journal of Rare Diseases, 2008, 3, 17.	1.2	188
27	Mitotane, Metyrapone, and Ketoconazole Combination Therapy as an Alternative to Rescue Adrenalectomy for Severe ACTH-Dependent Cushing's Syndrome. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 2796-2804.	1.8	187
28	Multidisciplinary management of acromegaly: A consensus. Reviews in Endocrine and Metabolic Disorders, 2020, 21, 667-678.	2.6	183
29	New insights in prolactin: pathological implications. Nature Reviews Endocrinology, 2015, 11, 265-275.	4.3	178
30	Prolactinomas resistant to standard doses of cabergoline: a multicenter study of 92 patients. European Journal of Endocrinology, 2012, 167, 651-662.	1.9	173
31	Growth Hormone, Insulin-Like Growth Factor-1, and the Kidney: Pathophysiological and Clinical Implications. Endocrine Reviews, 2014, 35, 234-281.	8.9	171
32	Factors predicting relapse of nonfunctioning pituitary macroadenomas after neurosurgery: a study of 142 patients. European Journal of Endocrinology, 2010, 163, 193-200.	1.9	167
33	Long-Term Results of Stereotactic Radiosurgery in Secretory Pituitary Adenomas. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 3400-3407.	1.8	164
34	Acromegaly at diagnosis in 3173 patients from the LiÃ¨ge Acromegaly Survey (LAS) Database. Endocrine-Related Cancer, 2017, 24, 505-518.	1.6	164
35	RecurrentPRKAR1AMutation in Acrodysostosis with Hormone Resistance. New England Journal of Medicine, 2011, 364, 2218-2226.	13.9	162
36	Germline AIP Mutations in Apparently Sporadic Pituitary Adenomas: Prevalence in a Prospective Single-Center Cohort of 443 Patients. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E663-E670.	1.8	157

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37	Cardiac Effects of Growth Hormone in Adults With Growth Hormone Deficiency. Circulation, 2003, 108, 2648-2652.	1.6	155
38	Clinical and genetic characterization of pituitary gigantism: an international collaborative study in 208 patients. Endocrine-Related Cancer, 2015, 22, 745-757.	1.6	155
39	Acromegaly and McCune-Albright Syndrome. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1955-1969.	1.8	149
40	Gross total resection or debulking of pituitary adenomas improves hormonal control of acromegaly by somatostatin analogs. European Journal of Endocrinology, 2005, 152, 61-66.	1.9	148
41	Pituitary tumour transforming gene (PTTG) expression correlates with the proliferative activity and recurrence status of pituitary adenomas: a clinical and immunohistochemical study. Clinical Endocrinology, 2006, 65, 536-543.	1.2	142
42	Diabetes in acromegaly, prevalence, risk factors, and evolution: data from the French Acromegaly Registry. European Journal of Endocrinology, 2011, 164, 877-884.	1.9	140
43	Management of clinically non-functioning pituitary adenoma. Annales D'Endocrinologie, 2015, 76, 239-247.	0.6	136
44	Cinacalcet Reduces Serum Calcium Concentrations in Patients with Intractable Primary Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 2766-2772.	1.8	134
45	Impact of Somatostatin Analogs on the Heart in Acromegaly: A Metaanalysis. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 1743-1747.	1.8	133
46	Acromegaly. Best Practice and Research in Clinical Endocrinology and Metabolism, 2009, 23, 555-574.	2.2	133
47	Changes in the management and comorbidities of acromegaly over three decades: the French Acromegaly Registry. European Journal of Endocrinology, 2017, 176, 645-655.	1.9	133
48	The diagnostic value of fine-needle aspiration biopsy under ultrasonography in nonfunctional thyroid nodules: A prospective study comparing cytologic and histologic findings. American Journal of Medicine, 1994, 97, 152-157.	0.6	128
