

Fady Hannah-Shmouni

List of Publications by Year in descending order

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Version: 2024-02-01

108
papers

1,673
citations

430874

18
h-index

345221

36
g-index

110
all docs

110
docs citations

110
times ranked

2432
citing authors

#	ARTICLE	IF	CITATIONS
1	Scoping review of COVID-19-related systematic reviews and meta-analyses: can we really have confidence in their results?. <i>Postgraduate Medical Journal</i> , 2022, 98, 372-379.	1.8	5
2	Hypertension and COVID-19: Updates from the era of vaccines and variants. <i>Journal of Clinical and Translational Endocrinology</i> , 2022, 27, 100285.	1.4	9
3	eP235: Interim results of the Vigilant observational study: clinical characteristics of creatine transporter deficiency. <i>Genetics in Medicine</i> , 2022, 24, S149.	2.4	0
4	Neurofibromatosis Type 1 Has a Wide Spectrum of Growth Hormone Excess. <i>Journal of Clinical Medicine</i> , 2022, 11, 2168.	2.4	6
5	USP13 genetics and expression in a family with thyroid cancer. <i>Endocrine</i> , 2022, 77, 281-290.	2.3	5
6	Update on the Genetics of Primary Aldosteronism and Aldosterone-Producing Adenomas. <i>Current Cardiology Reports</i> , 2022, 24, 1189-1195.	2.9	5
7	Management of mitochondrial diabetes in the era of novel therapies. <i>Journal of Diabetes and Its Complications</i> , 2021, 35, 107584.	2.3	25
8	Oxidative phosphorylation in creatine transporter deficiency. <i>NMR in Biomedicine</i> , 2021, 34, e4419.	2.8	4
9	Volumetric Modeling of Adrenal Gland Size in Primary Bilateral Macronodular Adrenocortical Hyperplasia. <i>Journal of the Endocrine Society</i> , 2021, 5, bvaa162.	0.2	7
10	First Somatic <i>PRKAR1A</i> Defect Associated With Mosaicism for Another <i>PRKAR1A</i> Mutation in a Patient With Cushing Syndrome. <i>Journal of the Endocrine Society</i> , 2021, 5, bvab007.	0.2	3
11	GPR101, an orphan G-protein coupled receptor, with roles in growth, puberty, and possibly appetite regulation. , 2021, , 79-88.		0
12	Cushing Syndrome in a Pediatric Patient With a <i>KCNJ5</i> Variant and Successful Treatment With Low-dose Ketoconazole. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, 1606-1616.	3.6	4
13	A case of Carney triad complicated by renal cell carcinoma and a germline <i>SDHA</i> pathogenic variant. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2021, 2021, .	0.5	2
14	Molecular Genetic and Genomic Alterations in Cushing's Syndrome and Primary Aldosteronism. <i>Frontiers in Endocrinology</i> , 2021, 12, 632543.	3.5	19
15	A Case of Carney Triad Complicated by Renal Cell Carcinoma and a Germline <i>SDHA</i> Pathogenic Variant. <i>Journal of the Endocrine Society</i> , 2021, 5, A985-A985.	0.2	0
16	Abnormal Pituitary Imaging and Associated Endocrine Dysfunctions in Erdheim-Chester Disease. <i>Journal of the Endocrine Society</i> , 2021, 5, A622-A622.	0.2	1
17	Contralateral Suppression Index Does Not Predict Clinical Cure in Patients Undergoing Surgery for Primary Aldosteronism. <i>Annals of Surgical Oncology</i> , 2021, 28, 7487-7495.	1.5	12
18	X-linked creatine transporter deficiency results in prolonged QTc and increased sudden death risk in humans and disease model. <i>Genetics in Medicine</i> , 2021, 23, 1864-1872.	2.4	8

