Frederic Castinetti

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Consensus on diagnosis and management of Cushing's disease: a guideline update. Lancet Diabetes and Endocrinology,the, 2021, 9, 847-875.	11.4	315
2	Long-term follow-up of ipilimumab-induced hypophysitis, a common adverse event of the anti-CTLA-4 antibody in melanoma. European Journal of Endocrinology, 2015, 172, 195-204.	3.7	232
3	Ketoconazole in Cushing's Disease: Is It Worth a Try?. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1623-1630.	3.6	231
4	Outcome of Gamma Knife Radiosurgery in 82 Patients with Acromegaly: Correlation with Initial Hypersecretion. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4483-4488.	3.6	209
5	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.	3.7	196
6	Genetics, diagnosis, management and future directions of research of phaeochromocytoma and paraganglioma: a position statement and consensus of the Working Group on Endocrine Hypertension of the European Society of Hypertension. Journal of Hypertension, 2020, 38, 1443-1456.	0.5	190
7	Gamma knife radiosurgery is a successful adjunctive treatment in Cushing's disease. European Journal of Endocrinology, 2007, 156, 91-98.	3.7	166
8	Long-Term Results of Stereotactic Radiosurgery in Secretory Pituitary Adenomas. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 3400-3407.	3.6	164
9	Ketoconazole revisited: a preoperative or postoperative treatment in Cushing's disease. European Journal of Endocrinology, 2008, 158, 91-99.	3.7	158
10	Merits and pitfalls of mifepristone in Cushing's syndrome. European Journal of Endocrinology, 2009, 160, 1003-1010.	3.7	141
11	Molecular mechanisms of pituitary organogenesis: In search of novel regulatory genes. Molecular and Cellular Endocrinology, 2010, 323, 4-19.	3.2	140
12	Outcomes of adrenal-sparing surgery or total adrenalectomy in phaeochromocytoma associated with multiple endocrine neoplasia type 2: an international retrospective population-based study. Lancet Oncology, The, 2014, 15, 648-655.	10.7	137
13	Management of clinically non-functioning pituitary adenoma. Annales D'Endocrinologie, 2015, 76, 239-247.	1.4	136
14	Pituitary carcinomas and aggressive pituitary tumours: merits and pitfalls of temozolomide treatment. Clinical Endocrinology, 2012, 76, 769-775.	2.4	125
15	Long-term prognosis of patients with pediatric pheochromocytoma. Endocrine-Related Cancer, 2014, 21, 17-25.	3.1	121
16	Temozolomide treatment can improve overall survival in aggressive pituitary tumors and pituitary carcinomas. European Journal of Endocrinology, 2017, 176, 769-777.	3.7	107
17	Complications Related to the Endoscopic Endonasal Transsphenoidal Approach for Nonfunctioning Pituitary Macroadenomas in 300 Consecutive Patients. World Neurosurgery, 2016, 89, 442-453.	1.3	101
18	Role of stereotactic radiosurgery in the management of pituitary adenomas. Nature Reviews Endocrinology, 2010, 6, 214-223.	9.6	99

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19	Prospective comparison of 68Ga-DOTATATE and 18F-FDOPA PET/CT in patients with various pheochromocytomas and paragangliomas with emphasis on sporadic cases. European Journal of Nuclear Medicine and Molecular Imaging, 2016, 43, 1248-1257.	6.4	96
20	French Endocrine Society Guidance on endocrine side effects of immunotherapy. Endocrine-Related Cancer, 2019, 26, G1-G18.	3.1	95
21	Pituitary Stem Cell Update and Potential Implications for Treating Hypopituitarism. Endocrine Reviews, 2011, 32, 453-471.	20.1	86
22	Natural history, treatment, and long-term follow up of patients with multiple endocrine neoplasia type 2B: an international, multicentre, retrospective study. Lancet Diabetes and Endocrinology,the, 2019, 7, 213-220.	11.4	86