49	Germline inactivating mutations of the aryl hydrocarbon receptor-interacting protein gene in a large cohort of sporadic acromegaly: mutations are found in a subset of young patients with macroadenomas. European Journal of Endocrinology, 2007, 157, 1-8.	1.9	127
50	Cardiovascular Effects of the Somatostatin Analog Octreotide in Acromegaly. Annals of Internal Medicine, 1990, 113, 921.	2.0	125
51	Testicular Anti-MÃ¼llerian Hormone Secretion Is Stimulated by Recombinant Human FSH in Patients with Congenital Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 724-728.	1.8	122
52	Prevalence and risk factors of impaired glucose tolerance and diabetes mellitus at diagnosis of acromegaly: a study in 148 patients. Pituitary, 2014, 17, 81-89.	1.6	122
53	Pregnancy outcomes following cabergoline treatment: extended results from a 12â€year observational study. Clinical Endocrinology, 2008, 68, 66-71.	1.2	120
54	Reference Values for IGF-I Serum Concentrations: Comparison of Six Immunoassays. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 3450-3458.	1.8	118

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55	Panhypopituitarism as a Model to Study the Metabolism of Dehydroepiandrosterone (DHEA) in Humans¹. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 2578-2585.	1.8	116
56	APRKAR1AMutation Associated with Primary Pigmented Nodular Adrenocortical Disease in 12 Kindreds. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 1943-1949.	1.8	116
57	The epidemiology, diagnosis and treatment of Prolactinomas: The old and the new. Best Practice and Research in Clinical Endocrinology and Metabolism, 2019, 33, 101290.	2.2	115
58	THERAPY OF ENDOCRINE DISEASE: Outcomes in patients with Cushing's disease undergoing transsphenoidal surgery: systematic review assessing criteria used to define remission and recurrence. European Journal of Endocrinology, 2015, 172, R227-R239.	1.9	114
59	Prevalence of Metabolic Syndrome in Adult Hypopituitary Growth Hormone (GH)-Deficient Patients Before and After GH Replacement. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 74-81.	1.8	113
60	Efficacy of the Long-Acting Octreotide Formulation (Octreotide-Lar) in Patients with Thyrotropin-Secreting Pituitary Adenomas. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 2849-2853.	1.8	112
61	Acromegaly and Pregnancy: A Retrospective Multicenter Study of 59 Pregnancies in 46 Women. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 4680-4687.	1.8	111
62	Clinical Pharmacokinetics of Octreotide. Clinical Pharmacokinetics, 1993, 25, 375-391.	1.6	110
63	Kallmannâ€™s Syndrome: A Comparison of the Reproductive Phenotypes in Men Carrying KAL1 and FGFR1/KAL2 Mutations. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 758-763.	1.8	109
64	Frequent Large GermlineHRPT2Deletions in a French National Cohort of Patients With Primary Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2013, 98, E403-E408.	1.8	107
65	Temozolomide treatment can improve overall survival in aggressive pituitary tumors and pituitary carcinomas. European Journal of Endocrinology, 2017, 176, 769-777.	1.9	107
66	Non-syndromic congenital hypogonadotropic hypogonadism: clinical presentation and genotypeâ€“phenotype relationships. European Journal of Endocrinology, 2010, 162, 835-851.	1.9	104
67	Normal Pituitary Hypertrophy as a Frequent Cause of Pituitary Incidentaloma: A Follow-Up Study. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 3009-3015.	1.8	101
68	Metabolic Syndrome in Cushingâ€™s Syndrome. Neuroendocrinology, 2010, 92, 96-101.	1.2	99
69	Pituitary Magnetic Resonance Imaging Findings Do Not Influence Surgical Outcome in Adrenocorticotropin-Secreting Microadenomas. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 3371-3376.	1.8	94
