## Henrik Falhammar

List of Publications by Year in descending order

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207 papers 6,747 citations

41 h-index 70 g-index

223 all docs 223 docs citations

times ranked

223

5635 citing authors

#	Article	IF	Citations
1	Current and Novel Treatment Strategies in Children with Congenital Adrenal Hyperplasia. Hormone Research in Paediatrics, 2023, 96, 560-572.	0.8	10
2	Congenital Adrenal Hyperplasiaâ€"Current Insights in Pathophysiology, Diagnostics, and Management. Endocrine Reviews, 2022, 43, 91-159.	8.9	182
3	Characteristics and health outcomes of patients hospitalised with hypomagnesaemia: a retrospective study from a single centre in the Northern Territory of Australia. Internal Medicine Journal, 2022, 52, 1544-1553.	0.5	4
4	Increased Prevalence of Fractures in Congenital Adrenal Hyperplasia: A Swedish Population-based National Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e475-e486.	1.8	17
5	Pressure relieving interventions for the management of diabetesâ€related foot ulcers: a study from the Northern Territory of Australia. ANZ Journal of Surgery, 2022, , .	0.3	4
6	The impact of adherence and therapy regimens on quality of life in patients with congenital adrenal hyperplasia. Clinical Endocrinology, 2022, 96, 666-679.	1.2	5
7	Comparison and outcomes of emergency department presentations with respiratory disorders among Australian indigenous and non-indigenous patients. BMC Emergency Medicine, 2022, 22, 11.	0.7	14
8	Pheochromocytomas and Abdominal Paragangliomas: A Practical Guidance. Cancers, 2022, 14, 917.	1.7	16
9	Adrenal trauma <i>experience </i> at a major tertiary centre in Sweden: Clinical and radiological findings. Clinical Endocrinology, 2022, 97, 28-35.	1.2	4
10	Current and Future Burdens of Heat-Related Hyponatremia: A Nationwide Register–Based Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2388-e2393.	1.8	5
11	Cytoâ€morphological features of parathyroid lesions: Fineâ€needle aspiration cytology series from an endocrine tumor referral center. Diagnostic Cytopathology, 2022, 50, 75-83.	0.5	6
12	CAH-X Syndrome: Genetic and Clinical Profile. Molecular Diagnosis and Therapy, 2022, 26, 293-300.	1.6	5
13	Age-dependent and sex-dependent disparity in mortality in patients with adrenal incidentalomas and autonomous cortisol secretion: an international, retrospective, cohort study. Lancet Diabetes and Endocrinology,the, 2022, 10, 499-508.	5.5	55
14	Prevalence and incidence of diabetes among Aboriginal people in remote communities of the Northern Territory, Australia: a retrospective, longitudinal data-linkage study. BMJ Open, 2022, 12, e059716.	0.8	17
15	Adrenal crises in adolescents and young adults. Endocrine, 2022, 77, 1-10.	1.1	9
16	Long-Term Outcomes of Congenital Adrenal Hyperplasia. Endocrinology and Metabolism, 2022, 37, 587-598.	1.3	13
17	Reproductive and Perinatal Outcomes in Women with Congenital Adrenal Hyperplasia: A Population-based Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e957-e965.	1.8	27
18	Ectopic ACTH- and/or CRH-Producing Pheochromocytomas. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 598-608.	1.8	19

