Aleixo M Muise

List of Publications by Year in descending order

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46918 54797 7,643 117 47 84 citations h-index g-index papers 120 120 120 10468 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	The Diagnostic Approach to Monogenic Very Early Onset Inflammatory Bowel Disease. Gastroenterology, 2014, 147, 990-1007.e3.	0.6	559
2	Common variants at five new loci associated with early-onset inflammatory bowel disease. Nature Genetics, 2009, 41, 1335-1340.	9.4	459
3	Interleukin-10 Receptor Signaling in Innate Immune Cells Regulates Mucosal Immune Tolerance and Anti-Inflammatory Macrophage Function. Immunity, 2014, 40, 706-719.	6.6	455
4	Interleukin 10 Receptor Signaling. Advances in Immunology, 2014, 122, 177-210.	1.1	239
5	Clinical outcome in IL-10– and IL-10 receptor–deficient patients with or without hematopoietic stem cell transplantation. Journal of Allergy and Clinical Immunology, 2013, 131, 825-830.e9.	1.5	236
6	Incidence, Outcomes, and Health Services Burden of Very Early Onset Inflammatory Bowel Disease. Gastroenterology, 2014, 147, 803-813.e7.	0.6	222
7	IL-10R Polymorphisms Are Associated with Very-early-onset Ulcerative Colitis. Inflammatory Bowel Diseases, 2013, 19, 115-123.	0.9	212
8	Loss of the Arp2/3 complex component ARPC1B causes platelet abnormalities and predisposes to inflammatory disease. Nature Communications, 2017, 8, 14816.	5.8	176
9	Mutations in Tetratricopeptide Repeat Domain 7A Result in a Severe Form of Very Early Onset Inflammatory Bowel Disease. Gastroenterology, 2014, 146, 1028-1039.	0.6	175
10	A eukaryotic transcriptional represser with carboxypeptidase activity. Nature, 1995, 378, 92-96.	13.7	161
11	NADPH oxidase complex and IBD candidate gene studies: identification of a rare variant in <i>NCF2</i> that results in reduced binding to RAC2. Gut, 2012, 61, 1028-1035.	6.1	158
12	Interleukin $1\hat{l}^2$ Mediates Intestinal Inflammation in Mice and Patients With Interleukin 10 Receptor Deficiency. Gastroenterology, 2016, 151, 1100-1104.	0.6	156
13	Eating disorders in adolescent boys: a review of the adolescent and young adult literature. Journal of Adolescent Health, 2003, 33, 427-435.	1.2	148
14	Human RIPK1 deficiency causes combined immunodeficiency and inflammatory bowel diseases. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 970-975.	3.3	130
15	Advances in Evaluation of Chronic Diarrhea in Infants. Gastroenterology, 2018, 154, 2045-2059.e6.	0.6	129
16	Defects in Nicotinamide-adenine Dinucleotide Phosphate Oxidase Genes NOX1 and DUOX2 in Very Early Onset Inflammatory Bowel Disease. Cellular and Molecular Gastroenterology and Hepatology, 2015, 1, 489-502.	2.3	127
17	Clinical Genomics in Inflammatory Bowel Disease. Trends in Genetics, 2017, 33, 629-641.	2.9	123
18	Listeria monocytogenes exploits efferocytosis to promote cell-to-cell spread. Nature, 2014, 509, 230-234.	13.7	118

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19	Reduced sodium/proton exchanger NHE3 activity causes congenital sodium diarrhea. Human Molecular Genetics, 2015, 24, 6614-6623.	1.4	111
20	Palmitoylation of NOD1 and NOD2 is required for bacterial sensing. Science, 2019, 366, 460-467.	6.0	109
21	Host and bacterial factors that regulate LC3 recruitment to <i><i>Listeria monocytogenes</i><ii>during the early stages of macrophage infection. Autophagy, 2013, 9, 985-995.</ii></i>	4.3	108
22	Variants in Nicotinamide Adenine Dinucleotide Phosphate Oxidase Complex Components Determine Susceptibility to Very Early Onset Inflammatory Bowel Disease. Gastroenterology, 2014, 147, 680-689.e2.	0.6	106
23	Regulation of Adipogenesis by a Transcriptional Repressor That Modulates MAPK Activation. Journal of Biological Chemistry, 2001, 276, 10199-10206.	1.6	105
