

Jukka Partanen

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.

212
papers

7,763
citations

49
h-index

80
g-index

225
ext. papers

8,694
ext. citations

5
avg, IF

5.16
L-index

#	Paper	IF	Citations
212	KIR gene content imputation from single-nucleotide polymorphisms in the Finnish population.. <i>PeerJ</i> , 2022 , 10, e12692	3.1	
211	How Communicating Polygenic and Clinical Risk for Atherosclerotic Cardiovascular Disease Impacts Health Behavior: an Observational Follow-up Study.. <i>Circulation Genomic and Precision Medicine</i> , 2022 , CIRCGEN121003459	5.2	4
210	PeptiCHIP: A Microfluidic Platform for Tumor Antigen Landscape Identification. <i>ACS Nano</i> , 2021 , 15, 15992-16010	10.1	10
209	Targeted RNA-Based Oxford Nanopore Sequencing for Typing 12 Classical HLA Genes. <i>Frontiers in Genetics</i> , 2021 , 12, 635601	4.5	1
208	Low ferritin levels appear to be associated with worsened health in male repeat blood donors. <i>Vox Sanguinis</i> , 2021 , 116, 1042-1050	3.1	0
207	Structural dissimilarity of partners' immune genes increases sperm viability in women's reproductive tract. <i>Journal of Evolutionary Biology</i> , 2021 , 34, 1125-1132	2.3	1
206	HLA RNA Sequencing With Unique Molecular Identifiers Reveals High Allele-Specific Variability in mRNA Expression. <i>Frontiers in Immunology</i> , 2021 , 12, 629059	8.4	0
205	Meta-Analysis of Genome-Wide Association and Gene Expression Studies Implicates Donor T Cell Function and Cytokine Pathways in Acute GvHD. <i>Frontiers in Immunology</i> , 2020 , 11, 19	8.4	3
204	FinDonor 10 000 study: a cohort to identify iron depletion and factors affecting it in Finnish blood donors. <i>Vox Sanguinis</i> , 2020 , 115, 36-46	3.1	7
203	Increasing accuracy of HLA imputation by a population-specific reference panel in a FinnGen biobank cohort. <i>NAR Genomics and Bioinformatics</i> , 2020 , 2, lqaa030	3.7	6
202	Gamete-level immunogenetic incompatibility in humans-towards deeper understanding of fertilization and infertility?. <i>Heredity</i> , 2020 , 125, 281-289	3.6	3
201	Review of Genetic Variation as a Predictive Biomarker for Chronic Graft-Versus-Host-Disease After Allogeneic Stem Cell Transplantation. <i>Frontiers in Immunology</i> , 2020 , 11, 575492	8.4	5
200	Post-copulatory genetic matchmaking: HLA-dependent effects of cervical mucus on human sperm function. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2020 , 287, 20201682	4.4	6
199	Increased MHC Matching by C4 Gene Compatibility in Unrelated Donor Hematopoietic Stem Cell Transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2019 , 25, 891-898	4.7	2
198	Attitudes of blood donors to their sample and data donation for biobanking. <i>European Journal of Human Genetics</i> , 2019 , 27, 1659-1667	5.3	8
197	Genomic prediction of relapse in recipients of allogeneic haematopoietic stem cell transplantation. <i>Leukemia</i> , 2019 , 33, 240-248	10.7	7
196	The effect of donation activity dwarfs the effect of lifestyle, diet and targeted iron supplementation on blood donor iron stores. <i>PLoS ONE</i> , 2019 , 14, e0220862	3.7	3