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19	Homozygous <i>SHBG</i> Variant (<i>rs6258</i>) Linked to Gonadotropin-Independent Precocious Puberty in a Young Girl. <i>Journal of the Endocrine Society</i> , 2021, 5, bvab125.	0.2	0
20	Pituitary Imaging Abnormalities and Related Endocrine Disorders in Erdheim-Chester Disease. <i>Cancers</i> , 2021, 13, 4126.	3.7	4
21	Dual molecular diagnoses in a neurometabolic specialty clinic. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 766-773.	1.2	8
22	Inherited Neuroendocrine Neoplasms. , 2021, , 409-459.		7
23	Survivorship Issues in Adult Patients With Histiocytic Neoplasms. <i>Journal of the National Comprehensive Cancer Network: JNCCN</i> , 2021, 19, 1312-1318.	4.9	1
24	Mosaicism for <i>KCNJ5</i> Causing Early-Onset Primary Aldosteronism due to Bilateral Adrenocortical Hyperplasia. <i>American Journal of Hypertension</i> , 2020, 33, 124-130.	2.0	20
25	An unexpected diagnosis of Graves' disease in an 81-year-old female with altered mental status. <i>Endocrine and Metabolic Science</i> , 2020, 1, 100055.	1.6	2
26	Insights into the Immunopathophysiology of Severe COVID-19 in Metabolic Disorders. <i>Annals of the National Academy of Medical Sciences (India)</i> , 2020, 56, 112-115.	0.3	1
27	Prevalence of Diabetes and Hypertension and Their Associated Risks for Poor Outcomes in Covid-19 Patients. <i>Journal of the Endocrine Society</i> , 2020, 4, bvaa102.	0.2	56
28	Prevalence of Hypothyroidism in Patients With Erdheim-Chester Disease. <i>JAMA Network Open</i> , 2020, 3, e2019169.	5.9	4
29	Hypertension Canada's 2020 Comprehensive Guidelines for the Prevention, Diagnosis, Risk Assessment, and Treatment of Hypertension in Adults and Children. <i>Canadian Journal of Cardiology</i> , 2020, 36, 596-624.	1.7	324
30	Hookah smoking and COVID-19: call for action. <i>Cmaj</i> , 2020, 192, E462-E462.	2.0	25
31	Association of BRAF V600E with Hypothalamic-Pituitary-Adrenal Axis Involvement in Erdheim-Chester Disease. <i>Endocrine and Metabolic Science</i> , 2020, 1, 100051.	1.6	2
32	Adrenocortical tumorigenesis: Lessons from genetics. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2020, 34, 101428.	4.7	36
33	Endocrine Conditions and COVID-19. <i>Hormone and Metabolic Research</i> , 2020, 52, 471-484.	1.5	34
34	A Gene-Based Classification of Primary Adrenocortical Hyperplasias. <i>Hormone and Metabolic Research</i> , 2020, 52, 133-141.	1.5	15
35	Assessment of Thyroid Function in Patients With Alkaptonuria. <i>JAMA Network Open</i> , 2020, 3, e201357.	5.9	12
36	Neurological manifestations of Erdheim-Chester Disease. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 497-506.	3.7	23

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37	Mass spectrometry-based steroid profiling in primary bilateral macronodular adrenocortical hyperplasia. <i>Endocrine-Related Cancer</i> , 2020, 27, 403-413.	3.1	13
38	ARMC5 variants in PRKAR1A-mutated patients modify cortisol levels and Cushing's syndrome. <i>Endocrine-Related Cancer</i> , 2020, 27, 509-517.	3.1	7
39	SAT-543 Human Hair Aldosterone Measurements for Evaluation of Primary Aldosteronism. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
40	SUN-713 Prevalence of Renal Cysts in Patients with Carney Complex. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
41	Targeting vulnerable atherosclerotic plaque via PET tracers aiming at cell surface overexpression of somatostatin receptors. <i>Biomedical Reports</i> , 2020, 13, 9.	2.0	2
42	Curative resection of an aldosteronoma causing primary aldosteronism in the second trimester of pregnancy. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2020, 2020, .	0.5	2
43	Commentary on A Rare Cause of Virilization, Short Stature, and Hypertension. <i>Clinical Chemistry</i> , 2020, 66, 1493-1494.	3.2	0
44	Inhibin A as a tumor marker for primary bilateral macronodular adrenal hyperplasia. <i>Endocrinology, Diabetes and Metabolism Case Reports</i> , 2020, 2020, .	0.5	0
45	MON-917 Carney Complex Due to a Contiguous Gene Deletion Syndrome (17q24.2-17q24.3). <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
46	A mitochondrial DNA D loop insertion detected almost exclusively in non-replicating tissues with maternal inheritance across three generations. <i>Mitochondrion</i> , 2019, 46, 298-301.	3.4	1
47	Comprehensive characterization of a Canadian cohort of von Hippel-Lindau disease patients. <i>Clinical Genetics</i> , 2019, 96, 461-467.	2.0	16
48	ARMC5 Variants and Risk of Hypertension in Blacks: MGRID Study. <i>Journal of the American Heart Association</i> , 2019, 8, e012508.	3.7	8
49	Expanding the Clinical Spectrum of LONP1-Related Mitochondrial Cytopathy. <i>Frontiers in Neurology</i> , 2019, 10, 981.	2.4	11
50	Resistant Hypertension. <i>Endocrinology and Metabolism Clinics of North America</i> , 2019, 48, 811-828.	3.2	5
51	Subspecialty training in adult inherited metabolic diseases: a call to action for unmet needs. <i>Lancet Diabetes and Endocrinology</i> , 2019, 7, 82-84.	11.4	7
52	Adrenal morphology and associated comorbidities in congenital adrenal hyperplasia. <i>Clinical Endocrinology</i> , 2019, 91, 247-255.	2.4	13
53	High expression of adrenal P450 aromatase (CYP19A1) in association with ARMC5-primary bilateral macronodular adrenocortical hyperplasia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 191, 105316.	2.5	13
54	Letter to the Editor: A Unique Case of Metastatic, Functional, Hereditary Paraganglioma Associated With an SDHC Germline Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 1158-1159.	3.6	0

