Marianne Antonius Jakobsen

List of Publications by Year
in descending order

27 papers	283 citations	840585 11 h-index	940416 16 g-index
28 all docs	28 docs citations	28 times ranked	634 citing authors

#	Article	IF	Citations
1	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, 330-337.	1.3	43
2	Molecular pathways in patients with systemic lupus erythematosus revealed by gene-centred DNA sequencing. Annals of the Rheumatic Diseases, 2021, 80, 109-117.	0.5	35
3	Clinical characteristics and real-life diagnostic approaches in all Danish children with hereditary angioedema. Orphanet Journal of Rare Diseases, 2017, 12, 55.	1.2	25
4	The use of nextâ€generation sequencing for the determination of rare blood group genotypes. Transfusion Medicine, 2019, 29, 162-168.	0.5	21
5	Microchimerism of male origin in a cohort of Danish girls. Chimerism, 2015, 6, 65-71.	0.7	18
6	Serum Concentration of the Growth Medium Markedly Affects Monocyte-Derived Dendritic Cells' Phenotype, Cytokine Production Profile and Capacities to Stimulate in MLR. Scandinavian Journal of Immunology, 2004, 60, 584-591.	1.3	17
7	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, 622-626.	1.0	17
8	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.	3.0	16
9	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.	0.7	16
10	STK4 Deficiency Impairs Innate Immunity and Interferon Production Through Negative Regulation of TBK1-IRF3 Signaling. Journal of Clinical Immunology, 2021, 41, 109-124.	2.0	16
11	Genetical Analysis of All <scp>D</scp>anish Patients Diagnosed with Chronic Granulomatous Disease. Scandinavian Journal of Immunology, 2012, 76, 505-511.	1.3	14
12	Impact of <scp>RHD</scp> genotyping on transfusion practice in <scp>Denmark</scp> and the <scp>United States</scp> and identification of novel <scp>RHD</scp> alleles. Transfusion, 2021, 61, 256-265.	0.8	9
13	Genetic susceptibility to angiotensin-converting enzyme-inhibitor induced angioedema: A systematic review and evaluation of methodological approaches. PLoS ONE, 2019, 14, e0224858.	1.1	8
14	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.	2.2	7
15	Male origin microchimerism and ovarian cancer. International Journal of Epidemiology, 2021, 50, 87-94.	0.9	6
16	A case of highâ€titer antiâ€<scp>D</scp> hemolytic disease of the newborn in which late onset and mild course is associated with the <scp>D</scp> variant, <scp>RHDâ€CE</scp>(9)â€<scp>D</scp>. Transfusion, 2014, 54, 2463-2467.	0.8	4
17	Characteristics of patients with familial Mediterranean fever in Denmark: a retrospective nationwide register-based cohort study. Scandinavian Journal of Rheumatology, 2020, 49, 489-497.	0.6	3
18	Pyrin Inflammasome Activation Abrogates Interleukinâ€1 Receptor Antagonist, Suggesting a New Mechanism Underlying Familial Mediterranean Fever Pathogenesis. Arthritis and Rheumatology, 2021, 73, 2116-2126.	2.9	3

#	Article	IF	Citations
19	Detection of Non-Î”GTNCF-1Mutations in Chronic Granulomatous Disease. Genetic Testing and Molecular Biomarkers, 2009, 13, 505-510.	0.3	2
20	Results of noninvasive prenatal RHD testing in Gestation Week 25 are not affected by maternal body mass index. Transfusion, 2018, 58, 2421-2425.	0.8	2
21	Acquired complement C1 esterase inhibitor deficiency in a patient with a rare SERPING1 variant with unknown significance. BMJ Case Reports, 2019, 12, e231122.	0.2	1
22	CD18 is redundant for the response to multiple vaccines: A case study. Pediatric Allergy and Immunology, 2019, 30, 136-139.	1.1	0
23	A novel ABO allele with a 21â€bp duplication identified in two unrelated European individuals with weak A expression. Transfusion Medicine, 2020, 30, 508-512.	0.5	0
24	Title is missing!. , 2019, 14, e0224858.		0
25	Title is missing!. , 2019, 14, e0224858.		0
26	Title is missing!. , 2019, 14, e0224858.		0
27	Title is missing!. , 2019, 14, e0224858.		0