24	Very Early Onset Inflammatory Bowel Disease: A Clinical Approach With a Focus on the Role of Genetics and Underlying Immune Deficiencies. Inflammatory Bowel Diseases, 2020, 26, 820-842.	0.9	100
25	Inherited p40phox deficiency differs from classic chronic granulomatous disease. Journal of Clinical Investigation, 2018, 128, 3957-3975.	3.9	99
26	Human TGF- \hat{l}^21 deficiency causes severe inflammatory bowel disease and encephalopathy. Nature Genetics, 2018, 50, 344-348.	9.4	95
27	Intestinal Inflammation and Dysregulated Immunity in Patients With Inherited Caspase-8 Deficiency. Gastroenterology, 2019, 156, 275-278.	0.6	92
28	Variants in TRIM22 That Affect NOD2 Signaling Are Associated With Very-Early-Onset Inflammatory Bowel Disease. Gastroenterology, 2016, 150, 1196-1207.	0.6	88
29	The Age of Gene Discovery in Very Early Onset Inflammatory Bowel Disease. Gastroenterology, 2012, 143, 285-288.	0.6	85
30	The <i> NOD2 </i> insC polymorphism is associated with worse outcome following ileal pouch-anal anastomosis for ulcerative colitis. Gut, 2013, 62, 1433-1439.	6.1	85
31	Very Early-onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2015, 21, 1166-1175.	0.9	82
32	Prevalence and Clinical Features of Inflammatory Bowel Diseases Associated With Monogenic Variants, Identified by Whole-Exome Sequencing in 1000 Children at a Single Center. Gastroenterology, 2020, 158, 2208-2220.	0.6	81
33	Clinical Genomics for the Diagnosis of Monogenic Forms of Inflammatory Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2021, 72, 456-473.	0.9	79
34	Infliximab Maintains Durable Response and Facilitates Catch-up Growth in Luminal Pediatric Crohn's Disease. Inflammatory Bowel Diseases, 2014, 20, 1177-1186.	0.9	78
35	Listeriolysin O Suppresses Phospholipase C-Mediated Activation of the Microbicidal NADPH Oxidase to Promote Listeria monocytogenes Infection. Cell Host and Microbe, 2011, 10, 627-634.	5.1	72
36	Higher Activity of the Inducible Nitric Oxide Synthase Contributes to Very Early Onset Inflammatory Bowel Disease. Clinical and Translational Gastroenterology, 2014, 5, e46.	1.3	71

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37	Single Nucleotide Polymorphisms That Increase Expression of the Guanosine Triphosphatase RAC1 Are Associated With Ulcerative Colitis. Gastroenterology, 2011, 141, 633-641.	0.6	67
38	Mucus sialylation determines intestinal host-commensal homeostasis. Cell, 2022, 185, 1172-1188.e28.	13.5	66
39	C-reactive protein (CRP), erythrocyte sedimentation rate (ESR) or both? A systematic evaluation in pediatric ulcerative colitis. Journal of Crohn's and Colitis, 2011, 5, 423-429.	0.6	63
40	Very Early Onset Inflammatory Bowel Disease Associated with Aberrant Trafficking of IL-10R1 and Cure by T Cell Replete Haploidentical Bone Marrow Transplantation. Journal of Clinical Immunology, 2014, 34, 331-339.	2.0	62
41	Infliximabâ€Induced Psoriasis and Psoriasiform Skin Lesions in Pediatric Crohn Disease and a Potential Association With ILâ€23 Receptor Polymorphisms. Journal of Pediatric Gastroenterology and Nutrition, 2013, 56, 512-518.	0.9	61
42	Transcriptional regulation by the \hat{l}^35 subunit of a heterotrimeric G protein during adipogenesis. EMBO Journal, 1999, 18, 4004-4012.	3.5	57
43	An ATG16L1-dependent pathway promotes plasma membrane repair and limits Listeria monocytogenes cell-to-cell spread. Nature Microbiology, 2018, 3, 1472-1485.	5.9	57
44	A Systematic Review of Monogenic Inflammatory Bowel Disease. Clinical Gastroenterology and Hepatology, 2022, 20, e653-e663.	2.4	57
45	Microvillus Inclusion Disease: Loss of Myosin Vb Disrupts Intracellular Traffic and Cell Polarity. Traffic, 2014, 15, 22-42.	1.3	56
46	Gain-of-function variants in SYK cause immune dysregulation and systemic inflammation in humans and mice. Nature Genetics, 2021, 53, 500-510.	9.4	56
