

Mã'nica R Gadelha

List of Publications by Year in descending order

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140
papers

6,290
citations

76322

40
h-index

82542

72
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144
all docs

144
docs citations

144
times ranked

3646
citing authors

#	ARTICLE	IF	CITATIONS
1	Consensus on diagnosis and management of Cushing's disease: a guideline update. <i>Lancet Diabetes and Endocrinology</i> , 2021, 9, 847-875.	11.4	315
2	Pasireotide versus continued treatment with octreotide or lanreotide in patients with inadequately controlled acromegaly (PAOLA): a randomised, phase 3 trial. <i>Lancet Diabetes and Endocrinology</i> , 2014, 2, 875-884.	11.4	309
3	The Role of the Aryl Hydrocarbon Receptor-Interacting Protein Gene in Familial and Sporadic Pituitary Adenomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008, 93, 2390-2401.	3.6	273
4	Systemic Complications of Acromegaly and the Impact of the Current Treatment Landscape: An Update. <i>Endocrine Reviews</i> , 2019, 40, 268-332.	20.1	226
5	A Consensus on the Diagnosis and Treatment of Acromegaly Comorbidities: An Update. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e937-e946.	3.6	207
6	Quantitative analysis of somatostatin receptor subtype (SSTR1 α) gene expression levels in somatotropinomas and non-functioning pituitary adenomas. <i>European Journal of Endocrinology</i> , 2007, 156, 65-74.	3.7	196
7	Multidisciplinary management of acromegaly: A consensus. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2020, 21, 667-678.	5.7	183
8	The Gene of the Ubiquitin-Specific Protease 8 Is Frequently Mutated in Adenomas Causing Cushing's Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, E997-E1004.	3.6	163
9	Quantitative analysis of somatostatin receptor subtypes (1 α) gene expression levels in somatotropinomas and correlation to in vivo hormonal and tumor volume responses to treatment with octreotide LAR. <i>European Journal of Endocrinology</i> , 2008, 158, 295-303.	3.7	160
10	A Pituitary Society update to acromegaly management guidelines. <i>Pituitary</i> , 2021, 24, 1-13.	2.9	158
11	Landscape of Familial Isolated and Young-Onset Pituitary Adenomas: Prospective Diagnosis in AIP Mutation Carriers. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, E1242-E1254.	3.6	144
12	BMI and Metabolic Profile in Patients With Prolactinoma Before and After Treatment With Dopamine Agonists. <i>Obesity</i> , 2011, 19, 800-805.	3.0	136
13	Novel pathway for somatostatin analogs in patients with acromegaly. <i>Trends in Endocrinology and Metabolism</i> , 2013, 24, 238-246.	7.1	126
14	Expression Analysis of Dopamine Receptor Subtypes in Normal Human Pituitaries, Nonfunctioning Pituitary Adenomas and Somatotropinomas, and the Association between Dopamine and Somatostatin Receptors with Clinical Response to Octreotide-LAR in Acromegaly. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 1931-1937.	3.6	120
15	Octreotide LAR vs. surgery in newly diagnosed patients with acromegaly: a randomized, open-label, multicentre study. <i>Clinical Endocrinology</i> , 2009, 70, 757-768.	2.4	108
16	Challenges in the diagnosis and management of acromegaly: a focus on comorbidities. <i>Pituitary</i> , 2016, 19, 448-457.	2.9	108
17	AIP expression in sporadic somatotropinomas is a predictor of the response to octreotide LAR therapy independent of SSTR2 expression. <i>Endocrine-Related Cancer</i> , 2012, 19, L25-L29.	3.1	100
18	Interpreting biochemical control response rates with first-generation somatostatin analogues in acromegaly. <i>Pituitary</i> , 2016, 19, 235-247.	2.9	93

