Pr Gerald Raverot

List of Publications by Year in descending order

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152 papers 8,134 citations

46 h-index 84 g-index

160 all docs

160 docs citations

160 times ranked 5726 citing authors

#	Article	IF	CITATIONS
1	Pituitary Society Delphi Survey: An international perspective on endocrine management of patients undergoing transsphenoidal surgery for pituitary adenomas. Pituitary, 2022, 25, 64-73.	1.6	7
2	Postoperative remission of non-invasive lactotroph pituitary tumor: A single-center experience. Annales D'Endocrinologie, 2022, 83, 1 -8.	0.6	5
3	First Case of Chronic Post-Traumatic Anterior Pituitary Dysfunction in a Professional Rugby Player: A Case Report. Annales D'Endocrinologie, 2022, , .	0.6	O
4	Maintenance of response to oral octreotide compared with injectable somatostatin receptor ligands in patients with acromegaly: a phase 3, multicentre, randomised controlled trial. Lancet Diabetes and Endocrinology,the, 2022, 10, 102-111.	5 . 5	23
5	Intratumoural spatial distribution of S100B + folliculostellate cells is associated with proliferation and expression of FSH and ERα in gonadotroph tumours. Acta Neuropathologica Communications, 2022, 10, 18.	2.4	5
6	Clinical Biology of the Pituitary Adenoma. Endocrine Reviews, 2022, 43, 1003-1037.	8.9	81
7	Consensus statement by the French Society of Endocrinology (SFE) and French Society of Pediatric Endocrinology & Samp; Diabetology (SFEDP) on diagnosis of Cushing's syndrome. Annales D'Endocrinologie, 2022, 83, 119-141.	0.6	23
8	Identification of predictive criteria for pathogenic variants of primary bilateral macronodular adrenal hyperplasia (PBMAH) gene <i>ARMC5</i> in 352 unselected patients. European Journal of Endocrinology, 2022, 187, 123-134.	1.9	18
9	Immunotherapy in aggressive pituitary tumors and carcinomas: a systematic review. Endocrine-Related Cancer, 2022, 29, 415-426.	1.6	20
10	Pituitary MRI Features in Acromegaly Resulting From Ectopic GHRH Secretion From a Neuroendocrine Tumor: Analysis of 30 Cases. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e3313-e3320.	1.8	7
11	Aggressive corticotroph tumors and carcinomas. Journal of Neuroendocrinology, 2022, 34, .	1.2	20
12	Second line treatment of acromegaly: Pasireotide or Pegvisomant?. Best Practice and Research in Clinical Endocrinology and Metabolism, 2022, 36, 101684.	2.2	11
13	Prolactin immunoassay: does the high-dose hook effect still exist?. Pituitary, 2022, 25, 653-657.	1.6	10
14	Pegvisomant in combination or pegvisomant alone after failure of somatostatin analogs in acromegaly patients: an observational French ACROSTUDY cohort study. Endocrine, 2021, 71, 158-167.	1.1	8
15	A key role for conservative treatment in the management of pituitary apoplexy. Endocrine, 2021, 71, 168-177.	1.1	7
16	Management of aggressive pituitary tumors. , 2021, , 485-497.		0
17	Pituitary Neoplasm Nomenclature Workshop: Does Adenoma Stand the Test of Time?. Journal of the Endocrine Society, 2021, 5, bvaa205.	0.1	31
18	Letter to the Editor From Helene Lasolle and Gérald Raverot: "USP8 and TP53 Drivers Are Associated With CNV in a Corticotroph Adenoma Cohort Enriched for Aggressive Tumors― Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3285-e3286.	1.8	1