195	Computational Analysis of HLA-presentation of Non-synonymous Recipient Mismatches Indicates Effect on the Risk of Chronic Graft-vs.-Host Disease After Allogeneic HSCT. <i>Frontiers in Immunology</i> , 2019 , 10, 1625	8.4	6
194	Exploring rare and low-frequency variants in the Saguenay-Lac-Saint-Jean population identified genes associated with asthma and allergy traits. <i>European Journal of Human Genetics</i> , 2019 , 27, 90-101	5.3	5
193	Immunomonitoring of MSC-Treated GvHD Patients Reveals Only Moderate Potential for Response Prediction but Indicates Treatment Safety. <i>Molecular Therapy - Methods and Clinical Development</i> , 2018 , 9, 109-118	6.4	15
192	Blood donors' preferences for blood donation for biomedical research. <i>Transfusion</i> , 2018 , 58, 1640-1646	6.9	10
191	Hidden genomic MHC disparity between HLA-matched sibling pairs in hematopoietic stem cell transplantation. <i>Scientific Reports</i> , 2018 , 8, 5396	4.9	5
190	Haematopoietic stem cell transplantation induces severe dysbiosis in intestinal microbiota of paediatric ALL patients. <i>Bone Marrow Transplantation</i> , 2017 , 52, 1479-1482	4.4	7
189	Genetic polymorphism related to monocyte-macrophage function is associated with graft-versus-host disease. <i>Scientific Reports</i> , 2017 , 7, 15666	4.9	14
188	Accuracy of Programs for the Determination of Human Leukocyte Antigen Alleles from Next-Generation Sequencing Data. <i>Frontiers in Immunology</i> , 2017 , 8, 1815	8.4	11
187	Retrospective analysis of capillary hemoglobin recovery in nearly 1 200 000 blood donor returns. <i>Blood Advances</i> , 2017 , 1, 961-967	7.8	7
186	In vitro Treg expansion favors the full-length splicing isoform of CTLA4. <i>Immunotherapy</i> , 2016 , 8, 541-553	3.8	
185	Graft Immune Cell Composition Associates with Clinical Outcome of Allogeneic Hematopoietic Stem Cell Transplantation in Patients with AML. <i>Frontiers in Immunology</i> , 2016 , 7, 523	8.4	18
184	Glycosylation pattern of anti-platelet IgG is stable during pregnancy and predicts clinical outcome in alloimmune thrombocytopenia. <i>British Journal of Haematology</i> , 2016 , 174, 310-20	4.5	62
183	Conflicting HLA assignment by three different typing methods due to the apparent loss of heterozygosity in the MHC region. <i>Hla</i> , 2016 , 87, 350-5	1.9	3
182	Medium-high resolution electrochemical genotyping of HLA-DQ2/DQ8 for detection of predisposition to coeliac disease. <i>Analytical and Bioanalytical Chemistry</i> , 2014 , 406, 2757-69	4.4	8
181	Donor Haplotype B of NK KIR Receptor Reduces the Relapse Risk in HLA-Identical Sibling Hematopoietic Stem Cell Transplantation of AML Patients. <i>Frontiers in Immunology</i> , 2014 , 5, 405	8.4	35
180	Gliadin antibodies in older population and neurological and psychiatric disorders. <i>Acta Neurologica Scandinavica</i> , 2013 , 127, 19-25	3.8	3
179	Lectin from <i>Erythrina cristagalli</i> supports undifferentiated growth and differentiation of human pluripotent stem cells. <i>Stem Cells and Development</i> , 2013 , 22, 707-16	4.4	15
178	Multicenter analyses demonstrate significant clinical effects of minor histocompatibility antigens on GvHD and GvL after HLA-matched related and unrelated hematopoietic stem cell transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2013 , 19, 1244-53	4.7	70