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19	Reduced risk for hospitalization due to hyponatraemia in lithium treated patients: A Swedish population-based case–control study. Journal of Psychopharmacology, 2021, 35, 184-189.	2.0	6
20	Top End Pulmonary Hypertension Study: Understanding Epidemiology, Therapeutic Gaps and Prognosis in Remote Australian Setting. Heart Lung and Circulation, 2021, 30, 507-515.	0.2	1
21	Association between lipid-lowering agents and severe hyponatremia: a population-based case–control study. European Journal of Clinical Pharmacology, 2021, 77, 747-755.	0.8	2
22	11C-Metomidate PET/CT Detected Multiple Ectopic Adrenal Rest Tumors in a Woman With Congenital Adrenal Hyperplasia. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e675-e679.	1.8	13
23	Association between newly initiated thiazide diuretics and hospitalization due to hyponatremia. European Journal of Clinical Pharmacology, 2021, 77, 1049-1055.	0.8	13
24	Metastatic Pheochromocytomas and Abdominal Paragangliomas. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e1937-e1952.	1.8	41
25	Bone Mass in Young Patients with Congenital Adrenal Hyperplasia due to 21-Hydroxylase Deficiency. Hormone Research in Paediatrics, 2021, 94, 1-8.	0.8	7
26	Magnesium: The recent research and developments. Advances in Food and Nutrition Research, 2021, 96, 193-218.	1.5	9
27	Assessment of medication adherence in children and adults with congenital adrenal hyperplasia and the impact of knowledge and selfâ€management. Clinical Endocrinology, 2021, 94, 753-764.	1.2	6
28	Genotype-Phenotype Correlation in Patients with Congenital Adrenal Hyperplasia due to 21-Hydroxylase Deficiency in Cuba. International Journal of Endocrinology, 2021, 2021, 1-6.	0.6	3
29	Clinical outcomes in 21-hydroxylase deficiency. Current Opinion in Endocrinology, Diabetes and Obesity, 2021, 28, 318-324.	1.2	10
30	Use of medical identification jewellery in children and young adults with adrenal insufficiency in Australia. Endocrine, 2021, 72, 539-545.	1.1	7
31	First insights into the genetics of 21â€hydroxylase deficiency in the Roma population. Clinical Endocrinology, 2021, 95, 41-46.	1.2	4
32	Quality of Life in Men With Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency. Frontiers in Endocrinology, 2021, 12, 626646.	1.5	8
33	Therapy options for adrenal insufficiency and recommendations for the management of adrenal crisis. Endocrine, 2021, 71, 586-594.	1.1	31
34	MANAGEMENT OF ENDOCRINE DISEASE: Gonadal dysfunction in congenital adrenal hyperplasia. European Journal of Endocrinology, 2021, 184, R85-R97.	1.9	38
35	Time-dependent association between selective serotonin reuptake inhibitors and hospitalization due to hyponatremia. Journal of Psychopharmacology, 2021, 35, 928-933.	2.0	7
36	Characteristics, Treatment, Outcomes, and Survival in Neuroendocrine G1 and G2 Pancreatic Tumors: Experiences From a Single Tertiary Referral Center. Frontiers in Endocrinology, 2021, 12, 657698.	1.5	3

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37	Clinical and Hormonal Profiles Correlate With Molecular Characteristics in Patients With 11β-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3714-e3724.	1.8	20
38	Rate of fracture in patients with glucocorticoid replacement therapy: a systematic review and meta-analysis. Endocrine, 2021, 74, 29-37.	1.1	17
39	Pituitary Apoplexy: A Retrospective Study of 33 Cases From a Single Center. Frontiers in Endocrinology, 2021, 12, 656950.	1.5	12
40	Institutional characterisation of water clear cell parathyroid adenoma: a rare entity often unrecognised by TC-99m-sestamibi scintigraphy. Pathology, 2021, 53, 852-859.	0.3	6
41	Prevalence of Nelson's syndrome after bilateral adrenalectomy in patients with cushing's disease: a systematic review and meta-analysis. Pituitary, 2021, 24, 797-809.	1.6	9
42	Nonâ€thiazide diuretics and hospitalization due to hyponatraemia: A populationâ€based caseâ€control study. Clinical Endocrinology, 2021, 95, 520-526.	1.2	2
43	Metastasis to the thyroid gland: Characterization and survival of an institutional series spanning 28 years. European Journal of Surgical Oncology, 2021, 47, 1364-1369.	0.5	15
44	The epidemiology of primary and secondary adrenal malignancies and associated adrenal insufficiency in hospitalised patients: an analysis of hospital admission data, NSW, Australia. BMC Endocrine Disorders, 2021, 21, 141.	0.9	4
45	Adrenal myelolipomas. Lancet Diabetes and Endocrinology, the, 2021, 9, 767-775.	5.5	49
46	Thyroid testing paradigm switch from thyrotropin to thyroid hormonesâ€"Future directions and opportunities in clinical medicine and research. Endocrine, 2021, 74, 285-289.	1.1	7
47	Orthostatic intolerance after bariatric surgery: A systematic review and metaâ€analysis. Clinical Obesity, 2021, 11, e12483.	1.1	4
48	Reduced expression of OXPHOS and DNA damage genes is linked to protection from microvascular complications in long-term type $1$ diabetes: the PROLONG study. Scientific Reports, 2021, $11$ , 20735.	1.6	7
49	Characteristics of In2G Variant in Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency. Frontiers in Endocrinology, 2021, 12, 788812.	1.5	4
50	Factors of importance for discontinuation of thiazides associated with hyponatremia in Sweden: A populationâ€based register study. Pharmacoepidemiology and Drug Safety, 2020, 29, 77-83.	0.9	3
51	Inverse association between glucose-lowering medications and severe hyponatremia: a Swedish population-based case-control study. Endocrine, 2020, 67, 579-586.	1.1	6
52	Stumbling broke the spleen and unveiled pheochromocytoma, which in turn broke the heart. Endocrine, 2020, 67, 727-728.	1.1	5
53	Bone mineral density and fractures in congenital adrenal hyperplasia: Findings from the dsdâ€LIFE study. Clinical Endocrinology, 2020, 92, 284-294.	1.2	29
54	P450 Oxidoreductase Deficiency: A Systematic Review and Meta-analysis of Genotypes, Phenotypes, and Their Relationships. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e42-e52.	1.8	19