70	Functional hypothalamic amenorrhoea: a partial and reversible gonadotrophin deficiency of nutritional origin. Clinical Endocrinology, 1999, 50, 229-235.	1.2	92
71	McCuneâ€“Albright syndrome and acromegaly: clinical studies and responses to treatment in five cases. European Journal of Endocrinology, 1994, 131, 229-234.	1.9	90
72	Long-Term Outcome of Patients with Acromegaly and Congestive Heart Failure. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 5308-5313.	1.8	89

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73	Current management practices for acromegaly: an international survey. Pituitary, 2011, 14, 125-133.	1.6	89
74	MAFA missense mutation causes familial insulinomatosis and diabetes mellitus. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 1027-1032.	3.3	88
75	Epithelial Sodium Channel Is a Key Mediator of Growth Hormone-Induced Sodium Retention in Acromegaly. Endocrinology, 2008, 149, 3294-3305.	1.4	86
76	Genetic mutations in sporadic pituitary adenomasâ€"what to screen for?. Nature Reviews Endocrinology, 2015, 11, 43-54.	4.3	86
77	Panhypopituitarism as a Model to Study the Metabolism of Dehydroepiandrosterone (DHEA) in Humans. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 2578-2585.	1.8	86
78	Rapid control of severe neoplastic hypercortisolism with metyrapone and ketoconazole. European Journal of Endocrinology, 2015, 172, 473-481.	1.9	84
79	Macroprolactinomas in Children and Adolescents: Factors Associated With the Response to Treatment in 77 Patients. Journal of Clinical Endocrinology and Metabolism, 2015, 100, 1177-1186.	1.8	83
80	Normosmic Congenital Hypogonadotropic Hypogonadism Due to TAC3/TACR3 Mutations: Characterization of Neuroendocrine Phenotypes and Novel Mutations. PLoS ONE, 2011, 6, e25614.	1.1	83
81	Differential gene expression profiles of invasive and non-invasive non-functioning pituitary adenomas based on microarray analysis. Endocrine-Related Cancer, 2010, 17, 361-371.	1.6	81
82	Clinical Characteristics and Outcome of Acromegaly Induced by Ectopic Secretion of Growth Hormone-Releasing Hormone (GHRH): A French Nationwide Series of 21 Cases. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 2093-2104.	1.8	81
83	Higher risk of death among MEN1 patients with mutations in the JunD interacting domain: a Groupe dâ€™Ã©tude des Tumeurs Endocrines (GTE) cohort study. Human Molecular Genetics, 2013, 22, 1940-1948.	1.4	81
84	Osteoblastic cells derived fronm isolated lesions of fibrous dysplasia contain activating somatic mutatuions of thje Gsî± gene. Human Molecular Genetics, 1995, 4, 1675-1676.	1.4	78
85	Pituitary MRI characteristics in 297 acromegaly patients based on T2-weighted sequences. Endocrine-Related Cancer, 2015, 22, 169-177.	1.6	78
86	AntimÃ¼llerian Hormone in Patients with Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 2696-2699.	1.8	77
87	Female Gonadal Function before and after Treatment of Acromegaly. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 4518-4525.	1.8	77
88	Type A Insulin Resistance Syndrome Revealing a Novel Lamin A Mutation. Diabetes, 2005, 54, 1873-1878.	0.3	75
89	Hepatobiliary and Pancreatic Neoplasms in Patients With McCune-Albright Syndrome. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E97-E101.	1.8	75
90	Effects of Human Recombinant Luteinizing Hormone and Follicle-Stimulating Hormone in Patients with Acquired Hypogonadotropic Hypogonadism: Study of Sertoli and Leydig Cell Secretions and Interactions. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 3239-3244.	1.8	72

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91	Macimorelin as a Diagnostic Test for Adult GH Deficiency. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 3083-3093.	1.8	71