177	Minor histocompatibility antigens as determinants for graft-versus-host disease after allogeneic haematopoietic stem cell transplantation. <i>International Journal of Immunogenetics</i> , 2013 , 40, 495-501	2.3	11
176	The duodenal microbiota composition of adult celiac disease patients is associated with the clinical manifestation of the disease. <i>Inflammatory Bowel Diseases</i> , 2013 , 19, 934-41	4.5	119
175	Interaction with intestinal epithelial cells promotes an immunosuppressive phenotype in <i>Lactobacillus casei</i> . <i>PLoS ONE</i> , 2013 , 8, e78420	3.7	6
174	Toll-like receptor gene polymorphisms confer susceptibility to graft-versus-host disease in allogeneic hematopoietic stem cell transplantation. <i>Scandinavian Journal of Immunology</i> , 2012 , 76, 336-43	3.4	19
173	Persistent duodenal intraepithelial lymphocytosis despite a long-term strict gluten-free diet in celiac disease. <i>American Journal of Gastroenterology</i> , 2012 , 107, 1563-9	0.7	86
172	Killer-cell immunoglobulin-like receptor gene profile predicts good molecular response to dasatinib therapy in chronic myeloid leukemia. <i>Experimental Hematology</i> , 2012 , 40, 906-913.e1	3.1	17
171	Association study of FUT2 (rs601338) with celiac disease and inflammatory bowel disease in the Finnish population. <i>Tissue Antigens</i> , 2012 , 80, 488-93		65
170	Endomysial antibodies predict celiac disease irrespective of the titers or clinical presentation. <i>World Journal of Gastroenterology</i> , 2012 , 18, 2511-6	5.6	23
169	Low-medium resolution HLA-DQ2/DQ8 typing for coeliac disease predisposition analysis by colorimetric assay. <i>Analytical and Bioanalytical Chemistry</i> , 2012 , 403, 807-19	4.4	9
168	Persistently positive gliadin antibodies without transglutaminase antibodies in the elderly: gluten intolerance beyond coeliac disease. <i>Digestive and Liver Disease</i> , 2011 , 43, 772-8	3.3	5
167	Gluten-sensitive hypertransaminasemia in celiac disease: an infrequent and often subclinical finding. <i>American Journal of Gastroenterology</i> , 2011 , 106, 1689-96	0.7	31
166	IgA-class autoantibodies against neuronal transglutaminase, TG6 in celiac disease: no evidence for gluten dependency. <i>Clinica Chimica Acta</i> , 2011 , 412, 1187-90	6.2	18
165	Antibodies against deamidated gliadin peptides in early-stage celiac disease. <i>Journal of Clinical Gastroenterology</i> , 2011 , 45, 673-8	3	22
164	Serodiagnostic assays for celiac disease based on the open or closed conformation of the autoantigen, transglutaminase 2. <i>Journal of Clinical Immunology</i> , 2011 , 31, 436-42	5.7	14
163	Secretor genotype (FUT2 gene) is strongly associated with the composition of Bifidobacteria in the human intestine. <i>PLoS ONE</i> , 2011 , 6, e20113	3.7	172
162	The severity of acute Puumala hantavirus infection does not predict the long-term outcome of patients. <i>Nephron Clinical Practice</i> , 2010 , 116, c89-94		11
161	Intestinal transglutaminase 2 specific antibody deposits in non-responsive coeliac disease. <i>Digestive and Liver Disease</i> , 2010 , 42, 692-7	3.3	7
160	Immunogenetic characteristics of patients with autoimmune gastritis. <i>World Journal of Gastroenterology</i> , 2010 , 16, 354-8	5.6	13