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19	Giant prolactinomas: the therapeutic approach. <i>Clinical Endocrinology</i> , 2013, 79, 447-456.	2.4	91
20	Somatostatin receptor ligands in the treatment of acromegaly. <i>Pituitary</i> , 2017, 20, 100-108.	2.9	91
21	Loss of Heterozygosity on Chromosome 11q13 in Two Families with Acromegaly/Gigantism Is Independent of Mutations of the Multiple Endocrine Neoplasia Type I Gene ¹ . <i>Journal of Clinical Endocrinology and Metabolism</i> , 1999, 84, 249-256.	3.6	80
22	Prevalence of sleep apnea and metabolic abnormalities in patients with acromegaly and analysis of cephalometric parameters by magnetic resonance imaging.. <i>European Journal of Endocrinology</i> , 2008, 158, 459-465.	3.7	77
23	Adverse effects of glucocorticoids: coagulopathy. <i>European Journal of Endocrinology</i> , 2015, 173, M11-M21.	3.7	72
24	Truncated somatostatin receptor variant sst5TMD4 confers aggressive features (proliferation,) Tj ETQq0 0 0 rgBT /Qyerlock 10 Tf 50 54.	7.2	72
25	Low Aryl Hydrocarbon Receptor-Interacting Protein Expression Is a Better Marker of Invasiveness in Somatotropinomas than Ki-67 and p53. <i>Neuroendocrinology</i> , 2011, 94, 39-48.	2.5	69
26	Controversial issues in the management of hyperprolactinemia and prolactinomas – An overview by the Neuroendocrinology Department of the Brazilian Society of Endocrinology and Metabolism. <i>Archives of Endocrinology and Metabolism</i> , 2018, 62, 236-263.	0.6	69
27	The genetic background of acromegaly. <i>Pituitary</i> , 2017, 20, 10-21.	2.9	65
28	Switching patients with acromegaly from octreotide to pasireotide improves biochemical control: crossover extension to a randomized, double-blind, Phase III study. <i>BMC Endocrine Disorders</i> , 2016, 16, 16.	2.2	63
29	Genetics of Pituitary Adenomas. <i>Frontiers of Hormone Research</i> , 2013, 41, 111-140.	1.0	61
30	Predictors of surgical outcome and early criteria of remission in acromegaly. <i>Endocrine</i> , 2018, 60, 415-422.	2.3	61
31	Efficacy of medical treatment in cushing's disease: a systematic review. <i>Clinical Endocrinology</i> , 2014, 80, 1-12.	2.4	59
32	Regulation of Aryl Hydrocarbon Receptor Interacting Protein (AIP) Protein Expression by MiR-34a in Sporadic Somatotropinomas. <i>PLoS ONE</i> , 2015, 10, e0117107.	2.5	59
33	Effect of pasireotide on glucose- and growth hormone-related biomarkers in patients with inadequately controlled acromegaly. <i>Endocrine</i> , 2016, 53, 210-219.	2.3	59
34	Ipilimumab-induced hypophysitis: review of the literature. <i>Journal of Endocrinological Investigation</i> , 2015, 38, 1159-1166.	3.3	56
35	MANAGEMENT OF ENDOCRINE DISEASE: Personalized medicine in the treatment of acromegaly. <i>European Journal of Endocrinology</i> , 2018, 178, R89-R100.	3.7	56
36	Ki-67 is a predictor of acromegaly control with octreotide LAR independent of SSTR2 status and relates to cytokeratin pattern. <i>European Journal of Endocrinology</i> , 2013, 169, 217-223.	3.7	55