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55	Diabetes during pregnancy and birthweight trends among Aboriginal and non-Aboriginal people in the Northern Territory of Australia over 30 years. The Lancet Regional Health - Western Pacific, 2020, 1, 100005.	1.3	14
56	Obstructive sleep apnoea and adherence to continuous positive airway therapy among Australian women. Internal Medicine Journal, 2020, , .	0.5	12
57	Riedel Thyroiditis. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e3469-e3481.	1.8	18
58	Liver nucleotide biosynthesis is linked to protection from vascular complications in individuals with long-term type 1 diabetes. Scientific Reports, 2020, 10, 11561.	1.6	8
59	Genetic and Biological Effects of ICAM-1 E469K Polymorphism in Diabetic Kidney Disease. Journal of Diabetes Research, 2020, 2020, 1-7.	1.0	5
60	Molecular diagnosis of patients with congenital adrenal hyperplasia due to 21-hydroxylase deficiency. BMC Endocrine Disorders, 2020, 20, 165.	0.9	6
61	Bone Mineral Density in Adults With Congenital Adrenal Hyperplasia: A Systematic Review and Meta-Analysis. Frontiers in Endocrinology, 2020, 11, 493.	1.5	32
62	Cardiovascular Manifestations and Complications of Pheochromocytomas and Paragangliomas. Journal of Clinical Medicine, 2020, 9, 2435.	1.0	54
63	Predictors of normalized HbA1c after gastric bypass surgery in subjects with abnormal glucose levels, a 2-year follow-up study. Scientific Reports, 2020, 10, 15127.	1.6	3
64	All-cause mortality following low-dose aspirin treatment for patients with high cardiovascular risk in remote Australian Aboriginal communities: an observational study. BMJ Open, 2020, 10, e030034.	0.8	6
65	Highly proliferative anal neuroendocrine carcinoma: molecular and clinical features of a rare, recurrent case in complete remission. BMC Gastroenterology, 2020, 20, 290.	0.8	4
66	The Success of a Screening Program Is Largely Dependent on Close Collaboration between the Laboratory and the Clinical Follow-Up of the Patients. International Journal of Neonatal Screening, 2020, 6, 68.	1.2	13
67	Update on the Swedish Newborn Screening for Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency. International Journal of Neonatal Screening, 2020, 6, 71.	1.2	19
68	Increased Plasma Soluble Interleukin-2 Receptor Alpha Levels in Patients With Long-Term Type 1 Diabetes With Vascular Complications Associated With IL2RA and PTPN2 Gene Polymorphisms. Frontiers in Endocrinology, 2020, 11, 575469.	1.5	4
69	Clinical Parameters Are More Likely to Be Associated with Thyroid Hormone Levels than with Thyrotropin Levels: A Systematic Review and Meta-Analysis. Thyroid, 2020, 30, 1695-1709.	2.4	73
70	Clinical outcomes and characteristics of P30L mutations in congenital adrenal hyperplasia due to 21-hydroxylase deficiency. Endocrine, 2020, 69, 262-277.	1.1	13
71	Current Management and Outcome of Pregnancies in Women With Adrenal Insufficiency: Experience from a Multicenter Survey. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2853-e2863.	1.8	30
72	Mucormycosis in a 40-year-old woman with diabetic ketoacidosis. Cmaj, 2020, 192, E431-E433.	0.9	8