47	Protein-Tyrosine Phosphatase Sigma Is Associated with Ulcerative Colitis. Current Biology, 2007, 17, 1212-1218.	1.8	53
48	Protein tyrosine phosphatase $largets$ apical junction complex proteins in the intestine and regulates epithelial permeability. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 693-698.	3.3	53
49	Are Children With Kawasaki Disease and Prolonged Fever at Risk for Macrophage Activation Syndrome?. Pediatrics, 2003, 112, e495-e497.	1.0	52
50	A Frameshift in CSF2RB Predominant Among Ashkenazi Jews Increases Risk for Crohn's Disease and Reduces Monocyte Signaling via GM-CSF. Gastroenterology, 2016, 151, 710-723.e2.	0.6	51
51	The Diaphanous-Related Formins Promote Protrusion Formation and Cell-to-Cell Spread of <i>Listeria monocytogenes </i> . Journal of Infectious Diseases, 2015, 211, 1185-1195.	1.9	49
52	TTC7A: Steward of Intestinal Health. Cellular and Molecular Gastroenterology and Hepatology, 2019, 7, 555-570.	2.3	48
53	Human <scp>ALPI</scp> deficiency causes inflammatory bowel disease and highlights a key mechanism of gut homeostasis. EMBO Molecular Medicine, 2018, 10, .	3.3	47
54	Diagnostic delay in Canadian children with inflammatory bowel disease is more common in Crohn's disease and associated with decreased height. Archives of Disease in Childhood, 2018, 103, 319-326.	1.0	45

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55	Interleukin-6 is associated with steroid resistance and reflects disease activity in severe pediatric ulcerative colitis. Journal of Crohn's and Colitis, 2013, 7, 916-922.	0.6	43
56	Mutations in Plasmalemma Vesicle Associated Protein Result in Sieving Protein-Losing Enteropathy Characterized by Hypoproteinemia, Hypoalbuminemia, and Hypertriglyceridemia. Cellular and Molecular Gastroenterology and Hepatology, 2015, 1, 381-394.e7.	2.3	43
57	Mucosa-Associated Ileal Microbiota in New-Onset Pediatric Crohn's Disease. Inflammatory Bowel Diseases, 2016, 22, 1533-1539.	0.9	43
58	Biliary Atresia With Choledochal Cyst: Implications for Classification. Clinical Gastroenterology and Hepatology, 2006, 4, 1411-1414.	2.4	39
59	Somatic mosaicism and common genetic variation contribute to the risk of very-early-onset inflammatory bowel disease. Nature Communications, 2020, 11, 995.	5.8	37
60	An Integrated Taxonomy for Monogenic Inflammatory Bowel Disease. Gastroenterology, 2022, 162, 859-876.	0.6	37
61	Drug Screen Identifies Leflunomide for Treatment of Inflammatory Bowel Disease Caused by TTC7A Deficiency. Gastroenterology, 2020, 158, 1000-1015.	0.6	36
62	Inflammatory Bowel Disease. Gastroenterology Clinics of North America, 2018, 47, 755-772.	1.0	34
63	Fatal autoimmunity in mice reconstituted with human hematopoietic stem cells encoding defective FOXP3. Blood, 2015, 125, 3886-3895.	0.6	33
64	Large Bâ€Cell Lymphoma in an Adolescent Patient With Interleukinâ€10 Receptor Deficiency and History of Infantile Inflammatory Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2016, 63, e15-7.	0.9	31
65	Diagnostic Delay Is Associated With Complicated Disease and Growth Impairment in Paediatric Crohn's Disease. Journal of Crohn's and Colitis, 2021, 15, 419-431.	0.6	30
66	Human autoinflammatory disease reveals ELF4 as a transcriptional regulator of inflammation. Nature Immunology, 2021, 22, 1118-1126.	7.0	30
67	Ankyrin repeat and zinc-finger domain-containing 1 mutations are associated with infantile-onset inflammatory bowel disease. Journal of Biological Chemistry, 2017, 292, 7904-7920.	1.6	29
68	Mutation spectrum of NOD2 reveals recessive inheritance as a main driver of Early Onset Crohn's Disease. Scientific Reports, 2021, 11, 5595.	1.6	29