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37	Prevalence of discordant GH and IGF-I levels in acromegalics at diagnosis, after surgical treatment and during treatment with octreotide LAR®. <i>Growth Hormone and IGF Research</i> , 2008, 18, 389-393.	1.1	53
38	Low Frequency of Cardiomyopathy Using Cardiac Magnetic Resonance Imaging in an Acromegaly Contemporary Cohort. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 4447-4455.	3.6	51
39	Dopamine receptor subtype 2 expression profile in nonfunctioning pituitary adenomas and <i>in vivo</i> response to cabergoline therapy. <i>Clinical Endocrinology</i> , 2015, 82, 739-746.	2.4	49
40	Apoplexy in nonfunctioning pituitary adenomas. <i>Pituitary</i> , 2018, 21, 138-144.	2.9	47
41	Pituitary apoplexy during treatment of cystic macroprolactinomas with cabergoline. <i>Pituitary</i> , 2008, 11, 287-292.	2.9	40
42	Acromegaly Secondary to Growth Hormone-releasing Hormone Secreted by an Incidentally Discovered Pheochromocytoma. <i>Endocrine Pathology</i> , 2007, 18, 46-52.	9.0	39
43	Acromegaly and pregnancy: a prospective study. <i>European Journal of Endocrinology</i> , 2014, 170, 301-310.	3.7	39
44	Determinants of morbidities and mortality in acromegaly. <i>Archives of Endocrinology and Metabolism</i> , 2020, 63, 630-637.	0.6	39
45	A meiotic recombination in a new isolated familial somatotropinoma kindred. <i>European Journal of Endocrinology</i> , 2004, 150, 643-648.	3.7	38
46	Lanreotide Autogel 120µg at extended dosing intervals in patients with acromegaly biochemically controlled with octreotide LAR: the LEAD study. <i>European Journal of Endocrinology</i> , 2015, 173, 313-323.	3.7	37
47	Somatic USP8 mutations are frequent events in corticotroph tumor progression causing Nelson's tumor. <i>European Journal of Endocrinology</i> , 2018, 178, 57-63.	3.7	37
48	Bone density and microarchitecture in endogenous hypercortisolism. <i>Clinical Endocrinology</i> , 2015, 83, 468-474.	2.4	36
49	Low frequency of cardiac arrhythmias and lack of structural heart disease in medically-naïve acromegaly patients: a prospective study at baseline and after 1 year of somatostatin analogs treatment. <i>Pituitary</i> , 2016, 19, 582-589.	2.9	36
50	Pasireotide for acromegaly: long-term outcomes from an extension to the Phase III PAOLA study. <i>European Journal of Endocrinology</i> , 2020, 182, 583.	3.7	36
51	Lycopene and Beta-Carotene Induce Growth Inhibition and Proapoptotic Effects on ACTH-Secreting Pituitary Adenoma Cells. <i>PLoS ONE</i> , 2013, 8, e62773.	2.5	35
52	The Future of Somatostatin Receptor Ligands in Acromegaly. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 297-308.	3.6	35
53	Definition and diagnosis of aggressive pituitary tumors. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2020, 21, 203-208.	5.7	33
54	Prevalence of gsp oncogene in somatotropinomas and clinically non-functioning pituitary adenomas: our experience. <i>Pituitary</i> , 2009, 12, 165-169.	2.9	32

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55	Management of acromegaly in Latin America: expert panel recommendations. <i>Pituitary</i> , 2010, 13, 168-175.	2.9	31
56	Cabergoline treatment in acromegaly: cons. <i>Endocrine</i> , 2014, 46, 220-225.	2.3	31
57	Pituitary Neoplasm Nomenclature Workshop: Does Adenoma Stand the Test of Time?. <i>Journal of the Endocrine Society</i> , 2021, 5, bvaa205.	0.2	31
58	Randomized Trial of Osilodrostat for the Treatment of Cushing Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e2882-e2895.	3.6	31
59	Splicing Machinery is Dysregulated in Pituitary Neuroendocrine Tumors and is Associated with Aggressiveness Features. <i>Cancers</i> , 2019, 11, 1439.	3.7	30
60	Use of late-night salivary cortisol to monitor response to medical treatment in Cushing's disease. <i>European Journal of Endocrinology</i> , 2020, 182, 207-217.	3.7	29
61	Machine Learning-based Prediction Model for Treatment of Acromegaly With First-generation Somatostatin Receptor Ligands. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, 2047-2056.	3.6	27
62	Risk factors and management of pasireotide-associated hyperglycemia in acromegaly. <i>Endocrine Connections</i> , 2020, 9, 1178-1190.	1.9	27
63	Treatment effectiveness of pasireotide on health-related quality of life in patients with Cushing's disease. <i>European Journal of Endocrinology</i> , 2014, 171, 89-98.	3.7	26
64	ZAC1 and SSTR2 Are Downregulated in Non-Functioning Pituitary Adenomas but Not in somatotropinomas. <i>PLoS ONE</i> , 2013, 8, e77406.	2.5	25
65	Prevalence of obstructive sleep apnea in patients with prolactinoma before and after treatment with dopamine agonists. <i>Pituitary</i> , 2014, 17, 441-449.	2.9	25
66	Pituitary Tumor Management in Pregnancy. <i>Endocrinology and Metabolism Clinics of North America</i> , 2015, 44, 181-197.	3.2	25
67	Pasireotide for the treatment of acromegaly. <i>Expert Opinion on Pharmacotherapy</i> , 2016, 17, 579-588.	1.8	24
68	Glucocorticoid use in patients with adrenal insufficiency following administration of the COVID-19 vaccine: a pituitary society statement. <i>Pituitary</i> , 2021, 24, 143-145.	2.9	24
69	Familial isolated pituitary adenomas experience at a single center: clinical importance of AIP mutation screening. <i>Arquivos Brasileiros De Endocrinologia E Metabologia</i> , 2010, 54, 698-704.	1.3	23
70	Growth of an aggressive tumor during pregnancy in an acromegalic patient. <i>Endocrine Journal</i> , 2012, 59, 313-319.	1.6	23
71	A paradigm shift in the medical treatment of acromegaly: from a "trial and error" to a personalized therapeutic decision-making process. <i>Clinical Endocrinology</i> , 2015, 83, 1-2.	2.4	23
72	Cavernous carotid artery pseudo-aneurysm treated by stenting in acromegalic patient. <i>Arquivos De Neuro-Psiquiatria</i> , 2003, 61, 459-462.	0.8	21