92	Growth hormone as a risk for premature mortality in healthy subjects: data from the Paris prospective study. BMJ: British Medical Journal, 1998, 316, 1132-1133.	2.4	70
93	Vascular reactivity in acromegalic patients: preliminary evidence for regional endothelial dysfunction and increased sympathetic vasoconstriction. Clinical Endocrinology, 2000, 53, 445-451.	1.2	70
94	Control of IGFâ€Ł levels with titrated dosing of lanreotide Autogel over 48Âweeks in patients with acromegaly. Clinical Endocrinology, 2008, 69, 299-305.	1.2	70
95	Primary hyperparathyroidism in pregnancy. Endocrine, 2013, 44, 591-597.	1.1	65
96	Cardiac Structure and Function in Cushing's Syndrome: A Cardiac Magnetic Resonance Imaging Study. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E2144-E2153.	1.8	65
97	Post-surgical management of non-functioning pituitary adenoma. Annales D'Endocrinologie, 2015, 76, 228-238.	0.6	65
98	National acromegaly registries. Best Practice and Research in Clinical Endocrinology and Metabolism, 2019, 33, 101264.	2.2	65
99	Clinical aspects of multiple endocrine neoplasia type 1. Nature Reviews Endocrinology, 2021, 17, 207-224.	4.3	64
100	Antimullerian Hormone in Patients with Hypogonadotropic Hypogonadism. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 2696-2699.	1.8	64
101	Cardiac Effects of Growth Hormone Treatment in Chronic Heart Failure: A Meta-Analysis. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 180-185.	1.8	63
102	Non-invasive Diagnostic Strategy in ACTH-dependent Cushingâ€™s Syndrome. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 3273-3284.	1.8	62
103	Biochemical characterization of a Ca2+/NAD(P)H-dependent H2O2 generator in human thyroid tissue. Biochimie, 1999, 81, 373-380.	1.3	61
104	Parental Origin of GsÎ± Mutations in the McCune-Albright Syndrome and in Isolated Endocrine Tumors. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 3007-3009.	1.8	61
105	Comparison of fast Fourier transform and autoregressive spectral analysis for the study of heart rate variability in diabetic patients. International Journal of Cardiology, 2005, 104, 307-313.	0.8	60
106	Prevalence and Incidence of Diabetes Mellitus in Adult Patients on Growth Hormone Replacement for Growth Hormone Deficiency: A Surveillance Database Analysis. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 2255-2261.	1.8	60
107	The Antigonadotropic Activity of a 19-Nor-Progesterone Derivative Is Exerted Both at the Hypothalamic and Pituitary Levels in Women. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 4191-4196.	1.8	57
108	No Evidence of a Detrimental Effect of Cabergoline Therapy on Cardiac Valves in Patients with Acromegaly. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E1714-E1719.	1.8	57

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109	SAGITÂ®: clinician-reported outcome instrument for managing acromegaly in clinical practiceâ€"development and results from a pilot study. Pituitary, 2016, 19, 39-49.	1.6	56
110	Anti-MÃ¼llerian Hormone and Ovarian Morphology in Women With Isolated Hypogonadotropic Hypogonadism/Kallmann Syndrome: Effects of Recombinant Human FSH. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 1102-1111.	1.8	55
111	McCune-Albright Syndrome and Acromegaly: Effects of Hypothalamopituitary Radiotherapy and/or Pegvisomant in Somatostatin Analog-Resistant Patients. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 4957-4961.	1.8	54
112	Endocrine Effects of the Tyrosine Kinase Inhibitor Vandetanib in Patients Treated for Thyroid Cancer. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 2741-2749.	1.8	54