159	Hippocampal sclerosis in refractory temporal lobe epilepsy is associated with gluten sensitivity. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2009 , 80, 626-30	5.5	27
158	Association of IL-10 and IL-10Rbeta gene polymorphisms with graft-versus-host disease after haematopoietic stem cell transplantation from an HLA-identical sibling donor. <i>BMC Immunology</i> , 2009 , 10, 24	3.7	23
157	Interspersed transcription chimeras: neglected pathological mechanism infiltrating gene accession queries?. <i>Journal of Biomedical Informatics</i> , 2009 , 42, 382-9	10.2	3
156	Cost-effective HLA typing with tagging SNPs predicts celiac disease risk haplotypes in the Finnish, Hungarian, and Italian populations. <i>Immunogenetics</i> , 2009 , 61, 247-56	3.2	47
155	The shared CTLA4-ICOS risk locus in celiac disease, IgA deficiency and common variable immunodeficiency. <i>Genes and Immunity</i> , 2009 , 10, 151-61	4.4	37
154	Donor-recipient mismatch for common gene deletion polymorphisms in graft-versus-host disease. <i>Nature Genetics</i> , 2009 , 41, 1341-4	36.3	75
153	Domestic and foreign donor candidates result in differential probability of matching minor histocompatibility antigens--relevance of selection for hematopoietic stem cell transplantation. <i>Tissue Antigens</i> , 2009 , 73, 236-41		1
152	Diagnosing mild enteropathy celiac disease: a randomized, controlled clinical study. <i>Gastroenterology</i> , 2009 , 136, 816-23	13.3	206
151	Association of genetic variation in inducible costimulator gene with outcome of kidney transplantation. <i>Transplantation</i> , 2009 , 87, 393-6	1.8	31
150	Genetic similarity of chromosome 6 between patients receiving hematopoietic stem cell transplantation and HLA matched sibling donors. <i>Haematologica</i> , 2009 , 94, 528-35	6.6	3
149	Full likelihood analysis of genetic risk with variable age at onset disease--combining population-based registry data and demographic information. <i>PLoS ONE</i> , 2009 , 4, e6836	3.7	3
148	Secretion of celiac disease autoantibodies after in vitro gliadin challenge is dependent on small-bowel mucosal transglutaminase 2-specific IgA deposits. <i>BMC Immunology</i> , 2008 , 9, 6	3.7	22
147	Functional network reconstruction reveals somatic stemness genetic maps and dedifferentiation-like transcriptome reprogramming induced by GATA2. <i>Stem Cells</i> , 2008 , 26, 1186-201	5.8	39
146	Association of chest radiography findings with host-related genetic factors in patients with nephropathia epidemica. <i>Scandinavian Journal of Infectious Diseases</i> , 2008 , 40, 254-8		14
145	Gluten-dependent small bowel mucosal transglutaminase 2-specific IgA deposits in overt and mild enteropathy coeliac disease. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2008 , 47, 436-42	2.8	51
144	Myosin IXB gene region and gluten intolerance: linkage to coeliac disease and a putative dermatitis herpetiformis association. <i>Journal of Medical Genetics</i> , 2008 , 45, 222-7	5.8	23
143	Lack of association between thrombosis-associated and cytokine candidate gene polymorphisms and acute rejection or vascular complications after kidney transplantation. <i>Nephrology Dialysis Transplantation</i> , 2008 , 23, 364-8	4.3	12
142	Performance of a new rapid whole blood coeliac test in adult patients with low prevalence of endomysial antibodies. <i>Digestive and Liver Disease</i> , 2007 , 39, 1057-63	3.3	31