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19	Graves' Disease during Immune Checkpoint Inhibitor Therapy (A Case Series and Literature Review). Cancers, 2021, 13, 1944.	1.7	21
20	Pegvisomant treatment in acromegaly in clinical practice: Final results of the French ACROSTUDY (312) Tj ETQ	q0 0 0 rgB1	Oyerlock 10
21	Aggressive pituitary tumours and pituitary carcinomas. Nature Reviews Endocrinology, 2021, 17, 671-684.	4.3	60
22	Somatostatin receptor ligands induce TSH deficiency in thyrotropin-secreting pituitary adenoma. European Journal of Endocrinology, 2021, 184, 1-8.	1.9	13
23	Cost-Utility of Acromegaly Pharmacological Treatments in a French Context. Frontiers in Endocrinology, 2021, 12, 745843.	1.5	4
24	Aggressive prolactinomas: how to manage?. Pituitary, 2020, 23, 70-77.	1.6	22
25	A Somatostatin Receptor Subtype-3 (SST3) Peptide Agonist Shows Antitumor Effects in Experimental Models of Nonfunctioning Pituitary Tumors. Clinical Cancer Research, 2020, 26, 957-969.	3.2	34
26	Current status and treatment modalities in metastases to the pituitary: a systematic review. Journal of Neuro-Oncology, 2020, 146, 219-227.	1.4	27
27	Frequency and Incidence of Carney Complex Manifestations: A Prospective Multicenter Study With a Three-Year Follow-Up. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e436-e446.	1.8	30
28	Definition and diagnosis of aggressive pituitary tumors. Reviews in Endocrine and Metabolic Disorders, 2020, 21, 203-208.	2.6	33
29	GNAS mutated thyroid carcinoma in a patient with Mc Cune Albright syndrome. Bone Reports, 2020, 13, 100299.	0.2	6
30	Chromosomal instability in the prediction of pituitary neuroendocrine tumors prognosis. Acta Neuropathologica Communications, 2020, 8, 190.	2.4	20
31	Evaluation of Nurses' and Patients' Overall Satisfaction with New and Previous Formulations of Octreotide Long-acting Release (Sandostatin LAR®): A French Observational Study. Advances in Therapy, 2020, 37, 3901-3915.	1.3	2
32	Treatment Options for Gonadotroph Tumors: Current State and Perspectives. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e3507-e3518.	1.8	15
33	Immune Landscape of Pituitary Tumors Reveals Association Between Macrophages and Gonadotroph Tumor Invasion. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 3459-3473.	1.8	23
34	Immunotherapy in Corticotroph and Lactotroph Aggressive Tumors and Carcinomas: Two Case Reports and a Review of the Literature. Journal of Personalized Medicine, 2020, 10, 88.	1.1	49
35	Anti-PD1 and Anti-PDL1-Induced Hypophysitis: A Cohort Study of 17 Patients with Longitudinal Follow-Up. Journal of Clinical Medicine, 2020, 9, 3280.	1.0	27
36	Metformin and everolimus in neuroendocrine tumours: A synergic effect?. Clinics and Research in Hepatology and Gastroenterology, 2020, 44, 954-960.	0.7	5

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37	Are aggressive pituitary tumors and carcinomas two sides of the same coin? Pathologists reply to clinician's questions. Reviews in Endocrine and Metabolic Disorders, 2020, 21, 243-251.	2.6	30
38	Aggressive Pituitary Adenomas and Carcinomas. Endocrinology and Metabolism Clinics of North America, 2020, 49, 505-515.	1.2	28
39	How to Classify Pituitary Neuroendocrine Tumors (PitNET)s in 2020. Cancers, 2020, 12, 514.	1.7	123
40	Evolution of macroprolactinomas during pregnancy: A cohort study of 85 pregnancies. Clinical Endocrinology, 2020, 92, 421-427.	1.2	9
41	Gonadotroph Tumors Show Subtype Differences that Might Have Implications for Therapy. Cancers, 2020, 12, 1012.	1.7	17
42	UrinaryÂfree cortisol: An automated immunoassay without extraction for diagnosis of Cushing's syndrome and followâ€up of patients treated by anticortisolic drugs. Clinical Endocrinology, 2020, 93, 76-78.	1.2	4
43	SST5 expression and USP8 mutation in functioning and silent corticotroph pituitary tumors. Endocrine Connections, 2020, 9, 243-253.	0.8	25
44	Pasireotide for acromegaly: long-term outcomes from an extension to the Phase III PAOLA study. European Journal of Endocrinology, 2020, 182, 583.	1.9	36
45	Efficacy and safety of dopamine agonists in patients treated with antipsychotics and presenting a macroprolactinoma. European Journal of Endocrinology, 2020, 183, 221-231.	1.9	8
46	Craniopharyngioma: Endocrinological Aspects After Surgery. , 2020, , 145-156.		1
47	Predicting early post-operative remission in pituitary adenomas: evaluation of the modified knosp classification. Pituitary, 2019, 22, 467-475.	1.6	27
48	Is Gross Total Resection Reasonable in Adults with Craniopharyngiomas with Hypothalamic Involvement?. World Neurosurgery, 2019, 129, e803-e811.	0.7	24
49	Emerging and Novel Treatments for Pituitary Tumors. Journal of Clinical Medicine, 2019, 8, 1107.	1.0	34
50	The Microenvironment of Pituitary Tumorsâ€"Biological and Therapeutic Implications. Cancers, 2019, 11, 1605.	1.7	44
51	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
52	Diagnosis, pathology, and management of TSH-secreting pituitary tumors. A single-center retrospective study of 20Âpatients from 1981 to 2014. Annales D'Endocrinologie, 2019, 80, 216-224.	0.6	9
53	Ommaya Reservoir System for the Treatment of Cystic Craniopharyngiomas: Surgical Results in a Series of 11 Adult Patients and Review of the Literature. World Neurosurgery, 2019, 132, e869-e877.	0.7	29
54	Modern neuro-ophthalmological evaluation of patients with pituitary disorders. Best Practice and Research in Clinical Endocrinology and Metabolism, 2019, 33, 101279.	2.2	8

