Marianne Antonius Jakobsen

List of Publications by Citations

27 papers	185 citations	9 h-index	13 g-index
28 ext. papers	228 ext. citations	4.2 avg, IF	2.96 L-index

#	Paper	IF	Citations
27	Peroxisome proliferator-activated receptor alpha, delta, gamma1 and gamma2 expressions are present in human monocyte-derived dendritic cells and modulate dendritic cell maturation by addition of subtype-specific ligands. *Scandinavian Journal of Immunology*, **2006**, 63, 330-7	3.4	38
26	Clinical characteristics and real-life diagnostic approaches in all Danish children with hereditary angioedema. *Orphanet Journal of Rare Diseases*, **2017**, 12, 55	4.2	19
25	Serum concentration of the growth medium markedly affects monocyte-derived dendritic cellsW phenotype, cytokine production profile and capacities to stimulate in MLR. *Scandinavian Journal of Immunology*, **2004**, 60, 584-91	3.4	15
24	Microchimerism of male origin in a cohort of Danish girls. *Chimerism*, **2015**, 6, 65-71		14
23	Testosterone treatment increases androgen receptor and aromatase gene expression in myotubes from patients with PCOS and controls, but does not induce insulin resistance. *Biochemical and Biophysical Research Communications*, **2014**, 451, 622-6	3.4	13
22	White blood cell mitochondrial DNA copy number is decreased in rheumatoid arthritis and linked with risk factors. A twin study. *Journal of Autoimmunity*, **2019**, 96, 142-146	15.5	12
21	The use of next-generation sequencing for the determination of rare blood group genotypes. *Transfusion Medicine*, **2019**, 29, 162-168	1.3	12
20	Genetical analysis of all Danish patients diagnosed with chronic granulomatous disease. *Scandinavian Journal of Immunology*, **2012**, 76, 505-11	3.4	11
19	Molecular pathways in patients with systemic lupus erythematosus revealed by gene-centred DNA sequencing. *Annals of the Rheumatic Diseases*, **2021**, 80, 109-117	2.4	10
18	Next Generation Sequencing-Based Fetal ABO Blood Group Prediction by Analysis of Cell-Free DNA from Maternal Plasma. *Transfusion Medicine and Hemotherapy*, **2020**, 47, 45-53	4.2	9
17	STK4 Deficiency Impairs Innate Immunity and Interferon Production Through Negative Regulation of TBK1-IRF3 Signaling. *Journal of Clinical Immunology*, **2021**, 41, 109-124	5.7	7
16	Male origin microchimerism and ovarian cancer. *International Journal of Epidemiology*, **2021**, 50, 87-94	7.8	6
15	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches. *PLoS ONE*, **2019**, 14, e0224858	3.7	4
14	A case of high-titer anti-D hemolytic disease of the newborn in which late onset and mild course is associated with the D variant, RHD-CE(9)-D. *Transfusion*, **2014**, 54, 2463-7	2.9	4
13	Impact of RHD genotyping on transfusion practice in Denmark and the United States and identification of novel RHD alleles. *Transfusion*, **2021**, 61, 256-265	2.9	3
12	Detection of non-DeltaGT NCF-1 mutations in chronic granulomatous disease. *Genetic Testing and Molecular Biomarkers*, **2009**, 13, 505-10	1.6	2
11	Characteristics of patients with familial Mediterranean fever in Denmark: a retrospective nationwide register-based cohort study. *Scandinavian Journal of Rheumatology*, **2020**, 49, 489-497	1.9	2

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10	Results of noninvasive prenatal RHD testing in Gestation Week 25 are not affected by maternal body mass index. *Transfusion*, **2018**, 58, 2421-2425	2.9	2
9	Altered Antibody Response to Epstein-Barr Virus in Patients With Rheumatoid Arthritis and Healthy Subjects Predisposed to the Disease. A Twin Study. *Frontiers in Immunology*, **2021**, 12, 650713	8.4	1
8	Pyrin Inflammasome Activation Abrogates Interleukin-1 Receptor Antagonist, Suggesting a New Mechanism Underlying Familial Mediterranean Fever Pathogenesis. *Arthritis and Rheumatology*, **2021**, 73, 2116-2126	9.5	1
7	A novel ABO allele with a 21-bp duplication identified in two unrelated European individuals with weak A expression. *Transfusion Medicine*, **2020**, 30, 508-512	1.3	
6	Acquired complement C1 esterase inhibitor deficiency in a patient with a rare variant with unknown significance. *BMJ Case Reports*, **2019**, 12,	0.9	
5	CD18 is redundant for the response to multiple vaccines: A case study. *Pediatric Allergy and Immunology*, **2019**, 30, 136-139	4.2	
4	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
3	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
2	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		
1	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches **2019**, 14, e0224858		