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73	Lactation Ketoacidosis: A Systematic Review of Case Reports. Medicina (Lithuania), 2020, 56, 299.	0.8	6
74	MON-183 Adrenal Androgen Control and Steroidal Side Effects in Adolescents and Adults with Congenital Adrenal Hyperplasia Treated with Glucocorticoids. Journal of the Endocrine Society, 2020, 4, .	0.1	0
75	Lipoadenoma of the Parathyroid Gland: Characterization of an Institutional Series Spanning 28ÂYears. Endocrine Pathology, 2020, 31, 156-165.	<b>5.</b> 2	13
76	Clinical features, complications, and outcomes of exogenous and endogenous catecholamineâ€triggered Takotsubo syndrome: A systematic review and metaâ€analysis of 156 published cases. Clinical Cardiology, 2020, 43, 459-467.	0.7	30
77	Authors' Response to Hennessey re: DOI: 10.1089/thy.2019.0535. Thyroid, 2020, 30, 1835-1836.	2.4	2
78	Adrenal crises in older patients. Lancet Diabetes and Endocrinology, the, 2020, 8, 628-639.	5 <b>.</b> 5	28
79	To Treat or Not to Treat Subclinical Hypothyroidism, What Is the Evidence?. Medicina (Lithuania), 2020, 56, 40.	0.8	25
80	Prevalence and Characteristics of Adrenal Tumors and Myelolipomas in Congenital Adrenal Hyperplasia: A Systematic Review and Meta-Analysis. Endocrine Practice, 2020, 26, 1351-1365.	1.1	37
81	Associations Between Antihypertensive Medications and Severe Hyponatremia: A Swedish Population–Based Case–Control Study. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e3696-e3705.	1.8	16
82	The effect of patientâ€managed stress dosing on electrolytes and blood pressure in acute illness in children with adrenal insufficiency. Clinical Endocrinology, 2020, 93, 97-103.	1.2	5
83	Sexual Orientation in Individuals With Congenital Adrenal Hyperplasia: A Systematic Review. Frontiers in Behavioral Neuroscience, 2020, 14, 38.	1.0	27
84	Lower extremity amputations and long-term outcomes in diabetic foot ulcers: A systematic review. World Journal of Diabetes, 2020, 11, 391-399.	1.3	34
85	OR25-05 Increased Overall Mortality and Cardiovascular Morbidity in Patients with Adrenal Incidentalomas and Autonomous Cortisol Secretion: Results of the ENS@T NAPACA-Outcome Study. Journal of the Endocrine Society, 2020, 4, .	0.1	0
86	Multiple cutaneous lesions and pulmonary cysts. European Journal of Internal Medicine, 2020, 76, 95-96.	1.0	O
87	Associations of proton pump inhibitors and hospitalization due to hyponatremia: A population–based case–control study. European Journal of Internal Medicine, 2019, 59, 65-69.	1.0	38
88	Pheochromocytoma- and paraganglioma-triggered Takotsubo syndrome. Endocrine, 2019, 65, 483-493.	1.1	38
89	Presentation, Treatment, Histology, and Outcomes in Adrenal Medullary Hyperplasia Compared With Pheochromocytoma. Journal of the Endocrine Society, 2019, 3, 1518-1530.	0.1	16
90	Sexuality in Males With Congenital Adrenal Hyperplasia Resulting From 21-Hydroxylase Deficiency. Journal of the Endocrine Society, 2019, 3, 1445-1456.	0.1	9

