

Minna K Karjalainen

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

26 papers	628 citations	14 h-index	25 g-index
28 ext. papers	810 ext. citations	6.7 avg, IF	3.15 L-index

#	Paper	IF	Citations
26	Integrative genetic, genomic and transcriptomic analysis of heat shock protein and nuclear hormone receptor gene associations with spontaneous preterm birth. <i>Scientific Reports</i> , 2021 , 11, 17115	4.9	2
25	Apolipoprotein A-I concentrations and risk of coronary artery disease: A Mendelian randomization study. <i>Atherosclerosis</i> , 2020 , 299, 56-63	3.1	27
24	Risk of spontaneous preterm birth and fetal growth associates with fetal SLIT2. <i>PLoS Genetics</i> , 2019 , 15, e1008107	6	13
23	Immature surfactant protein-B impairs the antioxidant capacity of HDL. <i>International Journal of Cardiology</i> , 2019 , 285, 53-58	3.2	5
22	Spontaneous premature birth as a target of genomic research. <i>Pediatric Research</i> , 2019 , 85, 422-431	3.2	14
21	NHLRC2 variants identified in patients with fibrosis, neurodegeneration, and cerebral angiomas (FINCA): characterisation of a novel cerebropulmonary disease. <i>Acta Neuropathologica</i> , 2018 , 135, 727-742	4.3	12
20	NKG2D gene variation and susceptibility to viral bronchiolitis in childhood. <i>Pediatric Research</i> , 2018 , 84, 451-457	3.2	3
19	Whole exome sequencing reveals HSPA1L as a genetic risk factor for spontaneous preterm birth. <i>PLoS Genetics</i> , 2018 , 14, e1007394	6	25
18	Expression of CPPED1 in human trophoblasts is associated with timing of term birth. <i>Journal of Cellular and Molecular Medicine</i> , 2018 , 22, 968-981	5.6	6
17	Genome-Wide Association Study of Polymorphisms Predisposing to Bronchiolitis. <i>Scientific Reports</i> , 2017 , 7, 41653	4.9	19
16	Mitochondrial hearing loss mutations among Finnish preterm and term-born infants. <i>Audiology Research</i> , 2017 , 7, 189	1.5	3
15	Haplotype of the Interleukin 17A gene is associated with osteitis after Bacillus Calmette-Guerin vaccination. <i>Scientific Reports</i> , 2017 , 7, 11691	4.9	10
14	Genome-wide association study of bronchopulmonary dysplasia: a potential role for variants near the CRP gene. <i>Scientific Reports</i> , 2017 , 7, 9271	4.9	11
13	Genetic Associations with Gestational Duration and Spontaneous Preterm Birth. <i>New England Journal of Medicine</i> , 2017 , 377, 1156-1167	59.2	183
12	CDHR3 gene variation and childhood bronchiolitis. <i>Journal of Allergy and Clinical Immunology</i> , 2017 , 140, 1469-1471.e7	11.5	7
11	Genes Encoding Vascular Endothelial Growth Factor A (VEGF-A) and VEGF Receptor 2 (VEGFR-2) and Risk for Bronchopulmonary Dysplasia. <i>Neonatology</i> , 2015 , 108, 53-9	4	17
10	CXCR3 Polymorphism and Expression Associate with Spontaneous Preterm Birth. <i>Journal of Immunology</i> , 2015 , 195, 2187-98	5.3	20

9	Polymorphisms of the gene encoding Kit ligand are associated with bronchopulmonary dysplasia. <i>Pediatric Pulmonology</i> , 2015 , 50, 260-270	3.5	3
8	A study of genes encoding cytokines (IL6, IL10, TNF), cytokine receptors (IL6R, IL6ST), and glucocorticoid receptor (NR3C1) and susceptibility to bronchopulmonary dysplasia. <i>BMC Medical Genetics</i> , 2014 , 15, 120	2.1	24
7	The genomics of preterm birth: from animal models to human studies. <i>Genome Medicine</i> , 2013 , 5, 34	14.4	70
6	IL10 polymorphisms, rhinovirus-induced bronchiolitis, and childhood asthma. <i>Journal of Allergy and Clinical Immunology</i> , 2013 , 131, 249-50	11.5	6
5	A potential novel spontaneous preterm birth gene, AR, identified by linkage and association analysis of X chromosomal markers. <i>PLoS ONE</i> , 2012 , 7, e51378	3.7	28
4	A study of collectin genes in spontaneous preterm birth reveals an association with a common surfactant protein D gene polymorphism. <i>Pediatric Research</i> , 2012 , 71, 93-9	3.2	23
3	Mapping a new spontaneous preterm birth susceptibility gene, IGF1R, using linkage, haplotype sharing, and association analysis. <i>PLoS Genetics</i> , 2011 , 7, e1001293	6	52
2	Genetic association of SP-C with duration of preterm premature rupture of fetal membranes and expression in gestational tissues. <i>Annals of Medicine</i> , 2009 , 41, 629-42	1.5	14
1	Haplotype analysis of ABCA3: association with respiratory distress in very premature infants. <i>Annals of Medicine</i> , 2008 , 40, 56-65	1.5	30