

Abdul Elkadri

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

1,753
citations

623734

14
h-index

839539

18
g-index

19
all docs

19
docs citations

19
times ranked

3518
citing authors

#	ARTICLE	IF	CITATIONS
1	Bone Marrow Suppression Associated With Celiac Disease in a 4-Year-Old Boy. <i>ACG Case Reports Journal</i> , 2021, 8, e00538.	0.4	1
2	Variants in <i>STXBP3</i> are Associated with Very Early Onset Inflammatory Bowel Disease, Bilateral Sensorineural Hearing Loss and Immune Dysregulation. <i>Journal of Crohn's and Colitis</i> , 2021, 15, 1908-1919.	1.3	7
3	North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition Position Paper on the Evaluation and Management for Patients With Very Early-Onset Inflammatory Bowel Disease. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2020, 70, 389-403.	1.8	79
4	Somatic mosaicism and common genetic variation contribute to the risk of very-early-onset inflammatory bowel disease. <i>Nature Communications</i> , 2020, 11, 995.	12.8	37
5	Prevalence and Clinical Features of Inflammatory Bowel Diseases Associated With Monogenic Variants, Identified by Whole-Exome Sequencing in 1000 Children at a Single Center. <i>Gastroenterology</i> , 2020, 158, 2208-2220.	1.3	81
6	Clinical Guideline Highlights for the Hospitalist: Clostridium difficile Infections in Children. <i>Journal of Hospital Medicine</i> , 2020, 15, 98-100.	1.4	0
7	A Systematic Review of Micronutrient Deficiencies in Pediatric Inflammatory Bowel Disease. <i>Inflammatory Bowel Diseases</i> , 2019, 25, 445-459.	1.9	22
8	Impaired antibacterial autophagy links granulomatous intestinal inflammation in Niemann-Pick disease type C1 and XIAP deficiency with NOD2 variants in Crohn's disease. <i>Gut</i> , 2017, 66, 1060-1073.	12.1	126
9	Loss of the Arp2/3 complex component ARPC1B causes platelet abnormalities and predisposes to inflammatory disease. <i>Nature Communications</i> , 2017, 8, 14816.	12.8	176
10	Ankyrin repeat and zinc-finger domain-containing 1 mutations are associated with infantile-onset inflammatory bowel disease. <i>Journal of Biological Chemistry</i> , 2017, 292, 7904-7920.	3.4	29
11	The operative management of children with complex perianal Crohn's disease. <i>Journal of Pediatric Surgery</i> , 2016, 51, 1993-1997.	1.6	11
12	Variants in TRIM22 That Affect NOD2 Signaling Are Associated With Very-Early-Onset Inflammatory Bowel Disease. <i>Gastroenterology</i> , 2016, 150, 1196-1207.	1.3	88
13	Defects in Nicotinamide-adenine Dinucleotide Phosphate Oxidase Genes NOX1 and DUOX2 in Very Early Onset Inflammatory Bowel Disease. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2015, 1, 489-502.	4.5	127
14	Mutations in Plasmalemma Vesicle Associated Protein Result in Sieving Protein-Losing Enteropathy Characterized by Hypoproteinemia, Hypoalbuminemia, and Hypertriglyceridemia. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2015, 1, 381-394.e7.	4.5	43
15	The role of surgery for children with perianal Crohn's disease. <i>Journal of Pediatric Surgery</i> , 2015, 50, 140-143.	1.6	15
16	Higher Activity of the Inducible Nitric Oxide Synthase Contributes to Very Early Onset Inflammatory Bowel Disease. <i>Clinical and Translational Gastroenterology</i> , 2014, 5, e46.	2.5	71
17	The Diagnostic Approach to Monogenic Very Early Onset Inflammatory Bowel Disease. <i>Gastroenterology</i> , 2014, 147, 990-1007.e3.	1.3	559
18	Mutations in Tetratricopeptide Repeat Domain 7A Result in a Severe Form of Very Early Onset Inflammatory Bowel Disease. <i>Gastroenterology</i> , 2014, 146, 1028-1039.	1.3	175

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19	Variants in Nicotinamide Adenine Dinucleotide Phosphate Oxidase Complex Components Determine Susceptibility to Very Early Onset Inflammatory Bowel Disease. <i>Gastroenterology</i> , 2014, 147, 680-689.e2.	1.3	106