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91	Amputations in patients with diabetic foot ulcer: a retrospective study from a single centre in the Northern Territory of Australia. ANZ Journal of Surgery, 2019, 89, 874-879.	0.3	18
92	Population data provide evidence against the presence of a set point for hemoglobin levels or tissue oxygen delivery. Physiological Reports, 2019, 7, e14153.	0.7	5
93	Sex-specific risks of death in patients hospitalized for hyponatremia: a population-based study. Endocrine, 2019, 66, 660-665.	1.1	13
94	Genome-Wide Association Study of Diabetic Kidney Disease Highlights Biology Involved in Glomerular Basement Membrane Collagen. Journal of the American Society of Nephrology: JASN, 2019, 30, 2000-2016.	3.0	135
95	Pyogenic hepatic abscess secondary to gastric perforation caused by an ingested fish bone. Medical Journal of Australia, 2019, 211, 451.	0.8	5
96	Carriers of a Classic CYP21A2 Mutation Have Reduced Mortality: A Population-Based National Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 6148-6154.	1.8	10
97	Adrenal Crisis. New England Journal of Medicine, 2019, 381, 852-861.	13.9	144
98	Increased Risk of Autoimmune Disorders in 21-Hydroxylase Deficiency: A Swedish Population-Based National Cohort Study. Journal of the Endocrine Society, 2019, 3, 1039-1052.	0.1	8
99	Glucocorticoid Regimens in the Treatment of Congenital Adrenal Hyperplasia: A Systematic Review and Meta-Analysis. Journal of the Endocrine Society, 2019, 3, 1227-1245.	0.1	48
100	Protective Effect of the <i>HIF-1A</i> Pro582Ser Polymorphism on Severe Diabetic Retinopathy. Journal of Diabetes Research, 2019, 2019, 1-8.	1.0	22
101	Adrenal insufficiency due to bilateral adrenal metastases – A systematic review and meta-analysis. Heliyon, 2019, 5, e01783.	1.4	25
102	Epidemiology of Pulmonary Hypertension at the Top End of Australia. Heart Lung and Circulation, 2019, 28, S65.	0.2	0
103	New cost-effective pleural procedure training: manikin-based model to increase the confidence and competency in trainee medical officers. Postgraduate Medical Journal, 2019, 95, 245-250.	0.9	2
104	Clinical perspectives in congenital adrenal hyperplasia due to $3\hat{l}^2$ -hydroxysteroid dehydrogenase type 2 deficiency. Endocrine, 2019, 63, 407-421.	1.1	54
105	Tramadol- and codeine-induced severe hyponatremia: A Swedish population-based case-control study. European Journal of Internal Medicine, 2019, 69, 20-24.	1.0	20
106	Adrenal Crisis. New England Journal of Medicine, 2019, 381, 2181-2183.	13.9	3
107	Acute suppurative thyroiditis with thyroid abscess in adults: clinical presentation, treatment and outcomes. BMC Endocrine Disorders, 2019, 19, 130.	0.9	41
108	Mortality in patients with diabetic foot ulcer: a retrospective study of 513 cases from a single Centre in the Northern Territory of Australia. BMC Endocrine Disorders, 2019, 19, 1.	0.9	140

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109	Antipsychotics and severe hyponatremia: A Swedish population–based case–control study. European Journal of Internal Medicine, 2019, 60, 71-77.	1.0	32
110	MANAGEMENT OF ENDOCRINE DISEASE: Diagnosis and management of the patient with non-classic CAH due to 21-hydroxylase deficiency. European Journal of Endocrinology, 2019, 180, R127-R145.	1.9	103
111	Lactation Ketoacidosis: A case series. Sultan Qaboos University Medical Journal, 2019, 19, 359.	0.3	6
112	11β-Hydroxylase Deficiency. , 2019, , 421-430.		2
113	Health status in 1040 adults with disorders of sex development (DSD): a European multicenter study. Endocrine Connections, 2018, 7, 466-478.	0.8	51
114	Riedel's thyroiditis: clinical presentation, treatment and outcomes. Endocrine, 2018, 60, 185-192.	1.1	38
115	Gonadal function in adult male patients with congenital adrenal hyperplasia. European Journal of Endocrinology, 2018, 178, 285-294.	1.9	57
116	Skeletal fragility induced by overtreatment of adrenal insufficiency. Endocrine, 2018, 59, 239-241.	1.1	6
117	Successful fertility outcome in a woman with 17É'â€hydroxylase deficiency. Clinical Endocrinology, 2018, 88, 607-609.	1.2	16
118	Two rare forms of congenital adrenal hyperplasia, $11\hat{l}^2$ hydroxylase deficiency and 17-hydroxylase/17,20-lyase deficiency, presenting with novel mutations. Hormones, 2018, 17, 127-132.	0.9	5
119	Differences in associations of antiepileptic drugs and hospitalization due to hyponatremia: A population–based case–control study. Seizure: the Journal of the British Epilepsy Association, 2018, 59, 28-33.	0.9	33
120	Initial clinical presentation and spectrum of pheochromocytoma: a study of 94 cases from a single center. Endocrine Connections, 2018, 7, 186-192.	0.8	74
121	Trends in surgery, hospital admissions and imaging for pituitary adenomas in Australia. Endocrine, 2018, 59, 373-382.	1.1	12
122	Differences in Associations of Antidepressants and Hospitalization Due to Hyponatremia. American Journal of Medicine, 2018, 131, 56-63.	0.6	47
123	Lactation ketoacidosis: case presentation and literature review. BMJ Case Reports, 2018, 2018, bcr-2017-223494.	0.2	10
124	Treatment and outcomes in pheochromocytomas and paragangliomas: a study of 110 cases from a single center. Endocrine, 2018, 62, 566-575.	1.1	30
125	Magnesium and Human Health: Perspectives and Research Directions. International Journal of Endocrinology, 2018, 2018, 1-17.	0.6	215
126	A Sickening Tale. New England Journal of Medicine, 2018, 379, 75-80.	13.9	2