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55	Genetics of Hypertension in African Americans and Others of African Descent. International Journal of Molecular Sciences, 2019, 20, 1081.	4.1	43
56	Artificial Intelligence and Machine Learning in Endocrinology and Metabolism: The Dawn of a New Era. Frontiers in Endocrinology, 2019, 10, 185.	3.5	35
57	Three cases of multi-generational Gaucher disease and colon cancer from an Ashkenazi Jewish family: A lesson for cascade screening. Molecular Genetics and Metabolism Reports, 2019, 18, 19-21.	1.1	2
58	Cover Image, Volume 91, Issue 2. Clinical Endocrinology, 2019, 91, i.	2.4	0
59	Whole-exome sequencing identifies a homozygous pathogenic variant in TAT in a girl with palmoplantar keratoderma. Molecular Genetics and Metabolism Reports, 2019, 21, 100534.	1.1	1
60	Hypophysitis: An update on the novel forms, diagnosis and management of disorders of pituitary inflammation. Best Practice and Research in Clinical Endocrinology and Metabolism, 2019, 33, 101371.	4.7	63
61	Severe cystic degeneration and intractable seizures in a newborn with molybdenum cofactor deficiency type B. Molecular Genetics and Metabolism Reports, 2019, 18, 11-13.	1.1	7
62	Growth hormone excess in neurofibromatosis 1. Genetics in Medicine, 2019, 21, 1254-1255.	2.4	13
63	MON-605 Hypothyroidism in Erdheim-Chester Disease: Experience from a Tertiary Care National Referral Center. Journal of the Endocrine Society, 2019, 3, .	0.2	1
64	Adrenocortical carcinoma and pulmonary embolism from tumoral extension. Endocrinology, Diabetes and Metabolism Case Reports, 2019, 2019, .	0.5	2
65	SAT-350 Comparative Proteomic Analysis of Various Forms of Bilateral Adrenocortical Hyperplasia. Journal of the Endocrine Society, 2019, 3, .	0.2	0
66	OR19-4 High Prevalence of Primary Hypothyroidism in Patients with Alkaptonuria Eighteen Years of Experience. Journal of the Endocrine Society, 2019, 3, .	0.2	0
67	OR02-6 Mass Spectrometry-Based Steroid Profiling Inprimary Bilateral Macronodular Adrenocortical Hyperplasia. Journal of the Endocrine Society, 2019, 3, .	0.2	0
68	Transcobalamin receptor defect: Identification of two new cases through positive newborn screening for propionic/methylmalonic aciduria and long-term outcome. American Journal of Medical Genetics, Part A, 2018, 176, 1411-1415.	1.2	13
69	Transplantation as disease modifying therapy in adults with inherited metabolic disorders. Journal of Inherited Metabolic Disease, 2018, 41, 885-896.	3.6	7
70	Coronary calcification in adults with Turner syndrome. Genetics in Medicine, 2018, 20, 664-668.	2.4	17
71	An update on the genetics of benign pituitary adenomas in children and adolescents. Current Opinion in Endocrine and Metabolic Research, 2018, 1, 19-24.	1.4	14
72	Successful Treatment of Estrogen Excess in Primary Bilateral Macronodular Adrenocortical Hyperplasia with Leuprolide Acetate. Hormone and Metabolic Research, 2018, 50, 124-132.	1.5	7