23	Desmopressin test during petrosal sinus sampling: a valuable tool to discriminate pituitary or ectopic ACTH-dependent Cushing's syndrome. European Journal of Endocrinology, 2007, 157, 271-277.	3.7	84
24	15 YEARS OF PARAGANGLIOMA: Imaging and imaging-based treatment of pheochromocytoma and paraganglioma. Endocrine-Related Cancer, 2015, 22, T135-T145.	3.1	84
25	T2-weighted MRI signal predicts hormone and tumor responses to somatostatin analogs in acromegaly. Endocrine-Related Cancer, 2016, 23, 871-881.	3.1	82
26	Comparison of Pheochromocytoma-Specific Morbidity and Mortality Among Adults With Bilateral Pheochromocytomas Undergoing Total Adrenalectomy vs Cortical-Sparing Adrenalectomy. JAMA Network Open, 2019, 2, e198898.	5.9	80
27	Pituitary stalk interruption syndrome in 83 patients: novel HESX1 mutation and severe hormonal prognosis in malformative forms. European Journal of Endocrinology, 2011, 164, 457-465.	3.7	77
28	Cabergoline for Cushing's disease: a large retrospective multicenter study. European Journal of Endocrinology, 2017, 176, 305-314.	3.7	77
29	A Novel Dysfunctional LHX4 Mutation with High Phenotypical Variability in Patients with Hypopituitarism. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 2790-2799.	3.6	73
30	Updates on the role of adrenal steroidogenesis inhibitors in Cushing's syndrome: a focus on novel therapies. Pituitary, 2016, 19, 643-653.	2.9	70
31	MANAGEMENT OF ENDOCRINE DISEASE: Immune check point inhibitors-induced hypophysitis. European Journal of Endocrinology, 2019, 181, R107-R118.	3.7	68
32	MANAGEMENT OF ENDOCRINE DISEASE: Management of Cushing's syndrome during pregnancy: solved and unsolved questions. European Journal of Endocrinology, 2018, 178, R259-R266.	3.7	67
33	Aggressive pituitary tumours and pituitary carcinomas. Nature Reviews Endocrinology, 2021, 17, 671-684.	9.6	60
34	A comprehensive review on MEN2B. Endocrine-Related Cancer, 2018, 25, T29-T39.	3.1	58
35	Radiotherapy and radiosurgery in acromegaly. Pituitary, 2009, 12, 3-10.	2.9	56
36	18F-FDOPA PET/CT imaging of insulinoma revisited. European Journal of Nuclear Medicine and Molecular Imaging, 2015, 42, 409-418.	6.4	54

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37	Bilateral adrenalectomy in the 21st century: when to use it for hypercortisolism?. Endocrine-Related Cancer, 2016, 23, R131-R142.	3.1	54
38	MANAGEMENT OF ENDOCRINE DISEASE: Outcome of adrenal sparing surgery in heritable pheochromocytoma. European Journal of Endocrinology, 2016, 174, R9-R18.	3.7	54
39	Medical Treatment of Cushing's Syndrome: Glucocorticoid Receptor Antagonists and Mifepristone. Neuroendocrinology, 2010, 92, 125-130.	2.5	53
40	Pharmacokinetic Evidence for Suboptimal Treatment of Adrenal Insufficiency with Currently Available Hydrocortisone Tablets. Clinical Pharmacokinetics, 2010, 49, 455-463.	3.5	53
41	65 YEARS OF THE DOUBLE HELIX: Genetics informs precision practice in the diagnosis and management of pheochromocytoma. Endocrine-Related Cancer, 2018, 25, T201-T219.	3.1	52
42	Persistent and recurrent hyperparathyroidism. Updates in Surgery, 2017, 69, 161-169.	2.0	50
43	DIAGNOSIS OF ENDOCRINE DISEASE: Pituitary stalk interruption syndrome: etiology and clinical manifestations. European Journal of Endocrinology, 2019, 181, R199-R209.	3.7	50
44	MECHANISMS IN ENDOCRINOLOGY: An update in the genetic aetiologies of combined pituitary hormone deficiency. European Journal of Endocrinology, 2016, 174, R239-R247.	3.7	49
45	Cushing's disease. Orphanet Journal of Rare Diseases, 2012, 7, 41.	2.7	46
46	Hepatic safety of ketoconazole in Cushing's syndrome: results of a Compassionate Use Programme in France. European Journal of Endocrinology, 2018, 178, 447-458.	3.7	46
47	The use of the glucocorticoid receptor antagonist mifepristone in Cushing's syndrome. Current Opinion in Endocrinology, Diabetes and Obesity, 2012, 19, 295-299.	2.3	45