69	Novel de novo mutations of the interleukin-10 receptor gene lead to infantile onset inflammatory bowel disease. Journal of Crohn's and Colitis, 2014, 8, 1551-1556.	0.6	28
70	Enhanced TH17 Responses in Patients with IL10 Receptor Deficiency and Infantile-onset IBD. Inflammatory Bowel Diseases, 2017, 23, 1950-1961.	0.9	28
71	Antiâ€₹NF, Infliximab, and Adalimumab Can Be Effective in Eosinophilic Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2013, 56, 492-497.	0.9	27
72	CARMIL2 Deficiency Presenting as Very Early Onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2019, 25, 1788-1795.	0.9	26

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73	Intensified Infliximab Induction is Associated with Improved Response and Decreased Colectomy in Steroid-Refractory Paediatric Ulcerative Colitis. Journal of Crohn's and Colitis, 2019, 13, 982-989.	0.6	26
74	<i>Natural History of </i> Very Early Onset Inflammatory Bowel Disease <i>in North America: A Retrospective Cohort Study</i> Inflammatory Bowel Diseases, 2021, 27, 295-302.	0.9	25
75	Enzymic characterization of a novel member of the regulatory B-like carboxypeptidase with transcriptional repression function: stimulation of enzymic activity by its target DNA. Biochemical Journal, 1999, 343, 341-345.	1.7	23
76	Low Levels of Procalcitonin During Episodes of Necrotizing Enterocolitis. Digestive Diseases and Sciences, 2007, 52, 2972-2976.	1.1	22
77	Rac2-Deficiency Leads to Exacerbated and Protracted Colitis in Response to Citrobacter rodentium Infection. PLoS ONE, 2013, 8, e61629.	1.1	22
78	Long-term outcomes for children with very early-onset colitis: Implications for surgical management. Journal of Pediatric Surgery, 2018, 53, 964-967.	0.8	22
79	Very early onset IBD: novel genetic aetiologies. Current Opinion in Allergy and Clinical Immunology, 2018, 18, 470-480.	1.1	19
80	Clinical Phenotypes and Outcomes in Monogenic Versus Non-monogenic Very Early Onset Inflammatory Bowel Disease. Journal of Crohn's and Colitis, 2022, 16, 1380-1396.	0.6	19
81	Replication of genetic variation in the MYO9B gene in Crohn's disease. Human Immunology, 2011, 72, 592-597.	1.2	17
82	A Novel Nonsense Mutation in the <i>EpCAM</i> Gene in a Patient With Congenital Tufting Enteropathy. Journal of Pediatric Gastroenterology and Nutrition, 2014, 58, 18-21.	0.9	17
83	Multilabel immunofluorescence and antigen reprobing on formalin-fixed paraffin-embedded sections: novel applications for precision pathology diagnosis. Modern Pathology, 2016, 29, 557-569.	2.9	17
84	Advanced Understanding of Monogenic Inflammatory Bowel Disease. Frontiers in Pediatrics, 2020, 8, 618918.	0.9	16
85	Monogenic Intestinal Epithelium Defects and the Development of Inflammatory Bowel Disease. Physiology, 2018, 33, 360-369.	1.6	15
86	Deficiency in X-linked inhibitor of apoptosis protein promotes susceptibility to microbial triggers of intestinal inflammation. Science Immunology, 2021, 6, eabf7473.	5.6	15
87	ARPC1B binds WASP to control actin polymerization and curtail tonic signaling in B cells. JCI Insight, 2021, 6, .	2.3	13
88	A CARD9 Polymorphism Is Associated with Decreased Likelihood of Persistent Conjugated Hyperbilirubinemia in Intestinal Failure. PLoS ONE, 2014, 9, e85915.	1.1	11
89	Novel CARMIL2 loss-of-function variants are associated with pediatric inflammatory bowel disease. Scientific Reports, 2021, 11, 5945.	1.6	11
90	Association between a Multi-Locus Genetic Risk Score and Inflammatory Bowel Disease. Bioinformatics and Biology Insights, 2013, 7, BBI.S11601.	1.0	9

#	Article	lF	Citations
91	Rac1 Polymorphisms and Thiopurine Efficacy in Children With Inflammatory Bowel Disease. Journal of Pediatric Gastroenterology and Nutrition, 2015, 61, 404-407.	0.9	9
92	NOX1 Regulates Collective and Planktonic Cell Migration: Insights From Patients With Pediatric-Onset IBD and NOX1 Deficiency. Inflammatory Bowel Diseases, 2020, 26, 1166-1176.	0.9	9