113	Management of hyperglycaemia in Cushing's disease: Expertsâ€™ proposals on the use of pasireotide. Diabetes and Metabolism, 2013, 39, 34-41.	1.4	54
114	Prognostic markers of survival after combined mitotane- and platinum-based chemotherapy in metastatic adrenocortical carcinoma. Endocrine-Related Cancer, 2010, 17, 797-807.	1.6	52
115	Signs and symptoms of acromegaly at diagnosis: the physicianâ€™s and the patientâ€™s perspectives in the ACRO-POLIS study. Endocrine, 2019, 63, 120-129.	1.1	51
116	A new FSHÎ² mutation in a 29-year-old woman with primary amenorrhea and isolated FSH deficiency: functional characterization and ovarian response to human recombinant FSH. European Journal of Endocrinology, 2010, 162, 633-641.	1.9	50
117	High-dose mitotane strategy in adrenocortical carcinoma: prospective analysis of plasma mitotane measurement during the first 3 months of follow-up. European Journal of Endocrinology, 2012, 166, 261-268.	1.9	50
118	Body Fluid Expansion in Acromegaly Is Related to Enhanced Epithelial Sodium Channel (ENaC) Activity. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 2127-2135.	1.8	49
119	Pathophysiology of Renal Calcium Handling in Acromegaly: What Lies behind Hypercalciuria?. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 2124-2133.	1.8	48
120	Management of nonfunctioning pituitary incidentaloma. Annales D'Endocrinologie, 2015, 76, 191-200.	0.6	48
121	Cabergoline in acromegaly. Pituitary, 2017, 20, 121-128.	1.6	48
122	Pitfall of Petrosal Sinus Sampling in a Cushing's Syndrome Secondary to Ectopic Adrenocorticotropin-Corticotropin Releasing Hormone (ACTH-CRH) Secretion. Journal of Clinical Endocrinology and Metabolism, 1998, 83, 305-308.	1.8	48
123	Prevalence ofKISS1 Receptormutations in a series of 603 patients with normosmic congenital hypogonadotrophic hypogonadism and characterization of novel mutations: a single-centre study. Human Reproduction, 2016, 31, 1363-1374.	0.4	47
124	The effect of somatostatin analogue on chiasmal dysfunction from pituitary macroadenomas. Journal of Neurosurgery, 1989, 71, 687-690.	0.9	46
125	Management of pituitary apoplexy. Expert Opinion on Pharmacotherapy, 2004, 5, 1287-1298.	0.9	46
126	Integrated Genomic Profiling Identifies Loss of Chromosome 11p Impacting Transcriptomic Activity in Aggressive Pituitary PRL Tumors. Brain Pathology, 2011, 21, 533-543.	2.1	46

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127	Insulin-like Peptide 3 (INSL3) in Men With Congenital Hypogonadotropic Hypogonadism/Kallmann Syndrome and Effects of Different Modalities of Hormonal Treatment: A Single-Center Study of 281 Patients. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E268-E275.	1.8	46
128	Impact of Successful Treatment of Acromegaly on Overnight Heart Rate Variability and Sleep Apnea. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 2925-2931.	1.8	46
129	Pituitary Apoplexy. Endocrinology and Metabolism Clinics of North America, 2015, 44, 199-209.	1.2	46
130	Decreased regional blood flow in patients with acromegaly. Clinical Endocrinology, 1998, 49, 725-731.	1.2	45
131	Long-term results of the surgical management of insulinoma patients with MEN1: a Groupe d'Ã©tude des Tumeurs Endocrines (GTE) retrospective study. European Journal of Endocrinology, 2015, 172, 309-319.	1.9	44
132	Long-term effects of pegvisomant on comorbidities in patients with acromegaly: a retrospective single-center study. European Journal of Endocrinology, 2015, 173, 693-702.	1.9	44
133	Very low frequency of germline GPR101 genetic variation and no biallelic defects with AIP in a large cohort of patients with sporadic pituitary adenomas. European Journal of Endocrinology, 2016, 174, 523-530.	1.9	44
