

Nilufer Ozdemir Kutbay

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

24 papers	144 citations	7 h-index	11 g-index
39 ext. papers	207 ext. citations	2.3 avg, IF	2.15 L-index

#	Paper	IF	Citations
24	Clinical presentations, metabolic abnormalities and end-organ complications in patients with familial partial lipodystrophy. <i>Metabolism: Clinical and Experimental</i> , 2017 , 72, 109-119	12.7	36
23	Dopamine Agonist-Induced Impulse Control Disorders in Patients With Prolactinoma: A Cross-Sectional Multicenter Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 2527-2534	5.6	24
22	Prevalence and predictors of gestational diabetes mellitus: a nationwide multicentre prospective study. <i>Diabetic Medicine</i> , 2019 , 36, 221-227	3.5	15
21	Determining residual adipose tissue characteristics with MRI in patients with various subtypes of lipodystrophy. <i>Diagnostic and Interventional Radiology</i> , 2017 , 23, 428-434	3.2	14
20	Renal complications of lipodystrophy: A closer look at the natural history of kidney disease. <i>Clinical Endocrinology</i> , 2018 , 89, 65-75	3.4	12
19	Effects of metformin and pioglitazone combination on apoptosis and AMPK/mTOR signaling pathway in human anaplastic thyroid cancer cells. <i>Journal of Biochemical and Molecular Toxicology</i> , 2020 , 34, e22547	3.4	10
18	Characteristics and Treatment Results of 5 Patients with Fibrous Dysplasia and Review of the Literature. <i>Case Reports in Endocrinology</i> , 2015 , 2015, 670809	1.2	10
17	Acromegaly is associated with high fibroblast growth factor-21 levels. <i>Journal of Endocrinological Investigation</i> , 2019 , 42, 53-60	5.2	5
16	A case of familial partial lipodystrophy caused by a novel lamin A/C (LMNA) mutation in exon 1 (D47N). <i>European Journal of Internal Medicine</i> , 2016 , 29, 37-9	3.9	4
15	The low levels of bone morphogenic protein-4 and its antagonist noggin in type 2 diabetes. <i>Hormones</i> , 2018 , 17, 247-253	3.1	4
14	A novel CYP11B1 mutation in a Turkish patient with 11 β hydroxylase deficiency: An association with the severe hypokalemia leading to rhabdomyolysis. <i>Hormones</i> , 2016 , 15, 300-302	3.1	3
13	Cardiac phenotype in familial partial lipodystrophy. <i>Clinical Endocrinology</i> , 2021 , 94, 1043-1053	3.4	3
12	A new type of familial partial lipodystrophy: distinctive fat distribution and proteinuria. <i>Endocrine Research</i> , 2018 , 43, 258-263	1.9	1
11	A case of dyskeratosis congenita associated with hypothyroidism and hypogonadism. <i>Hormones</i> , 2016 , 15, 297-299	3.1	1
10	Identifying Clinical Characteristics of Hypoparathyroidism in Turkey: HIOPARATURK-NET Study. <i>Calcified Tissue International</i> , 2021 , 1	3.9	1
9	A case of idiopathic granulomatous hypophysitis. <i>Hormones</i> , 2017 , 16, 331-332	3.1	0
8	AN UNUSUAL CASE OF ACQUIRED PARTIAL LIPODYSTROPHY PRESENTING WITH. <i>Acta Endocrinologica</i> , 2019 , -5, 129-130	0.9	0

7	Biochemical characteristics and calcium and PTH levels of patients with high normal and elevated serum 25(OH)D levels in Turkey: DeVIT-TOX survey. <i>Archives of Osteoporosis</i> , 2021 , 16, 138	2.9	0
6	Adult Nesidioblastosis With Hypoglycemia Mimicking an Insulinoma: A Challenging Case. <i>International Surgery</i> , 2017 , 102, 324-327	0.1	
5	Nevus-like lesions on the lip and the foot. <i>European Journal of Internal Medicine</i> , 2018 , 53, e6-e7	3.9	
4	Magnetic resonance spectroscopy to assess hepatic steatosis in patients with lipodystrophy. <i>Turkish Journal of Gastroenterology</i> , 2020 , 31, 588-595	1	
3	Magnetic resonance spectroscopy to assess hepatic steatosis in patients with lipodystrophy. <i>Turkish Journal of Gastroenterology</i> , 2020 , 31, 588-595	1	
2	Approach to transgender individuals. <i>The Egyptian Journal of Internal Medicine</i> , 2017 , 29, 1-4	0.5	
1	Two challenging cases of pituitaryoma. <i>Hormones</i> , 2021 , 20, 813-818	3.1	