48	A Combined Dexamethasone Desmopressin Test as an Early Marker of Postsurgical Recurrence in Cushing's Disease. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 1897-1903.	3.6	44
49	Outcome of multimodal therapy in operated acromegalic patients, a study in 115 patients. Clinical Endocrinology, 2013, 78, 263-270.	2.4	44
50	Delayed diagnosis of Sheehan's syndrome in a developed country: a retrospective cohort study. European Journal of Endocrinology, 2013, 169, 431-438.	3.7	43
51	Preoperative imaging for focused parathyroidectomy: making a good strategy even better. European Journal of Endocrinology, 2015, 172, 519-526.	3.7	40
52	The risks of overlooking the diagnosis of secreting pituitary adenomas. Orphanet Journal of Rare Diseases, 2016, 11, 135.	2.7	39
53	Approach to the Patient Treated with Steroidogenesis Inhibitors. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 2114-2123.	3.6	39
54	SFE/SFEDP adrenal insufficiency French consensus: Introduction and handbook. Annales D'Endocrinologie, 2018, 79, 1-22.	1.4	38

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55	Combined pituitary hormone deficiency: current and future status. Journal of Endocrinological Investigation, 2015, 38, 1-12.	3.3	37
56	Three Novel Heterozygous Point Mutations of <i>NR3C1</i> Causing Glucocorticoid Resistance. Human Mutation, 2016, 37, 794-803.	2.5	34
57	Risk Profile of the RET A883F Germline Mutation: An International Collaborative Study. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 2069-2074.	3.6	34
58	A conservative management is preferable in milder forms of pituitary tumor apoplexy. Journal of Endocrinological Investigation, 2011, 34, 502-9.	3.3	32
59	Significant prevalence of NR3C1 mutations in incidentally discovered bilateral adrenal hyperplasia: results of the French MUTA-GR Study. European Journal of Endocrinology, 2018, 178, 411-423.	3.7	31
60	Non-functioning pituitary adenoma: When and how to operate? What pathologic criteria for typing?. Annales D'Endocrinologie, 2015, 76, 220-227.	1.4	30
61	A registry-based study of thyroid paraganglioma: histological and genetic characteristics. Endocrine-Related Cancer, 2015, 22, 191-204.	3.1	29
62	ISL1 Is Necessary for Maximal Thyrotrope Response to Hypothyroidism. Molecular Endocrinology, 2015, 29, 1510-1521.	3.7	28
63	Quantitative 18F-DOPA PET/CT in pheochromocytoma: the relationship between tumor secretion and its biochemical phenotype. European Journal of Nuclear Medicine and Molecular Imaging, 2018, 45, 278-282.	6.4	28
64	MRI follow-up is unnecessary in patients with macroprolactinomas and long-term normal prolactin levels on dopamine agonist treatment. European Journal of Endocrinology, 2017, 176, 323-328.	3.7	27
65	18F-FDOPA PET/CT Imaging of MAX-Related Pheochromocytoma. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 1574-1582.	3.6	27
66	Long-term control of a MEN1 prolactin secreting pituitary carcinoma after temozolomide treatment. Annales D'Endocrinologie, 2012, 73, 225-229.	1.4	26
67	Bilateral neck exploration in patients with primary hyperparathyroidism and discordant imaging results: a single-centre study. European Journal of Endocrinology, 2014, 170, 719-725.	3.7	26
68	Endocrine side-effects of new anticancer therapies: Overall monitoring and conclusions. Annales D'Endocrinologie, 2018, 79, 591-595.	1.4	26
69	PITX2 AND PITX1 Regulate Thyrotroph Function and Response to Hypothyroidism. Molecular Endocrinology, 2011, 25, 1950-1960.	3.7	25
70	Pathological and Genetic Characterization of Bilateral Adrenomedullary Hyperplasia in a Patient with Germline MAX Mutation. Endocrine Pathology, 2017, 28, 302-307.	9.0	25
71	Positron Emission Tomography Imaging in Medullary Thyroid Carcinoma: Time for Reappraisal?. Thyroid, 2021, 31, 151-155.	4.5	25
