

# Avishay Lahad

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4617928/publications.pdf>

Version: 2024-02-01

14  
papers

344  
citations

1039880

9  
h-index

1058333

14  
g-index

14  
all docs

14  
docs citations

14  
times ranked

836  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in <i>STN1</i> cause Coats plus syndrome and are associated with genomic and telomere defects. <i>Journal of Experimental Medicine</i> , 2016, 213, 1429-1440.	4.2	100
2	Endocrine abnormalities in ataxia telangiectasia: findings from a national cohort. <i>Pediatric Research</i> , 2016, 79, 889-894.	1.1	63
3	Individualized Dynamics in the Gut Microbiota Precede Crohn's Disease Flares. <i>American Journal of Gastroenterology</i> , 2019, 114, 1142-1151.	0.2	50
4	Congenital protein losing enteropathy: an inborn error of lipid metabolism due to DGAT1 mutations. <i>European Journal of Human Genetics</i> , 2016, 24, 1268-1273.	1.4	37
5	Current therapy of pediatric Crohn's disease. <i>World Journal of Gastrointestinal Pathophysiology</i> , 2015, 6, 33.	0.5	24
6	Elevated IgM levels as a marker for a unique phenotype in patients with Ataxia telangiectasia. <i>BMC Pediatrics</i> , 2018, 18, 185.	0.7	15
7	Impaired IL-10 Receptor-mediated Suppression in Monocyte From Patients With Crohn Disease. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018, 66, 779-784.	0.9	12
8	Genetic and Structural Analysis of a SKIV2L Mutation Causing Tricho-hepato-enteric Syndrome. <i>Digestive Diseases and Sciences</i> , 2018, 63, 1192-1199.	1.1	11
9	Long-term nutritional and gastrointestinal aspects in patients with ataxia telangiectasia. <i>Nutrition</i> , 2018, 46, 48-52.	1.1	10
10	Cerebral and portal vein thrombosis, macrocephaly and atypical absence seizures in Glycosylphosphatidyl inositol deficiency due to a PIGM promoter mutation. <i>Molecular Genetics and Metabolism</i> , 2019, 128, 151-161.	0.5	9
11	Longitudinal changes in bone mineral density in children with inflammatory bowel diseases. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2020, 109, 1026-1032.	0.7	7
12	Relationships between Clinical Presentation, Serology, Histology, and Duodenal Deposits of Tissue Transglutaminase Antibodies in Pediatric Celiac Disease. <i>Digestive Diseases</i> , 2018, 36, 369-376.	0.8	3
13	Resting Energy Expenditure in Patients With Familial Dysautonomia. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2019, 68, 422-427.	0.9	2
14	Glycogen Storage Disease type IA refractory to cornstarch: Can next generation sequencing offer a solution?. <i>European Journal of Medical Genetics</i> , 2022, , 104518.	0.7	1