Anna Monica Rosaria Bianco

List of Publications by Year
in descending order

40	833	567247 15	501174 28
papers	citations	h-index	g-index

41	41	41	1715
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Type I interferon-mediated autoinflammation due to DNase II deficiency. Nature Communications, 2017, 8, 2176.	12.8	164
2	Genetics of inflammatory bowel disease from multifactorial to monogenic forms. World Journal of Gastroenterology, 2015, 21, 12296.	3.3	113
3	Curcumin and Inflammatory Bowel Disease: Potential and Limits of Innovative Treatments. Molecules, 2014, 19, 21127-21153.	3.8	105
4	Spectrum ofFANCA mutations in Italian Fanconi anemia patients: Identification of six novel alleles and phenotypic characterization of the S858R variant. Human Mutation, 2003, 22, 338-339.	2.5	35
5	Lovastatinâ€induced apoptosis is modulated by geranylgeraniol in a neuroblastoma cell line. International Journal of Developmental Neuroscience, 2012, 30, 451-456.	1.6	33
6	Diagnostic Approach to Monogenic Inflammatory Bowel Disease in Clinical Practice: A Ten-Year Multicentric Experience. Inflammatory Bowel Diseases, 2020, 26, 720-727.	1.9	32
7	Mevalonate kinase deficiency and IBD: shared genetic background. Gut, 2014, 63, 1367-1368.	12.1	30
8	Genetic profile of patients with early onset inflammatory bowel disease. Gene, 2018, 645, 18-29.	2.2	29
9	The diagnostic challenge of very early-onset enterocolitis in an infant with XIAP deficiency. BMC Pediatrics, 2015, 15, 208.	1.7	25
10	Iron signature in asbestos-induced malignant pleural mesothelioma: A population-based autopsy study. Journal of Toxicology and Environmental Health - Part A: Current Issues, 2016, 79, 129-141.	2.3	24
11	Genetic profiling of autoinflammatory disorders in patients with periodic fever: a prospective study. Pediatric Rheumatology, 2015, 13, 11.	2.1	22
12	Expression and association data strongly support JARID2 involvement in nonsyndromic cleft lip with or without cleft palate. Human Mutation, 2010, 31, 794-800.	2.5	19
13	Database tools in genetic diseases research. Genomics, 2013, 101, 75-85.	2.9	18
14	LDL receptor cDNA sequence analysis in familial hypercholesterolemia patients: 5 novel mutations with high prevalence in families originating from southern Italy. Human Mutation, 2001, 17, 433-433.	2.5	17
15	Unusual splice site mutations disrupt FANCA exon 8 definition. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2014, 1842, 1052-1058.	3.8	16
16	A common genetic background could explain early-onset Crohnâ€™s disease. Medical Hypotheses, 2012, 78, 520-522.	1.5	15
17	Novel NOD2 Mutation in Early-Onset Inflammatory Bowel Phenotype. Inflammatory Bowel Diseases, 2018, 24, 1204-1212.	1.9	13
18	Inflammation profile of four early onset Crohn patients. Gene, 2012, 493, 282-285.	2.2	12

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19	DEFB1 gene 5â€² untranslated region (UTR) polymorphisms in inflammatory bowel diseases. Clinics, 2012, 67, 395-398.	1.5	11
20	GNE-related thrombocytopenia: evidence for a mutational hotspot in the ADP/substrate domain of the GNE bifunctional enzyme. Haematologica, 2022, 107, 750-754.	3.5	11
21	Genetic and Functional Profiling of Crohn's Disease: Autophagy Mechanism and Susceptibility to Infectious Diseases. BioMed Research International, 2013, 2013, 1-11.	1.9	10
22	A G to C transversion at the last nucleotide of exon 25 of the MYH9 gene results in a missense mutation rather than in a splicing defect. European Journal of Medical Genetics, 2010, 53, 256-260.	1.3	9
23	Novel Missense Mutation in the NOD2 Gene in a Patient with Early Onset Ulcerative Colitis: Causal or Chance Association?. International Journal of Molecular Sciences, 2014, 15, 3834-3841.	4.1	9
24	Neither hereditary periodic fever nor periodic fever, aphthae, pharingitis, adenitis: Undifferentiated periodic fever in a tertiary pediatric center. World Journal of Clinical Pediatrics, 2018, 7, 49-55.	2.1	9
25	Protective Role of BST2 Polymorphisms in Mother-to-Child Transmission of HIV-1 and Adult AIDS Progression. Journal of Acquired Immune Deficiency Syndromes (1999), 2016, 72, 237-241.	2.1	8
26	Altered pattern of tumor necrosis factor-alpha production in peripheral blood monocytes from Crohn's disease. World Journal of Gastroenterology, 2016, 22, 9117.	3.3	7
27	The Genetic Diagnosis of Ultrarare DEEs: An Ongoing Challenge. Genes, 2022, 13, 500.	2.4	7
28	Family history in early-onset inflammatory bowel disease. Journal of Gastroenterology, 2013, 48, 144-144.	5.1	5
29	Putative modifier genes in mevalonate kinase deficiency. Molecular Medicine Reports, 2016, 13, 3181-3189.	2.4	4
30	Fever tree revisited: From malaria to autoinflammatory diseases. World Journal of Clinical Pediatrics, 2015, 4, 106.	2.1	4
31	The effect of clodronate on a mevalonate kinase deficiency cellular model. Inflammation Research, 2012, 61, 1363-1367.	4.0	3
32	Could the <scp>MED13</scp> mutations manifest as a <scp>Kabuki</scp>â€like syndrome?. American Journal of Medical Genetics, Part A, 2021, 185, 584-590.	1.2	3
33	Letter to the Editor. Cell Biochemistry and Function, 2012, 30, 176-176.	2.9	2
34	Two-gene mutation in a single patient: Biochemical and functional analysis for a correct interpretation of exome results. Molecular Medicine Reports, 2015, 12, 6128-6132.	2.4	2
35	The Challenge of Next Generation Sequencing in a Boy With Severe Mononucleosis and EBV-related Lymphoma. Journal of Pediatric Hematology/Oncology, 2018, 40, e323-e326.	0.6	2
36	Familial hypogammaglobulinemia with high RTE and naÃ¯ve T lymphocytes. Inflammation Research, 2019, 68, 901-904.	4.0	2

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37	Serum amyloid A and cholesterol: a pivotal role on inflammation. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2012, 19, 163-164.	3.0	1
38	MYO5B Gene Mutations. Journal of Pediatric Gastroenterology and Nutrition, 2022, 74, .	1.8	1
39	Very-early onset IBD in male as expression of XIAP mutation. Digestive and Liver Disease, 2014, 46, e91.	0.9	0
40	The Glycosylation Nexus Enigma: new clues to the role of the GNE gene in the pathogenesis of inherited thrombocytopenia in both isolated and myopathy-associated forms. Hamostaseologie, 2021, 41, .	1.9	0