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127	Extensive Bilateral Adrenal Rest Testicular Tumors in a Patient With $3\hat{l}^2$ -Hydroxysteroid Dehydrogenase Type 2 Deficiency. Journal of the Endocrine Society, 2018, 2, 513-517.	0.1	10
128	Bilateral Adrenalectomy in Congenital Adrenal Hyperplasia: A Systematic Review and Meta-Analysis. Journal of Clinical Endocrinology and Metabolism, 2018, 103, 1767-1778.	1.8	36
129	Psychological adjustment, quality of life, and self-perceptions of reproductive health in males with congenital adrenal hyperplasia: a systematic review. Endocrine, 2018, 62, 3-13.	1.1	34
130	Variations in the management of acute illness in children with congenital adrenal hyperplasia: An audit of three paediatric hospitals. Clinical Endocrinology, 2018, 89, 577-585.	1.2	20
131	Adrenal Crises in Children: Perspectives and Research Directions. Hormone Research in Paediatrics, 2018, 89, 341-351.	0.8	39
132	Rescue pre-operative treatment with Lugol's solution in uncontrolled Graves' disease. Endocrine Connections, 2017, 6, 200-205.	0.8	11
133	Clinical perspectives in congenital adrenal hyperplasia due to $11\hat{l}^2$ -hydroxylase deficiency. Endocrine, 2017, 55, 19-36.	1.1	99
134	Adrenal crises: perspectives and research directions. Endocrine, 2017, 55, 336-345.	1.1	61
135	Lugol's solution and other iodide preparations: perspectives and research directions in Graves' disease. Endocrine, 2017, 58, 467-473.	1.1	29
136	Hospitalisation in Children with Adrenal Insufficiency and Hypopituitarism: Is There a Differential Burden between Boys and Girls and between Age Groups?. Hormone Research in Paediatrics, 2017, 88, 339-346.	0.8	16
137	Cost-effectiveness of stroke care in Aboriginal and non-Aboriginal patients: an observational cohort study in the Northern Territory of Australia. BMJ Open, 2017, 7, e015033.	0.8	10
138	Increased mortality in patients with adrenal incidentalomas and autonomous cortisol secretion: a 13-year retrospective study from one center. Endocrine, 2017, 58, 267-275.	1.1	76
139	Swyer-James-MacLeod syndrome-a rare diagnosis presented through two adult patients. Respirology Case Reports, 2017, 5, e00245.	0.3	12
140	Dopa-testotoxicosis: disruptive hypersexuality in hypogonadal men with prolactinomas treated with dopamine agonists. Endocrine, 2017, 55, 618-624.	1.1	39
141	Are carriers of <i>CYP21A2</i> mutations less vulnerable to psychological stress? A population-based national cohort study. Clinical Endocrinology, 2017, 86, 317-324.	1.2	8
142	Frequency of Cushing's syndrome due to ACTH-secreting adrenal medullary lesions: a retrospective study over 10 years from a single center. Endocrine, 2017, 55, 296-302.	1.1	30
143	Reduced Frequency of Biological and Increased Frequency of Adopted Children in Males With 21-Hydroxylase Deficiency: A Swedish Population-Based National Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 4191-4199.	1.8	50
144	Pleural Lipomatosis Masquerading as Pleural Mass With Effusion. Chest, 2017, 152, A539.	0.4	1