134	Metastatic Potential and Survival of Duodenal and Pancreatic Tumors in Multiple Endocrine Neoplasia Type 1. Annals of Surgery, 2020, 272, 1094-1101.	2.1	44
135	RAPID IMPROVEMENT IN SLEEP APNOEA OF ACROMEGALY AFTER SHORT-TERM TREATMENT WITH SOMATOSTATIN ANALOGUE SMS 201-995. Lancet, The, 1986, 327, 1270-1271.	6.3	43
136	Genomic Alterations and Complex Subclonal Architecture in Sporadic GH-Secreting Pituitary Adenomas. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 1929-1939.	1.8	43
137	Efficacy and tolerability of the long-acting somatostatin analog lanreotide in acromegaly. A 12-month multicenter study of 58 acromegalic patients. French Multicenter Study Group on Lanreotide in Acromegaly. Pituitary, 2000, 2, 269-276.	1.6	42
138	Diagnostic tests for Cushing's syndrome differ from published guidelines: data from ERCUSYN. European Journal of Endocrinology, 2017, 176, 613-624.	1.9	42
139	Rapidly Reversible Myocardial Edema in Patients with Acromegaly: Assessment with Ultrafast T2 Mapping in a Single-Breath-Hold MRI Sequence. American Journal of Roentgenology, 2008, 190, 1576-1582.	1.0	40
140	Acromegaly. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2014, 124, 197-219.	1.0	40
141	Worse Healthâ€Related Quality of Life at longâ€term followâ€up in patients with Cushing's disease than patients with cortisol producing adenoma. Data from the <scp>ERCUSYN</scp>. Clinical Endocrinology, 2018, 88, 787-798.	1.2	40
142	Sex-Related Differences in Lactotroph Tumor Aggressiveness Are Associated With a Specific Gene-Expression Signature and Genome Instability. Frontiers in Endocrinology, 2018, 9, 706.	1.5	40
143	Cardiovascular findings and management in Turner syndrome: insights from a French cohort. European Journal of Endocrinology, 2012, 167, 517-522.	1.9	39
144	Hypothalamic-Pituitary-Ovarian Axis Reactivation by Kisspeptin-10 in Hyperprolactinemic Women With Chronic Amenorrhea. Journal of the Endocrine Society, 2017, 1, 1362-1371.	0.1	38

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145	Adrenal GIPR expression and chromosome 19q13 microduplications in GIP-dependent Cushingâ€™s syndrome. JCI Insight, 2017, 2, .	2.3	38
146	Preoperative medical treatment in Cushingâ€™s syndrome: frequency of use and its impact on postoperative assessment: data from ERCUSYN. European Journal of Endocrinology, 2018, 178, 399-409.	1.9	37
147	The 2016â€“2019 ImmunoTOX assessment board report of collaborative management of immune-related adverse events, an observational clinical study. European Journal of Cancer, 2020, 130, 39-50.	1.3	37
148	The effect of subcutaneous infusion versus subcutaneous injections of a somatostatin analogue (SMS) Tj ETQq0 0 0 rgBT /Overlock 10 T	1.9	36
149	68Ga-Exendin-4 PET/CT Detects Insulinomas in Patients With Endogenous Hyperinsulinemic Hypoglycemia in MEN-1. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5843-5852.	1.8	36
150	Non-responsiveness of serum gonadotropins and testosterone to pulsatile GnRH in hemochromatosis suggesting a pituitary defect. European Journal of Endocrinology, 1993, 128, 351-354.	1.9	35
151	Congenital hypogonadotropic hypogonadism in females: Clinical spectrum, evaluation and genetics. Annales D'Endocrinologie, 2010, 71, 158-162.	0.6	34
152	Growth Hormone Response to Oral Glucose Load: From Normal to Pathological Conditions. Neuroendocrinology, 2019, 108, 244-255.	1.2	34
153	How can we minimise the use of regular oral corticosteroids in asthma?. European Respiratory Review, 2020, 29, 190085.	3.0	34
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155	Severe hyponatremia as a frequent revealing sign of hypopituitarism after 60 years of age. European Journal of Endocrinology, 2003, 149, 177-178.	1.9	33
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