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73	Primary hypophysitis and other autoimmune disorders of the sellar and suprasellar regions. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 335-347.	5.7	34
74	Not quite type 1 or type 2, what now? Review of monogenic, mitochondrial, and syndromic diabetes. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 35-52.	5.7	25
75	An overview of inborn errors of metabolism manifesting with primary adrenal insufficiency. Reviews in Endocrine and Metabolic Disorders, 2018, 19, 53-67.	5.7	13
76	Adrenal Cortex; Physiology. , 2018, , 1-7.		1
77	65 YEARS OF THE DOUBLE HELIX: Endocrine tumour syndromes in children and adolescents. Endocrine-Related Cancer, 2018, 25, T221-T244.	3.1	10
78	Genetic Disorders of Adrenocortical Function. Endocrinology, 2018, , 727-763.	0.1	0
79	Application of Whole Exome Sequencing in the Clinical Diagnosis and Management of Inherited Cardiovascular Diseases in Adults. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	55
80	Genetics of Congenital Adrenal Hyperplasia. Endocrinology and Metabolism Clinics of North America, 2017, 46, 435-458.	3.2	56
81	Obesity and the diagnostic accuracy for primary aldosteronism. Journal of Clinical Hypertension, 2017, 19, 790-797.	2.0	10
82	Revisiting the prevalence of nonclassic congenital adrenal hyperplasia in US Ashkenazi Jews and Caucasians. Genetics in Medicine, 2017, 19, 1276-1279.	2.4	90
83	Thyroid Hormone Therapy for Older Adults with Subclinical Hypothyroidism. New England Journal of Medicine, 2017, 377, e20.	27.0	39
84	MEN4 and CDKN1B mutations: the latest of the MEN syndromes. Endocrine-Related Cancer, 2017, 24, T195-T208.	3.1	136
85	Metanephrines for Evaluating Palpitations and Flushing. JAMA - Journal of the American Medical Association, 2017, 318, 385.	7.4	11
86	Management of primary aldosteronism in patients with adrenal hemorrhage following adrenal vein sampling: A brief review with illustrative cases. Journal of Clinical Hypertension, 2017, 19, 1372-1376.	2.0	3
87	Obesity and the Diagnostic Accuracy for Primary Aldosteronism. Canadian Journal of Diabetes, 2017, 41, S29.	0.8	0
88	Coronary Atherosclerosis in Females with Turner Syndrome. Canadian Journal of Diabetes, 2017, 41, S30.	0.8	0
89	Pain in the groin: mycotic aneurysm after transcatheter aortic valve replacement. BMJ Case Reports, 2017, 2017, bcr-2017-220626.	0.5	0
90	Large pituitary gland with an expanding lesion in the context of neurofibromatosis 1. BMJ Case Reports, 2017, 2017, bcr-2017-222411.	0.5	5

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91	Enlarging hypermetabolic nodule: benign non-functional adrenocortical adenoma. BMJ Case Reports, 2017, 2017, bcr-2017-220820.	0.5	2
92	Genetics of Benign Adrenocortical Tumors. , 2017, , 31-53.		0
93	Genetic Disorders of Adrenocortical Function. Endocrinology, 2017, , 1-37.	0.1	0
94	Alterations of Phosphodiesterases in Adrenocortical Tumors. Frontiers in Endocrinology, 2016, 7, 111.	3.5	21
95	Cortisol in the Evaluation of Adrenal Insufficiency. JAMA - Journal of the American Medical Association, 2016, 316, 535.	7.4	5
96	Genetics of gigantism and acromegaly. Growth Hormone and IGF Research, 2016, 30-31, 37-41.	1.1	40
97	Flushing in (neuro)endocrinology. Reviews in Endocrine and Metabolic Disorders, 2016, 17, 373-380.	5.7	52
98	Rheumatologic and Imaging Manifestations of Thyroid Acropachy. Arthritis and Rheumatology, 2016, 68, 1636-1636.	5.6	2
99	Genetic Disorders of Adrenocortical Function. Endocrinology, 2016, , 1-37.	0.1	0
100	Across the globe in 4 months. Journal of Hypertension, 2015, 33, 891-893.	0.5	1
101	The Genetic Challenges and Opportunities in Advanced Heart Failure. Canadian Journal of Cardiology, 2015, 31, 1338-1350.	1.7	7
102	Teaching Video Neuro <i>Images</i> : Spasmodic dysphonia preceding idiopathic parkinsonism. Neurology, 2014, 82, e55.	1.1	0
103	Acylcarnitine profile in thyroid disease. Clinical Biochemistry, 2013, 46, 180-183.	1.9	10
104	Safety of statin therapy in patients with mitochondrial diseases. Journal of Clinical Lipidology, 2013, 7, 182.	1.5	3
105	Increased Prevalence of Hypertension in Young Adults with High Heteroplasmy Levels of the MELAS m.3243A>G Mutation. JIMD Reports, 2013, 12, 17-23.	1.5	13
106	Recurrent exercise-induced rhabdomyolysis. Cmaj, 2012, 184, 426-430.	2.0	16
107	ARMCS variants and risk of hypertension in African Americans: Minority Health-GRID study. Endocrine Abstracts, 0, , .	0.0	1
108	Comparative proteomic analysis of different bilateral adrenocortical hyperplasia. Endocrine Abstracts, 0, , .	0.0	0