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55	Clinical, Pathological, and Molecular Factors of Aggressiveness in Lactotroph Tumours. Neuroendocrinology, 2019, 109, 70-76.	1.2	44
56	Pheochromocytoma, paragangliomas, and pituitary adenoma. Medicine (United States), 2019, 98, e16594.	0.4	5
57	Classification of Pituitary Neuroendocrine Tumors (PitNets). , 2019, , 176-184.		3
58	Pasireotide: A potential therapeutic alternative for resistant prolactinoma. Annales D'Endocrinologie, 2019, 80, 84-88.	0.6	38
59	Bilateral adrenalectomy in Cushing's disease: Altered long-term quality of life compared to other treatment options. Annales D'Endocrinologie, 2019, 80, 32-37.	0.6	17
60	Signs and symptoms of acromegaly at diagnosis: the physician's and the patient's perspectives in the ACRO-POLIS study. Endocrine, 2019, 63, 120-129.	1.1	51
61	Evaluation of the Efficacy and Safety of Switching to Pasireotide in Patients With Acromegaly Inadequately Controlled With First-Generation Somatostatin Analogs. Frontiers in Endocrinology, 2019, 10, 931.	1.5	21
62	Cabergoline in severe ectopic or occult Cushing's syndrome. European Journal of Endocrinology, 2019, 181, K1-K9.	1.9	8
63	Confirmation of a new therapeutic option for aggressive or dopamine agonist-resistant prolactin pituitary neuroendocrine tumors. European Journal of Endocrinology, 2019, 181, C1-C3.	1.9	20
64	Pathological markers of somatotroph pituitary neuroendocrine tumors predicting the response to medical treatment. Minerva Endocrinologica, 2019, 44, 129-136.	1.7	14
65	Diagnosis and Clinical Management of Aggressive Pituitary Tumors. , 2019, , 294-300.		1
66	Ectopic acromegaly due to growth hormone-releasing hormone producing bronchial carcinoid causing somatotroph hyperplasia and partial pituitary insufficiency. Polish Archives of Internal Medicine, 2019, 129, 208-210.	0.3	4
67	Centralization errors in comparative genomic hybridization array analysis of pituitary tumor samples. Genes Chromosomes and Cancer, 2018, 57, 320-328.	1.5	2
68	Hepatic safety of ketoconazole in Cushing's syndrome: results of a Compassionate Use Programme in France. European Journal of Endocrinology, 2018, 178, 447-458.	1.9	46
69	Aggressive pituitary tumours and carcinomas: two sides of the same coin?. European Journal of Endocrinology, 2018, 178, C7-C9.	1.9	54
70	Physiopathology, Diagnosis, and Treatment of Nonfunctioning Pituitary Adenomas. Endocrinology, 2018, , 93-128.	0.1	1
71	MANAGEMENT OF ENDOCRINE DISEASE Hyperandrogenic states in women: pitfalls in laboratory diagnosis. European Journal of Endocrinology, 2018, 178, R141-R154.	1.9	19
72	Prognostic factors of regrowth in nonfunctioning pituitary tumors. Pituitary, 2018, 21, 176-182.	1.6	28