72	Lanreotide for the treatment of acromegaly. Advances in Therapy, 2009, 26, 600-612.	2.9	24

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73	Diagnosis and preoperative imaging of multiple endocrine neoplasia type 2: current status and future directions. Clinical Endocrinology, 2014, 81, 317-328.	2.4	24
74	Corepressors TLE1 and TLE3 Interact with HESX1 and PROP1. Molecular Endocrinology, 2010, 24, 754-765.	3.7	23
75	High-throughput splicing assays identify missense and silent splice-disruptive POU1F1 variants underlying pituitary hormone deficiency. American Journal of Human Genetics, 2021, 108, 1526-1539.	6.2	23
76	Consensus statement by the French Society of Endocrinology (SFE) and French Society of Pediatric Endocrinology & Diabetology (SFEDP) on diagnosis of Cushing's syndrome. Annales D'Endocrinologie, 2022, 83, 119-141.	1.4	23
77	Looking beyond the thyroid: advances in the understanding of pheochromocytoma and hyperparathyroidism phenotypes in MEN2 and of non-MEN2 familial forms. Endocrine-Related Cancer, 2018, 25, T15-T28.	3.1	22
78	Clinical lessons learned in constitutional hypopituitarism from two decades of experience in a large international cohort. Clinical Endocrinology, 2021, 94, 277-289.	2.4	22
79	Copy number variations alter methylation and parallel IGF2 overexpression in adrenal tumors. Endocrine-Related Cancer, 2015, 22, 953-967.	3.1	21
80	Early 18F-FDOPA PET/CT imaging after carbidopa premedication as a valuable diagnostic option in patients with insulinoma. European Journal of Nuclear Medicine and Molecular Imaging, 2019, 46, 686-695.	6.4	21
81	MEN2-related pheochromocytoma: current state of knowledge, specific characteristics in MEN2B, and perspectives. Endocrine, 2020, 69, 496-503.	2.3	21
82	Pre-surgical medical treatment, a major prognostic factor for long-term remission in acromegaly. Pituitary, 2018, 21, 615-623.	2.9	20
83	Acromegaly in Carney complex. Pituitary, 2019, 22, 456-466.	2.9	20
84	Pheochromocytoma surgery without systematic preoperative pharmacological preparation: insights from a referral tertiary center experience. Surgical Endoscopy and Other Interventional Techniques, 2021, 35, 728-735.	2.4	20
85	Postoperative followâ€up of Cushing's disease using cortisol, desmopressin and coupled dexamethasoneâ€desmopressin tests: a headâ€toâ€head comparison. Clinical Endocrinology, 2015, 83, 216-222.	2.4	19
86	The penetrance of MEN2 pheochromocytoma is not only determined by RET mutations. Endocrine-Related Cancer, 2017, 24, L63-L67.	3.1	19
87	Active Cushing syndrome patients have increased ectopic fat deposition and bone marrow fat content compared to cured patients and healthy subjects: a pilot 1H-MRS study. European Journal of Endocrinology, 2018, 179, 307-317.	3.7	19
88	Medical management of Cushing's disease: When and how?. Journal of Neuroendocrinology, 2022, 34, e13120.	2.6	19
89	GPR101 Mutations are not a Frequent Cause of Congenital Isolated Growth Hormone Deficiency. Hormone and Metabolic Research, 2016, 48, 389-393.	1.5	18
90	Cushing Syndrome Is Associated With Subclinical LV Dysfunction and Increased Epicardial Adipose Tissue. Journal of the American College of Cardiology, 2018, 72, 2276-2277.	2.8	18

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91	Surgical indications for pituitary tumors during pregnancy: a literature review. Pituitary, 2020, 23, 189-199.	2.9	18
92	The risks of medical treatment of prolactinoma. Annales D'Endocrinologie, 2021, 82, 15-19.	1.4	18
93	Radiotherapy as a tool for the treatment of Cushing's disease. European Journal of Endocrinology, 2019, 180, D9-D18.	3.7	18
94	Which patients with acromegaly are treated with pegvisomant? An overview of methodology and baseline data in ACROSTUDY. European Journal of Endocrinology, 2009, 161, S11-S17.	3.7	17
