

# Nivedita Patni

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5232466/publications.pdf>

Version: 2024-02-01

26  
papers

1,129  
citations

759233

12  
h-index

642732

23  
g-index

26  
all docs

26  
docs citations

26  
times ranked

1430  
citing authors

#	ARTICLE	IF	CITATIONS
1	The Diagnosis and Management of Lipodystrophy Syndromes: A Multi-Society Practice Guideline. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 4500-4511.	3.6	323
2	YouTube As a Source of Information on the H1N1 Influenza Pandemic. <i>American Journal of Preventive Medicine</i> , 2010, 38, e1-e3.	3.0	269
3	Congenital generalized lipodystrophies—new insights into metabolic dysfunction. <i>Nature Reviews Endocrinology</i> , 2015, 11, 522-534.	9.6	195
4	Lipodystrophies, dyslipidaemias and atherosclerotic cardiovascular disease. <i>Pathology</i> , 2019, 51, 202-212.	0.6	67
5	A Novel Generalized Lipodystrophy-Associated Progeroid Syndrome Due to Recurrent Heterozygous LMNA p.T10I Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 1005-1014.	3.6	47
6	Bi-allelic POLR3A Loss-of-Function Variants Cause Autosomal-Recessive Wiedemann-Rautenstrauch Syndrome. <i>American Journal of Human Genetics</i> , 2018, 103, 968-975.	6.2	43
7	Regional Body Fat Changes and Metabolic Complications in Children With Dunnigan Lipodystrophy-Causing LMNA Variants. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 1099-1108.	3.6	27
8	A Novel Syndrome of Generalized Lipodystrophy Associated With Pilocytic Astrocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 3603-3606.	3.6	21
9	The prevalence and etiology of extreme hypertriglyceridemia in children: Data from a tertiary children's hospital. <i>Journal of Clinical Lipidology</i> , 2018, 12, 305-310.	1.5	20
10	Juvenile-onset generalized lipodystrophy due to a novel heterozygous missense LMNA mutation affecting lamin C. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 2517-2521.	1.2	17
11	Type 1 hyperlipoproteinemia in a child with large homozygous deletion encompassing GPIHBP1. <i>Journal of Clinical Lipidology</i> , 2016, 10, 1035-1039.e2.	1.5	16
12	Assessment of risk and prophylaxis for deep vein thrombosis and pulmonary embolism in medically ill patients during their early days of hospital stay at a tertiary care center in a developing country. <i>Vascular Health and Risk Management</i> , 2009, 5, 643.	2.3	15
13	Orlistat Therapy for Children With Type 1 Hyperlipoproteinemia: A Randomized Clinical Trial. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 2403-2407.	3.6	13
14	Postmortem Findings in a Young Man With Congenital Generalized Lipodystrophy, Type 4 Due to CAVIN1 Mutations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 957-960.	3.6	12
15	Lipodystrophy for the Diabetologist—What to Look For. <i>Current Diabetes Reports</i> , 2022, 22, 461-470.	4.2	12
16	Elevated Î±-fetoprotein levels in Van Wyk-Grumbach syndrome: a case report and review of literature. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2012, 25, 761-7.	0.9	8
17	An additional case of NÃ©stor-Guillermo progeria syndrome diagnosed in early childhood. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 2399-2402.	1.2	8
18	A novel autosomal recessive lipodystrophy syndrome due to homozygous LMNA variant. <i>Journal of Medical Genetics</i> , 2020, 57, 422-426.	3.2	4

#	ARTICLE	IF	CITATIONS
19	Case Studies in Pediatric Lipid Disorders and Their Management. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 3605-3620.	3.6	4
20	Diazoxide for the Treatment of Hypoglycemia Resulting From Dumping Syndrome in a Child. Journal of the Endocrine Society, 2019, 3, 1357-1360.	0.2	3
21	Worsening hypertriglyceridemia with oral contraceptive pills in an adolescent with HIV-associated lipodystrophy: a case report and review of the literature. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1247-51.	0.9	2
22	Effect of recombinant human insulin-like growth factor 1 therapy in a child with 3-M syndrome-1 with CUL7 gene mutation. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 1609-1612.	0.9	2
23	Should children with chronic diarrhea be referred to a lipid clinic?. Journal of Clinical Lipidology, 2018, 12, 1099-1101.	1.5	1
24	Case 3: Hypoglycemia in an Infant with Cholestasis. Pediatrics in Review, 2019, 40, 488-490.	0.4	0
25	Diet-Responsive Hypercholesterolemia With Cardiofaciocutaneous Syndrome Type 3. Journal of the Endocrine Society, 2021, 5, A308-A308.	0.2	0
26	A 16-Year-Old Girl with Polyuria, Polydipsia, and a New-Onset Rash Around Her Elbows and Knees. Pediatric Annals, 2016, 45, e317-8.	0.8	0