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73	The Role of Temozolomide in the Treatment of a Patient With a Pure Silent Pituitary Somatotroph Carcinoma. <i>Endocrine Practice</i> , 2013, 19, e145-e149.	2.1	21
74	Treatment escape reduces the effectiveness of cabergoline during long-term treatment of acromegaly in monotherapy or in association with first-generation somatostatin receptor ligands. <i>Clinical Endocrinology</i> , 2018, 88, 889-895.	2.4	21
75	Management of pituitary incidentaloma. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2019, 33, 101268.	4.7	21
76	Clinical significance of filamin A in patients with acromegaly and its association with somatostatin and dopamine receptor profiles. <i>Scientific Reports</i> , 2019, 9, 1122.	3.3	21
77	Evaluation of the Efficacy and Safety of Switching to Pasireotide in Patients With Acromegaly Inadequately Controlled With First-Generation Somatostatin Analogs. <i>Frontiers in Endocrinology</i> , 2019, 10, 931.	3.5	21
78	Acromegalic patients lost to follow-up: a pilot study. <i>Pituitary</i> , 2013, 16, 245-250.	2.9	20
79	A review on the diagnosis and treatment of patients with clinically nonfunctioning pituitary adenoma by the Neuroendocrinology Department of the Brazilian Society of Endocrinology and Metabolism. <i>Archives of Endocrinology and Metabolism</i> , 2016, 60, 374-390.	0.6	20
80	Balance Control and Peripheral Muscle Function in Aging: A Comparison Between Individuals with Acromegaly and Healthy Subjects. <i>Journal of Aging and Physical Activity</i> , 2017, 25, 218-227.	1.0	20
81	Utility of [18F] fluoro-2-deoxy-d-glucose positron emission tomography in the localization of ectopic ACTH-secreting tumors. <i>Pituitary</i> , 2009, 12, 380-383.	2.9	19
82	Experience with pegvisomant treatment in acromegaly in a single Brazilian tertiary reference center: efficacy, safety and predictors of response. <i>Archives of Endocrinology and Metabolism</i> , 2016, 60, 479-485.	0.6	19
83	On the Functional Capacity and Quality of Life of Patients with Acromegaly: Are They Candidates for Rehabilitation Programs?. <i>Journal of Physical Therapy Science</i> , 2013, 25, 1497-1501.	0.6	18
84	Posture and balance control in patients with acromegaly: Results of a cross-sectional study. <i>Gait and Posture</i> , 2014, 40, 154-159.	1.4	18
85	AIP mutations in Brazilian patients with sporadic pituitary adenomas: a single-center evaluation. <i>Endocrine Connections</i> , 2017, 6, 914-925.	1.9	18
86	Molecular evidence and clinical importance of β -arrestins expression in patients with acromegaly. <i>Journal of Cellular and Molecular Medicine</i> , 2018, 22, 2110-2116.	3.6	18
87	Expression of Retinoblastoma Protein in Human Growth Hormone-Secreting Pituitary Adenomas. <i>Endocrine Pathology</i> , 2005, 16, 053-062.	9.0	17
88	Osteosarcoma and acromegaly: A case report and review of the literature. <i>Journal of Endocrinological Investigation</i> , 2006, 29, 1006-1011.	3.3	17
89	Optic pathways tuberculoma mimicking glioma: case report. <i>World Neurosurgery</i> , 2003, 60, 349-353.	1.3	16
90	Sellar and suprasellar mixed germ cell tumor mimicking a pituitary adenoma. <i>Pituitary</i> , 2011, 14, 345-350.	2.9	16