95	Genetic causes of combined pituitary hormone deficiencies in humans. Annales D'Endocrinologie, 2012, 73, 53-55.	1.4	17
96	Heterozygous LHX3 mutations may lead to a mild phenotype of combined pituitary hormone deficiency. European Journal of Human Genetics, 2019, 27, 216-225.	2.8	17
97	Primary hyperparathyroidism as first manifestation in multiple endocrine neoplasia type 2A: an international multicenter study. Endocrine Connections, 2020, 9, 489-497.	1.9	17
98	Adrenal Myelolipoma: An Unusual Cause of Bilateral Highly 18F-FDG-Avid Adrenal Masses. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 2577-2578.	3.6	16
99	Patients lost to follow-up in acromegaly: results of the ACROSPECT study. European Journal of Endocrinology, 2014, 170, 791-797.	3.7	16
100	Lack of functional remission in Cushing's syndrome. Endocrine, 2018, 61, 518-525.	2.3	16
101	Does first-line surgery still have its place in the treatment of acromegaly?. Annales D'Endocrinologie, 2009, 70, 107-112.	1.4	15
102	Value of 123I/99mTc-sestamibi parathyroid scintigraphy with subtraction SPECT/CT in primary hyperparathyroidism for directing minimally invasive parathyroidectomy. American Journal of Surgery, 2019, 217, 108-113.	1.8	15
103	Identifying the Deleterious Effect of Rare LHX4 Allelic Variants, a Challenging Issue. PLoS ONE, 2015, 10, e0126648.	2.5	15
104	An observational study on adrenal insufficiency in a French tertiary centre: Real life versus theory. Annales D'Endocrinologie, 2015, 76, 1-8.	1.4	14
105	Contemporary review of large adrenal tumors in a tertiary referral center. Anticancer Research, 2014, 34, 2581-8.	1.1	14
106	Gamma Knife radiosurgery for hypothalamic hamartoma preserves endocrine functions. Epilepsia, 2017, 58, 72-76.	5.1	13
107	Functioning gonadotroph adenoma with severe ovarian hyperstimulation syndrome: A new emergency in pituitary adenoma surgery? Surgical considerations and literature review. Annales D'Endocrinologie, 2019, 80, 122-127.	1.4	13
108	Osilodrostat in Cushing's disease: The risk of delayed adrenal insufficiency should be carefully monitored. Clinical Endocrinology, 2023, 98, 629-630.	2.4	13

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109	Persistent cortisol response to desmopressin predicts recurrence of Cushing's disease in patients with post-operative corticotropic insufficiency. European Journal of Endocrinology, 2020, 182, 489-498.	3.7	13
110	Group 4: Replacement therapy for adrenal insufficiency. Annales D'Endocrinologie, 2017, 78, 525-534.	1.4	12
111	Increased Risk of Persistent Glucose Disorders After Control of Acromegaly. Journal of the Endocrine Society, 2017, 1, 1531-1539.	0.2	12
112	lgG4 hypophysitis: Diagnosis and management. Presse Medicale, 2020, 49, 104016.	1.9	12
113	Pituitary adenoma in patients with multiple endocrine neoplasia type 1: a cohort study. European Journal of Endocrinology, 2021, 185, 863-873.	3.7	12
114	Auto-immune thyroid dysfunction induced by tyrosine kinase inhibitors in a patient with recurrent chordoma. BMC Cancer, 2016, 16, 679.	2.6	11
115	Prospective evaluation of ⁶⁸ Gaâ€ <scp>DOTATATE PET</scp> / <scp>CT</scp> in limited disease neuroendocrine tumours and/or elevated serum neuroendocrine biomarkers. Clinical Endocrinology, 2018, 89, 155-163.	2.4	11
116	Germinal defects of SDHx genes in patients with isolated pituitary adenoma. European Journal of Endocrinology, 2020, 183, 369-379.	3.7	11
117	Evaluation of an individualized education program in pituitary diseases: a pilot study. European Journal of Endocrinology, 2020, 183, 551-559.	3.7	11
118	Large Adrenal Incidentalomas Require a Dedicated Diagnostic Procedure. Endocrine Practice, 2019, 25, 669-677.	2.1	9
119	Meningiomas in patients with long-term exposition to progestins: Characteristics and outcome. Neurochirurgie, 2021, 67, 556-563.	1.2	9