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145	Hospital Admission Patterns in Children with CAH: Admission Rates and Adrenal Crises Decline with Age. International Journal of Endocrinology, 2016, 2016, 1-7.	0.6	22
146	Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency Presenting as Adrenal Incidentaloma: A Systematic Review and Meta-Analysis. Endocrine Practice, 2016, 22, 736-752.	1.1	43
147	Adrenocortical cancer: mortality, hormone secretion, proliferation and urine steroids – experience from a single centre spanning three decades. BMC Endocrine Disorders, 2016, 16, 15.	0.9	27
148	Relationship between depression and diabetes in pregnancy: A systematic review. World Journal of Diabetes, 2016, 7, 554.	1.3	39
149	Clinical Outcomes in Adrenal Incidentaloma: Experience From one Center. Endocrine Practice, 2015, 21, 870-877.	1.1	64
150	Congenital Adrenal Hyperplasia, Polycystic Ovary Syndrome and criminal behavior: A Swedish population based study. Psychiatry Research, 2015, 229, 953-959.	1.7	12
151	Nonclassic congenital adrenal hyperplasia due to 21-hydroxylase deficiency: clinical presentation, diagnosis, treatment, and outcome. Endocrine, 2015, 50, 32-50.	1.1	93
152	Ileal neuroendocrine tumors and heart: not only valvular consequences. Endocrine, 2015, 48, 743-755.	1.1	9
153	Limited value of long-term biochemical follow-up in patients with adrenal incidentalomas-a retrospective cohort study. BMC Endocrine Disorders, 2015, 15, 6.	0.9	22
154	Congenital adrenal hyperplasia and risk for psychiatric disorders in girls and women born between 1915 and 2010: A total population study. Psychoneuroendocrinology, 2015, 60, 195-205.	1.3	96
155	Increased Cardiovascular and Metabolic Morbidity in Patients With 21-Hydroxylase Deficiency: A Swedish Population-Based National Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2015, 100, 3520-3528.	1.8	153
156	Biochemical and genetic diagnosis of 21-hydroxylase deficiency. Endocrine, 2015, 50, 306-314.	1.1	62
157	Pregnancy and neonatal outcomes in Indigenous Australians with diabetes in pregnancy. World Journal of Diabetes, 2015, 6, 880.	1.3	14
158	Increased Mortality in Patients With Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E2715-E2721.	1.8	138
159	Quality of life, social situation, and sexual satisfaction, in adult males with congenital adrenal hyperplasia. Endocrine, 2014, 47, 299-307.	1.1	34
160	68Ga-DOTA-TOC-PET/CT detects heart metastases from ileal neuroendocrine tumors. Endocrine, 2014, 47, 169-176.	1.1	20
161	Increased Psychiatric Morbidity in Men With Congenital Adrenal Hyperplasia due to 21-Hydroxylase Deficiency. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E554-E560.	1.8	78
162	Suboptimal Psychosocial Outcomes in Patients With Congenital Adrenal Hyperplasia: Epidemiological Studies in a Nonbiased National Cohort in Sweden. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 1425-1432.	1.8	86

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163	GP4-3: Epigenetic analyses of the insulin-like growth factor binding protein 1 gene in diabetes and diabetic nephropathy. Growth Hormone and IGF Research, 2014, 24, S48.	0.5	0
164	Epigenetic analyses of the insulin-like growth factor binding protein 1 gene in type 1 diabetes and diabetic nephropathy. Clinical Epigenetics, 2014, $6$ , $10$ .	1.8	45
165	Non-functioning adrenal incidentalomas caused by 21-hydroxylase deficiency or carrier status?. Endocrine, 2014, 47, 308-314.	1.1	31
166	Thyrotoxic periodic paralysis: clinical and molecular aspects. Endocrine, 2013, 43, 274-84.	1.1	33
167	One hundred years of congenital adrenal hyperplasia in Sweden: a retrospective, population-based cohort study. Lancet Diabetes and Endocrinology,the, 2013, 1, 35-42.	5.5	141
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