93	Unrelated donor hematopoietic stem cell transplantation for infantile enteropathy due to <scp> L< scp>å€10 <scp> L< scp> </scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp></scp>	0.5	8
94	A Chromosomal Duplication Encompassing Interleukin-33 Causes a Novel Hyper IgE Phenotype Characterized by Eosinophilic Esophagitis and Generalized Autoimmunity. Gastroenterology, 2022, 163, 510-513.e3.	0.6	8
95	Apical junction complex proteins and ulcerative colitis: a focus on the <i>PTPRS </i> gene. Expert Review of Molecular Diagnostics, 2008, 8, 465-477.	1.5	7
96	Predictive Prenatal Diagnosis for Infantileâ€onset Inflammatory Bowel Disease Because of Interleukinâ€10 Signalling Defects. Journal of Pediatric Gastroenterology and Nutrition, 2021, 72, 276-281.	0.9	7
97	Variants in <i>STXBP3</i> are Associated with Very Early Onset Inflammatory Bowel Disease, Bilateral Sensorineural Hearing Loss and Immune Dysregulation. Journal of Crohn's and Colitis, 2021, 15, 1908-1919.	0.6	7
98	Very Early Onset IBD: How Very Different â€~on Average'?. Journal of Crohn's and Colitis, 2017, 11, jjw217.	0.6	6
99	The risk of myocardial infarction in HIV-infected patients receiving HAART: a case report. International Journal of STD and AIDS, 2001, 12, 612-613.	0.5	5
100	Cutting Edge: NOX2 NADPH Oxidase Controls Infection by an Intracellular Bacterial Pathogen through Limiting the Type 1 IFN Response. Journal of Immunology, 2021, 206, 323-328.	0.4	5
101	Whipple disease mimicking inflammatory bowel disease. Intestinal Research, 2021, 19, 119-125.	1.0	5
102	The E3 ubiquitin ligase UBR5 interacts with TTC7A and may be associated with very early onset inflammatory bowel disease. Scientific Reports, 2020, 10, 18648.	1.6	4
103	Novel Exonic Deletions in TTC7A in a Newborn with Multiple Intestinal Atresia and Combined Immunodeficiency. Journal of Clinical Immunology, 2019, 39, 616-619.	2.0	3
104	Application of Whole Exome Sequencing in Congenital Secretory Diarrhea Diagnosis. Journal of Pediatric Gastroenterology and Nutrition, 2019, 68, e106-e108.	0.9	2
105	Utilization of Whole Exome Sequencing Data to Identify Clinically Relevant Pharmacogenomic Variants in Pediatric Inflammatory Bowel Disease. Clinical and Translational Gastroenterology, 2020, 11, e00263.	1.3	1
106	Multisystem Autoimmune Inflammatory Disease, Including Colitis, Due to Inborn Error of Immunity. Pediatrics, 2021, 148, e2021050614.	1.0	1
107	Platelet VPS16B is dependent on VPS33B expression, as determined in two siblings with arthrogryposis, renal dysfunction and cholestasis (ARC) syndrome. Journal of Thrombosis and Haemostasis, 2022, , .	1.9	1
108	NADPH oxidase complex and IBD Candidate Gene studies. Inflammatory Bowel Diseases, 2011, 17, S8.	0.9	0

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109	The Authors' reply: Figure 1. Gut, 2012, 61, 1097.2-1098.	6.1	O
110	Inflammatory Bowel Disease in Primary Immunodeficiencies. , 2018, , 167-181.		0
111	Very Early Onset Inflammatory Bowel Disease (VEOIBD)., 2019,, 383-404.		O
112	Pediatric Diarrheal Disorders. , 2020, , 143-157.		0
113	2019 Harry Shwachman Award. Journal of Pediatric Gastroenterology and Nutrition, 2020, 70, 405-405.	0.9	O
114	Sequencing and Mapping IBD Genes to Individual Causative Variants and Their Clinical Relevance. , 2019, , 117-139.		0
115	A Machine Learning Approach to Identifying Causal Monogenic Variants in Inflammatory Bowel Disease. , 2022, 1, 171-179.		0
116	Valosin-containing protein-regulated endoplasmic reticulum stress causes NOD2-dependent inflammatory responses. Scientific Reports, 2022, 12, 3906.	1.6	0
117	Histopathological Features of Monogenic Inflammatory Bowel Disease: Sub-Analysis of Systematic Review. , 2022, , .		O