Avishay Lahad

List of Publications by Year
in descending order

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