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91	Rotation thromboelastometry and the hypercoagulable state in Cushing's syndrome. <i>Clinical Endocrinology</i> , 2014, 81, 657-664.	2.4	16
92	Brazilian multicenter study on pegvisomant treatment in acromegaly. <i>Archives of Endocrinology and Metabolism</i> , 2019, 63, 328-336.	0.6	16
93	Accuracy of microcystic aspect on T2-weighted MRI for the diagnosis of silent corticotroph adenomas. <i>Clinical Endocrinology</i> , 2020, 92, 145-149.	2.4	16
94	Pulmonary function testing and chest tomography in patients with acromegaly. <i>Multidisciplinary Respiratory Medicine</i> , 2013, 8, 70.	1.5	14
95	Recommendations of the Neuroendocrinology Department of the Brazilian Society of Endocrinology and Metabolism for the diagnosis of Cushing's disease in Brazil. <i>Archives of Endocrinology and Metabolism</i> , 2016, 60, 267-286.	0.6	14
96	Long-Term Remission of Acromegaly after Octreotide Withdrawal Is an Uncommon and Frequently Unsustainable Event. <i>Neuroendocrinology</i> , 2017, 104, 273-279.	2.5	14
97	Acromegaly. <i>Endocrinology and Metabolism Clinics of North America</i> , 2020, 49, 475-486.	3.2	14
98	Resistance to octreotide LAR in acromegalic patients with high SSTR2 expression: analysis of AIP expression. <i>Arquivos Brasileiros De Endocrinologia E Metabologia</i> , 2012, 56, 501-506.	1.3	13
99	Etiologic aspects and management of acromegaly. <i>Arquivos Brasileiros De Endocrinologia E Metabologia</i> , 2005, 49, 626-640.	1.3	12
100	The NETting of pituitary adenoma: a gland illusion. <i>Pituitary</i> , 2022, 25, 349-351.	2.9	12
101	Growth hormone isoforms in acromegalic patients before and after treatment with octreotide LAR. <i>Growth Hormone and IGF Research</i> , 2010, 20, 87-92.	1.1	11
102	What is the effect of peripheral muscle fatigue, pulmonary function, and body composition on functional exercise capacity in acromegalic patients?. <i>Journal of Physical Therapy Science</i> , 2015, 27, 719-724.	0.6	11
103	Physical exercise improves functional capacity and quality of life in patients with acromegaly: a 12-week follow-up study. <i>Endocrine</i> , 2019, 66, 301-309.	2.3	11
104	Collision sellar lesions: coexistence of pituitary adenoma and Rathke cleft cyst—a single-center experience. <i>Endocrine</i> , 2020, 68, 174-181.	2.3	11
105	Approach to the Patient: Differential Diagnosis of Cystic Sellar Lesions. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 1751-1758.	3.6	11
106	Insulin-like growth factor (IGF)-I, IGF binding protein-3, and prostate cancer: correlation with gleason score. <i>International Braz J Urol: Official Journal of the Brazilian Society of Urology</i> , 2015, 41, 110-115.	1.5	10
107	gsp Mutation Is Not a Molecular Biomarker of Long-Term Response to First-Generation Somatostatin Receptor Ligands in Acromegaly. <i>Cancers</i> , 2021, 13, 4857.	3.7	10
108	A Subcutaneous Octreotide Hydrogel Implant for the Treatment of Acromegaly. <i>Endocrine Practice</i> , 2012, 18, 870-881.	2.1	9