120	Controversies about the systematic preoperative pharmacological treatment before pheochromocytoma or paraganglioma surgery. European Journal of Endocrinology, 2022, 186, D17-D24.	3.7	9
121	Introduction to expert opinion on endocrine complications of new anticancer therapies. Annales D'Endocrinologie, 2018, 79, 535-538.	1.4	8
122	Discordant biological parameters of remission in acromegaly do not increase the risk of hypertension or diabetes: a study with the Liege Acromegaly Survey database. Endocrine, 2020, 70, 134-142.	2.3	8
123	Radiation techniques in aggressive pituitary tumours and carcinomas. Reviews in Endocrine and Metabolic Disorders, 2020, 21, 287-292.	5.7	8
124	Recurrence-Free Survival Analysis in Locally Advanced Pheochromocytoma: First Appraisal. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 2726-2737.	3.6	8
125	Pituitary apoplexy after somatostatin analogue administration: coincidental or causative?. Clinical Endocrinology, 2014, 81, 471-473.	2.4	7
126	A monocentric experience of growth hormone replacement therapy in adult patients. Annales D'Endocrinologie, 2014, 75, 176-183.	1.4	7

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127	Transcranial approach in giant pituitary adenomas: results and outcome in a modern series. Journal of Neurosurgical Sciences, 2020, 64, 25-36.	0.6	7
128	Gamma Knife radiosurgery in pituitary adenomas: Why, who, and how to treat?. Discovery Medicine, 2010, 10, 107-11.	0.5	7
129	Current and Emerging Medical Therapies in Pituitary Tumors. Journal of Clinical Medicine, 2022, 11, 955.	2.4	7
130	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.	3.6	7
131	Lessons from monogenic causes of growth hormone deficiency. Annales D'Endocrinologie, 2017, 78, 77-79.	1.4	6
132	Clinical management of difficult to treat macroprolactinomas. Expert Review of Endocrinology and Metabolism, 2019, 14, 179-192.	2.4	6
133	Risk stratification of adrenal masses by [¹⁸ F]FDG PET/CT: Changing tactics. Clinical Endocrinology, 2021, 94, 133-140.	2.4	6
134	Synergistic cortisol suppression by ketoconazole–osilodrostat combination therapy. Endocrinology, Diabetes and Metabolism Case Reports, 2021, 2021, .	0.5	6
135	Impact of Cushing's syndrome on fertility and pregnancy. Annales D'Endocrinologie, 2022, 83, 188-190.	1.4	6
136	Pre-term birth in women exposed to Cushing's disease: the baby-cush study. European Journal of Endocrinology, 2021, 184, 469-476.	3.7	5
137	Long-term outcome of macroprolactinomas. Annales D'Endocrinologie, 2016, 77, 641-648.	1.4	4
138	Implications of SDHB genetic testing in patients with sporadic pheochromocytoma. Langenbeck's Archives of Surgery, 2017, 402, 787-798.	1.9	4
139	Characterization of adrenocortical tumors by 18F-FDG PET/CT: Does steroid hormone hypersecretion status modify the uptake pattern?. Surgical Oncology, 2018, 27, 231-235.	1.6	4
140	An Open-Label, Analgesic Efficacy and Safety of Pituitary Radiosurgery for Patients With Opioid-Refractory Pain: Study Protocol for a Randomized Controlled Trial. Neurosurgery, 2018, 83, 146-153.	1.1	4
141	Psychological impact of von Hippel-Lindau genetic screening in patients with a previous history of hemangioblastoma of the central nervous system. Journal of Psychosocial Oncology, 2018, 36, 624-634.	1.2	4
142	Tumor multifocality with vagus nerve involvement as a phenotypic marker of <i>SDHD</i> mutation in patients with head and neck paragangliomas: A ¹⁸ Fâ€FDOPA PET/CT study. Head and Neck, 2019, 41, 1565-1571.	2.0	4
143	Thyroiditis and immune check point inhibitors: the postâ€marketing experience using the French National Pharmacovigilance database. Fundamental and Clinical Pharmacology, 2019, 33, 239-240.	1.9	4
144	Commentary: The Impact of Insulin-Like Growth Factor Index and Biologically Effective Dose on Outcomes After Stereotactic Radiosurgery for Acromegaly: Cohort Study. Neurosurgery, 2020, 87, E301-E302.	1.1	4