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73	Triple-A syndrome: a wide spectrum of adrenal dysfunction. European Journal of Endocrinology, 2018, 178, 199-207.	1.9	38
74	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
75	Dysregulation of cell cycle in animal models and human neuroendocrine pituitary tumors (PitNET). Cell Cycle, 2018, 17, 917-917.	1.3	0
76	European Society of Endocrinology Clinical Practice Guidelines for the management of aggressive pituitary tumours and carcinomas. European Journal of Endocrinology, 2018, 178, G1-G24.	1.9	387
77	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
78	35e Congrès de la Société française d'endocrinologie (SFE) Nancy 2018. Annales D'Endocrinologie, 2018, 79, 191.	0.6	0
79	ALK7 expression in prolactinoma is associated with reduced prolactin and increased proliferation. Endocrine-Related Cancer, 2018, 25, 795-806.	1.6	10
80	Unusual neurologic presentation of aseptic abscesses syndrome. Neurology: Neuroimmunology and NeuroInflammation, 2018, 5, e469.	3.1	1
81	Physiopathology, Diagnosis, and Treatment of Nonfunctioning Pituitary Adenomas. Endocrinology, 2018, , 1-37.	0.1	0
82	MicroRNAs in pituitary tumors. Molecular and Cellular Endocrinology, 2017, 456, 51-61.	1.6	39
83	From pituitary adenoma to pituitary neuroendocrine tumor (PitNET): an International Pituitary Pathology Club proposal. Endocrine-Related Cancer, 2017, 24, C5-C8.	1.6	262
84	Transdifferentiation of Neuroendocrine Cells. American Journal of Surgical Pathology, 2017, 41, 849-853.	2.1	19
85	Temozolomide treatment can improve overall survival in aggressive pituitary tumors and pituitary carcinomas. European Journal of Endocrinology, 2017, 176, 769-777.	1.9	107
86	Criteria for the definition of Pituitary Tumor Centers of Excellence (PTCOE): A Pituitary Society Statement. Pituitary, 2017, 20, 489-498.	1.6	233
87	Reconsidering olfactory bulb magnetic resonance patterns in Kallmann syndrome. Annales D'Endocrinologie, 2017, 78, 455-461.	0.6	7
88	Risk of Recurrence in Pituitary Neuroendocrine Tumors: A Prospective Study Using a Five-Tiered Classification. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 3368-3374.	1.8	112
89	Risks and Benefits of Endoscopic Transsphenoidal Surgery for Nonfunctioning Pituitary Adenomas in Patients of the Ninth Decade. World Neurosurgery, 2017, 106, 315-321.	0.7	14
90	Silent somatotroph tumour revisited from a study of 80 patients with and without acromegaly and a review of the literature. European Journal of Endocrinology, 2017, 176, 195-201.	1.9	33

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91	An old retrocardiac mass fortuitously reclassified as paraganglioma. Annales D'Endocrinologie, 2016, 77, 668-669.	0.6	0
92	Therapeutic innovations in endocrine diseases – partÂ3Â: temozolomide and future therapeutics for aggressive pituitary tumors and carcinomas. Presse Medicale, 2016, 45, e211-e216.	0.8	9
93	Performance of the 4-mg intravenous dexamethasone suppression test in differentiating Cushing disease from pseudo-Cushing syndrome. Annales D'Endocrinologie, 2016, 77, 30-36.	0.6	7
94	Establishment of revised diagnostic cut-offs for adrenal laboratory investigation using the new Roche Diagnostics Elecsys® Cortisol II assay. Annales D'Endocrinologie, 2016, 77, 620-622.	0.6	39
95	T2-weighted MRI signal predicts hormone and tumor responses to somatostatin analogs in acromegaly. Endocrine-Related Cancer, 2016, 23, 871-881.	1.6	82
96	Differential Effects of PI3K and Dual PI3K/mTOR Inhibition in Rat Prolactin-Secreting Pituitary Tumors. Molecular Cancer Therapeutics, 2016, 15, 1261-1270.	1.9	19
97	Dramatic change in skin color after bilateral adrenalectomy in Cushing's disease. Annales D'Endocrinologie, 2016, 77, 623-624.	0.6	0
98	Pituitary siderosis: the dark side of the pituitary. Lancet Diabetes and Endocrinology, the, 2016, 4, 374.	5.5	2
99	Clinicopathological prognostic and theranostic markers in pituitary tumors. Minerva Endocrinologica, 2016, 41, 377-89.	1.7	12
100	Pituitary Involvement in Granulomatosis With Polyangiitis. Medicine (United States), 2015, 94, e748.	0.4	61
101	Dilated Cardiomyopathy Revealing Cushing Disease. Medicine (United States), 2015, 94, e2011.	0.4	16
102	Deregulation of miR-183 and KIAAO101 in Aggressive and Malignant Pituitary Tumors. Frontiers in Medicine, 2015, 2, 54.	1.2	33
103	Expression of estrogen receptor alpha is associated with prolactin pituitary tumor prognosis and supports the sex-related difference in tumor growth. European Journal of Endocrinology, 2015, 172, 791-801.	1.9	76
104	A Prognostic Clinicopathologic Classification of Pituitary Endocrine Tumors. Endocrinology and Metabolism Clinics of North America, 2015, 44, 11-18.	1.2	29
105	Pituitary MRI characteristics in 297 acromegaly patients based on T2-weighted sequences. Endocrine-Related Cancer, 2015, 22, 169-177.	1.6	78
106	Biological and radiological exploration and management of non-functioning pituitary adenoma. Annales D'Endocrinologie, 2015, 76, 201-209.	0.6	25
107	Macroprolactinaemia: a biological diagnostic strategy from the study of 222 patients. European Journal of Endocrinology, 2015, 172, 687-695.	1.9	9
108	Downregulation of miR-410 targeting the cyclin B1 gene plays a role in pituitary gonadotroph tumors. Cell Cycle, 2015, 14, 2590-2597.	1.3	46