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109	Somatotropinomas inadequately controlled with octreotide may over-respond to pasireotide: the importance of dose adjustment to achieve long-term biochemical control. <i>Hormones</i> , 2017, 16, 84-91.	1.9	9
110	Cyclin A in nonfunctioning pituitary adenomas. <i>Endocrine</i> , 2020, 70, 380-387.	2.3	8
111	Novel therapies for acromegaly. <i>Endocrine Connections</i> , 2020, 9, R274-R285.	1.9	8
112	Prolactinomas. <i>Presse Medicale</i> , 2021, 50, 104080.	1.9	8
113	Acromegaly and Non-Hodgkin's Lymphoma. <i>Endocrine Practice</i> , 1998, 4, 279-281.	2.1	7
114	Tumor Deletion Mapping of Chromosomal Region 13q14 in 43 Growth Hormone Secreting Pituitary Adenomas. <i>Endocrine</i> , 2005, 28, 131-136.	2.2	7
115	Hematologic neoplasias and acromegaly. <i>Pituitary</i> , 2011, 14, 377-381.	2.9	7
116	Computed tomography airway lumen volumetry in patients with acromegaly: Association with growth hormone levels and lung function. <i>Journal of Medical Imaging and Radiation Oncology</i> , 2017, 61, 591-599.	1.8	7
117	Pituitary MRI Features in Acromegaly Resulting From Ectopic GHRH Secretion From a Neuroendocrine Tumor: Analysis of 30 Cases. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e3313-e3320.	3.6	7
118	New and emerging pharmacological treatment options for acromegaly. <i>Expert Opinion on Pharmacotherapy</i> , 2021, 22, 1615-1623.	1.8	6
119	Identification of mutant K-RAS in pituitary macroadenoma. <i>Pituitary</i> , 2021, 24, 746-753.	2.9	6
120	Pituitary MRI Standard and Advanced Sequences: Role in the Diagnosis and Characterization of Pituitary Adenomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 1431-1440.	3.6	6
121	Management of hypopituitarism: a perspective from the Brazilian Society of Endocrinology and Metabolism. <i>Archives of Endocrinology and Metabolism</i> , 2021, 65, 212-230.	0.6	5
122	Growth hormone receptor exon 3 isoforms may have no importance in the clinical setting of multiethnic Brazilian acromegaly patients. <i>Pituitary</i> , 2016, 19, 375-380.	2.9	4
123	The effectiveness of a therapist-oriented home rehabilitation program for a patient with acromegaly: A case study. <i>Journal of Bodywork and Movement Therapies</i> , 2019, 23, 634-642.	1.2	4
124	The Glittre Activities of Daily Living Test in patients with acromegaly: Associations with hand function and health-related quality of life. <i>Journal of Back and Musculoskeletal Rehabilitation</i> , 2021, 34, 441-451.	1.1	4
125	Osilodrostat for the treatment of Cushing's disease: efficacy, stability, and persistence – Authors' reply. <i>Lancet Diabetes and Endocrinology</i> , 2022, 10, 385-387.	11.4	4
126	A review of Cushing's disease treatment by the Department of Neuroendocrinology of the Brazilian Society of Endocrinology and Metabolism. <i>Archives of Endocrinology and Metabolism</i> , 2018, 62, 87-105.	0.6	3

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127	Letter to the Editor: "Our Response to COVID-19 as Endocrinologists and Diabetologists" Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2661-e2662.	3.6	3
128	GH and IGF-I levels and tumor shrinkage in response to first generation somatostatin receptor ligands in acromegaly: a comparative study between two reference centers for pituitary diseases in Brazil. Endocrine, 2021, 74, 146-154.	2.3	3
129	Safety and Efficacy of Switching Injected Peptide Long-Acting Somatostatin Receptor Ligands to Once Daily Oral Paltusotine: ACROBAT Edge Phase 2 Study. Journal of the Endocrine Society, 2021, 5, A526-A527.	0.2	3
130	Telomerase expression in clinically non-functioning pituitary adenomas. Endocrine, 2021, 72, 208-215.	2.3	2
131	Growth hormone-releasing hormone-secreting pulmonary neuroendocrine tumor associated with pituitary hyperplasia and somatotropinoma. Archives of Endocrinology and Metabolism, 2021, 65, 648-663.	0.6	2
132	Current opinion on the diagnosis and management of non-functioning pituitary adenomas. Expert Review of Endocrinology and Metabolism, 2021, 16, 309-320.	2.4	2
133	Germ cell tumor presenting as sellar mass with suprasellar extension and long history of hypopituitarism. Neuroendocrinology Letters, 2010, 31, 306-9.	0.2	2
134	Authors'™ Response: Isolated Familial Somatotropinomas: Does the Disease Map to 11q13 or to 2p16?. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 4921-4921.	3.6	1
135	Cyclic ACTH-secreting thymic carcinoid: a case report and review of the literature. Archives of Endocrinology and Metabolism, 2021, 65, 512-516.	0.6	1
136	Apoplexy in sporadic pituitary adenomas: a single referral center experience and AIP mutation analysis. Archives of Endocrinology and Metabolism, 2021, 65, 295-304.	0.6	1
137	Current reliability of the Immulite® assay for measurement of serum IGF-1 in the Brazilian adult population. Archives of Endocrinology and Metabolism, 2015, 59, 195-196.	0.6	1
138	SAT-433 Long-Acting Pasireotide Provides Clinical Benefit to Patients with Uncontrolled Acromegaly over Continued Treatment with First-Generation Somatostatin Analogues (SSAs): Results from Phase 3b, Open-Label Study. Journal of the Endocrine Society, 2019, 3, .	0.2	1
139	Evidence-based guidelines in acromegaly: implications on the clinic. Expert Review of Endocrinology and Metabolism, 2016, 11, 171-175.	2.4	0
140	Clinical and functional variables can predict general fatigue in patients with acromegaly: an explanatory model approach. Archives of Endocrinology and Metabolism, 2019, 63, 235-240.	0.6	0