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145	Adrenal Crisis May Occur Even In Patients With Asymptomatic Covid-19. Endocrine Practice, 2020, 26, 929-930.	2.1	4
146	Lack of delayed neurocognitive side effects of Gamma Knife radiosurgery in acromegaly: the Later-Ac study. European Journal of Endocrinology, 2022, 186, 37-44.	3.7	4
147	Osilodrostat for the treatment of Cushing's disease: efficacy, stability, and persistence – Authors' reply. Lancet Diabetes and Endocrinology,the, 2022, 10, 385-387.	11.4	4
148	Gamma Knife for Cushing disease — time for a reappraisal?. Nature Reviews Endocrinology, 2017, 13, 628-629.	9.6	3
149	Letter to the Editor: "Medullary Thyroid Carcinoma in MEN2A: ATA Moderate- or High-Risk RET Mutations Do Not Predict Disease Aggressiveness― Journal of Clinical Endocrinology and Metabolism, 2017, 102, 3557-3558.	3.6	3
150	Letter to the Editor: "Why We Should Still Treat by Neurosurgery Patients With Cushing Disease and a Normal or Inconclusive Pituitary MRI― Journal of Clinical Endocrinology and Metabolism, 2019, 104, 5791-5792.	3.6	3
151	Exploring the link between tumour metabolism and succinate dehydrogenase deficiency: A ¹⁸ Fâ€FDOPA PET/CT study in head and neck paragangliomas. Clinical Endocrinology, 2019, 91, 879-884.	2.4	3
152	Acromegaly in remission: a view from the partner. European Journal of Endocrinology, 2021, 185, K19-K23.	3.7	3
153	Familial hypocalciuric hypercalcemia: the challenge of diagnosis. Endocrine, 2022, 75, 646-649.	2.3	3
154	Radiosurgery: A Useful First-Line Treatment of Prolactinomas?. World Neurosurgery, 2010, 74, 103-104.	1.3	2
155	Medical management of adrenocortical carcinoma: Current recommendations, new therapeutic options and future perspectives. Annales D'Endocrinologie, 2021, 82, 52-58.	1.4	2
156	Women's perceptions of femininity after craniopharyngioma: a qualitative study. Clinical Endocrinology, 2021, 94, 880-887.	2.4	2
157	Congenital pituitary hormone deficiencies: role ofLHX3/LHX4genes. Expert Review of Endocrinology and Metabolism, 2008, 3, 751-760.	2.4	1
158	Disease of Adrenal Glands. International Journal of Endocrinology, 2015, 2015, 1-2.	1.5	1
159	Combined Pituitary Hormone Deficiency. , 2016, , 177-194.		1
160	Physiopathology, Diagnosis, and Treatment of Nonfunctioning Pituitary Adenomas. Endocrinology, 2018, , 93-128.	0.1	1
161	Genes important in the fetal development of the pituitary. Current Opinion in Endocrine and Metabolic Research, 2018, 1, 9-12.	1.4	1
162	Fluctuation analysis of postoperative secretory status in patients operated for acromegaly. Annales D'Endocrinologie, 2020, 81, 11-17.	1.4	1

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163	Comparison of 68Ga-Dotatate PET/CT and 18F-FDOPA PET/CT for the diagnosis of pancreatic neuroendocrine tumors in a MEN1 patient. Annales D'Endocrinologie, 2020, 81, 39-43.	1.4	1
164	Coexistence of Endocrine Side Effects of Immunotherapy in Clinical Practice. Endocrinology, 2021, , 405-411.	0.1	1
165	Role of growth hormone in hepatic and intestinal triglyceride-rich lipoprotein metabolism. Journal of Clinical Lipidology, 2021, 15, 712-723.	1.5	1
166	Letter to the Editor from Soghomonian et al.: "Epicardial and Pericardial Adiposity Without Myocardial Steatosis in Cushing Syndrome― Journal of Clinical Endocrinology and Metabolism, 2022, 107, e434-e435.	3.6	1
167	Cushingâ \in ™s disease: role of preoperative and primary medical therapy. Pituitary, 0, , .	2.9	1
168	La chirurgie de première intention a-t-elle encore une place dans le traitement de l'acromégalie�. Annales D'Endocrinologie, 2009, 70, e23-e28.	1.4	0
169	Adenomi ipofisari. EMC - AKOS - Trattato Di Medicina, 2010, 12, 1-12.	0.0	0
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