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109	Management of clinically non-functioning pituitary adenoma. Annales D'Endocrinologie, 2015, 76, 239-247.	0.6	136
110	<i>HMGA1</i> -pseudogene expression is induced in human pituitary tumors. Cell Cycle, 2015, 14, 1471-1475.	1.3	48
111	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
112	MANAGEMENT OF ENDOCRINE DISEASE: Clinicopathological classification and molecular markers of pituitary tumours for personalized therapeutic strategies. European Journal of Endocrinology, 2014, 170, R121-R132.	1.9	81
113	Expression of somatostatin receptors, SSTR2A and SSTR5, in 108 endocrine pituitary tumors using immunohistochemical detection with new specific monoclonal antibodies. Human Pathology, 2014, 45, 71-77.	1.1	79
114	Ketoconazole in Cushing's Disease: Is It Worth a Try?. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1623-1630.	1.8	231
115	Pasireotide versus continued treatment with octreotide or lanreotide in patients with inadequately controlled acromegaly (PAOLA): a randomised, phase 3 trial. Lancet Diabetes and Endocrinology,the, 2014, 2, 875-884.	5.5	309
116	Giant prolactinomas in women. European Journal of Endocrinology, 2014, 170, 31-38.	1.9	64
117	miR-23b and miR-130b expression is downregulated in pituitary adenomas. Molecular and Cellular Endocrinology, 2014, 390, 1-7.	1.6	78
118	Non-functioning pituitary macro-incidentalomas benefit from early surgery before becoming symptomatic. Clinical Neurology and Neurosurgery, 2013, 115, 2514-2520.	0.6	25
119	Silent GH pituitary tumor: Diagnostic and therapeutic challenges. Annales D'Endocrinologie, 2013, 74, 491-495.	0.6	15
120	A new prognostic clinicopathological classification of pituitary adenomas: a multicentric case–control study of 410 patients with 8Âyears post-operative follow-up. Acta Neuropathologica, 2013, 126, 123-135.	3.9	395
121	Aggressive Silent GH Pituitary Tumor Resistant to Multiple Treatments, Including Temozolomide. Cancer Investigation, 2013, 31, 190-196.	0.6	32
122	18F-FDG PET/CT Findings in a Patient With Isolated Intracranial Rosai-Dorfman Disease. Clinical Nuclear Medicine, 2013, 38, e50-e52.	0.7	17
123	Altered MicroRNA Expression Profile in Human Pituitary GH Adenomas: Down-Regulation of miRNA Targeting HMGA1, HMGA2, and E2F1. Journal of Clinical Endocrinology and Metabolism, 2012, 97, E1128-E1138.	1.8	136
124	Hypothalamo-pituitary sarcoidosis: a multicenter study of 24 patients. QJM - Monthly Journal of the Association of Physicians, 2012, 105, 981-995.	0.2	116
125	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
126	Acromegaly induced by ectopic secretion of GHRH: A review 30 years after GHRH discovery. Annales D'Endocrinologie, 2012, 73, 497-502.	0.6	64

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127	Pituitary carcinomas and aggressive pituitary tumours: merits and pitfalls of temozolomide treatment. Clinical Endocrinology, 2012, 76, 769-775.	1.2	125
128	New targeted therapies in pituitary carcinoma resistant to temozolomide. Pituitary, 2012, 15, 37-43.	1.6	87
129	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
130	Evidence of improved surgical outcome following endoscopy for nonfunctioning pituitary adenoma removal. Neurosurgical Focus, 2011, 30, E11.	1.0	126
131	Ectopic ACTH Syndrome in Children and Adolescents. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 1213-1222.	1.8	63
132	Integrated Genomic Profiling Identifies Loss of Chromosome 11p Impacting Transcriptomic Activity in Aggressive Pituitary PRL Tumours., 2011,, P3-69-P3-69.		0
133	Cabergoline therapy of paraneoplastic cushing syndrome in children. Pediatric Blood and Cancer, 2010, 55, 590-591.	0.8	3
134	Clinical Characteristics and Therapeutic Responses in Patients with Germ-Line <i>AIP </i> Mutations and Pituitary Adenomas: An International Collaborative Study. Journal of Clinical Endocrinology and Metabolism, 2010, 95, E373-E383.	1.8	323
135	Clinical, hormonal and molecular characterization of pituitary ACTH adenomas without (silent) Tj ETQq1 1 0.7843 35-43.	14 rgBT /C 1.9	Overlock 10 94
136	Carcinome hypophysaire. , 2010, , 441-445.		2
137	Prognostic Factors in Prolactin Pituitary Tumors: Clinical, Histological, and Molecular Data from a Series of 94 Patients with a Long Postoperative Follow-Up. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1708-1716.	1.8	144
138	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
139	Sex hormone-binding globulin gene expression in the liver: Drugs and the metabolic syndrome. Molecular and Cellular Endocrinology, 2010, 316, 53-59.	1.6	100
140	Proliferation markers of human pituitary tumors: Contribution of a genome-wide transcriptome approach. Molecular and Cellular Endocrinology, 2010, 326, 30-39.	1.6	27
141	Secondary deterioration of visual field during cabergoline treatment for macroprolactinoma. Clinical Endocrinology, 2009, 70, 588-592.	1.2	42
142	Predicting Visual Outcome After Treatment of Pituitary Adenomas With Optical Coherence Tomography. American Journal of Ophthalmology, 2009, 147, 64-70.e2.	1.7	135
143	Aryl Hydrocarbon Receptor-Interacting Protein Gene Mutations in Familial Isolated Pituitary Adenomas: Analysis in 73 Families. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 1891-1896.	1.8	283
144	Familial pituitary adenomas with a heterogeneous functional pattern: Clinical and genetic features. Journal of Endocrinological Investigation, 2007, 30, 787-790.	1.8	5

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145	A diagnostic marker set for invasion, proliferation, and aggressiveness of prolactin pituitary tumors. Endocrine-Related Cancer, 2007, 14, 887-900.	1.6	146
146	Mitotane Has an Estrogenic Effect on Sex Hormone-Binding Globulin and Corticosteroid-Binding Globulin in Humans. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 2165-2170.	1.8	75
147	Sox3 expression in undifferentiated spermatogonia is required for the progression of spermatogenesis. Developmental Biology, 2005, 283, 215-225.	0.9	142
148	Nuclear receptors Sf1 and Dax1 function cooperatively to mediate somatic cell differentiation during testis development. Development (Cambridge), 2005, 132, 2415-2423.	1.2	81
149	Influence of SHBGGene Pentanucleotide TAAAA Repeat and D327N Polymorphism on Serum Sex Hormone-Binding Globulin Concentration in Hirsute Women. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 917-924.	1.8	109
150	X-Linked Sex-Determining Region Y Box 3 (SOX3) Gene Mutations Are Uncommon in Men with Idiopathic Oligoazoospermic Infertility. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 4146-4148.	1.8	19
151	Sox3 Is Required for Gonadal Function, but Not Sex Determination, in Males and Females. Molecular and Cellular Biology, 2003, 23, 8084-8091.	1.1	168
152	Neoadjuvant B-RAF and MEK Inhibitor Targeted Therapy for Adult Papillary Craniopharyngiomas: A New Treatment Paradigm. Frontiers in Endocrinology, 0, 13, .	1.5	14