141	T cell regeneration in pediatric allogeneic stem cell transplantation. <i>Bone Marrow Transplantation</i> , 2007 , 39, 149-56	4.4	25
140	Persistent small bowel mucosal villous atrophy without symptoms in coeliac disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2007 , 25, 1237-45	6.1	118
139	N-glycan structures and associated gene expression reflect the characteristic N-glycosylation pattern of human hematopoietic stem and progenitor cells. <i>Experimental Hematology</i> , 2007 , 35, 1279-92 ^{3.1}		46
138	Heme oxygenase 1 gene polymorphisms and outcome of renal transplantation. <i>International Journal of Immunogenetics</i> , 2007 , 34, 253-7	2.3	15
137	Geographic distribution of cervical cancer-associated human leucocyte antigens and cervical cancer incidence in Finland. <i>International Journal of STD and AIDS</i> , 2007 , 18, 672-9	1.4	8
136	Diagnosis of acute renal allograft rejection by analyzing whole blood mRNA expression of lymphocyte marker molecules. <i>Transplantation</i> , 2007 , 83, 791-8	1.8	25
135	Killer-cell immunoglobulin-like receptor ligand compatibility in the outcome of Finnish unrelated donor hematopoietic stem cell transplantation. <i>Transplant Immunology</i> , 2007 , 18, 62-6	1.7	7
134	Genetic variation in ICOS regulates mRNA levels of ICOS and splicing isoforms of CTLA4. <i>Molecular Immunology</i> , 2007 , 44, 1644-51	4.3	18
133	Resurrection of gliadin antibodies in coeliac disease. Deamidated gliadin peptide antibody test provides additional diagnostic benefit. <i>Scandinavian Journal of Gastroenterology</i> , 2007 , 42, 1428-33	2.4	68
132	Endomysial antibody-negative coeliac disease: clinical characteristics and intestinal autoantibody deposits. <i>Gut</i> , 2006 , 55, 1746-53	19.2	172
131	Transcriptional profiling reflects shared and unique characters for CD34+ and CD133+ cells. <i>Stem Cells and Development</i> , 2006 , 15, 839-51	4.4	28
130	Genetic background of type I protein C deficiency in Finland. <i>Thrombosis Research</i> , 2006 , 118, 603-9	8.2	8
129	The impact of donor cytokine gene polymorphisms on the incidence of cytomegalovirus infection after kidney transplantation. <i>Transplant Immunology</i> , 2006 , 16, 258-62	1.7	17
128	HLA-DRB1, -DQB1 alleles in head and neck carcinoma patients. <i>Tissue Antigens</i> , 2006 , 67, 237-40		8
127	Immunoglobulin A autoantibodies against transglutaminase 2 in the small intestinal mucosa predict forthcoming coeliac disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2006 , 24, 541-52	6.1	128
126	Diagnostic significance of HLA-DQ typing in patients with previous coeliac disease diagnosis based on histology alone. <i>Alimentary Pharmacology and Therapeutics</i> , 2006 , 24, 1395-402	6.1	28
125	Global gene expression profile of human cord blood-derived CD133+ cells. <i>Stem Cells</i> , 2006 , 24, 631-41	5.8	91
124	Celiac disease: from inflammation to atrophy: a long-term follow-up study. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2005 , 41, 44-8	2.8	41

123	Small-bowel mucosal transglutaminase 2-specific IgA deposits in coeliac disease without villous atrophy: a prospective and randomized clinical study. <i>Scandinavian Journal of Gastroenterology</i> , 2005 , 40, 564-72	2.4	124
122	Genetic diversity of KIR natural killer cell markers in populations from France, Guadeloupe, Finland, Senegal and Réunion. <i>Tissue Antigens</i> , 2005 , 66, 267-76		88
121	The DR4-DQ8 haplotype and a specific T cell receptor Vbeta T cell subset are associated with absence of allergy to Can f 1. <i>Clinical and Experimental Allergy</i> , 2005 , 35, 797-803	4.1	18
120	Collection of autologous blood for bone marrow donation: how useful is it?. <i>Bone Marrow Transplantation</i> , 2005 , 35, 1035-9	4.4	17
119	Cytokine gene polymorphisms and genetic association with coeliac disease in the Finnish population. <i>Scandinavian Journal of Immunology</i> , 2005 , 61, 51-6	3.4	20
118	Protein S gene polymorphisms Pro626 and nt2698--no correlation to free protein S levels or protein S activities. <i>Thrombosis and Haemostasis</i> , 2005 , 94, 1340-1	7	3
117	T cell epitope-containing peptides of the major dog allergen Can f 1 as candidates for allergen immunotherapy. <i>Journal of Immunology</i> , 2005 , 175, 3614-20	5.3	36
116	Candidate gene region 2q33 in European families with coeliac disease. <i>Tissue Antigens</i> , 2004 , 63, 212-22		38
115	Characterization a novel HLA-B40 allele with serological Bw4 motif, HLA-B*4047, in the Finnish population and confirmation of B*270503 allele. <i>Tissue Antigens</i> , 2004 , 63, 595-7		8
114	Genetic association of coeliac disease susceptibility to polymorphisms in the ICOS gene on chromosome 2q33. <i>Genes and Immunity</i> , 2004 , 5, 85-92	4.4	48
113	High birth weight is associated with human leukocyte antigen (HLA) DRB1*13 in full-term infants. <i>International Journal of Immunogenetics</i> , 2004 , 31, 21-6		19
112	Villous tip intraepithelial lymphocytes as markers of early-stage coeliac disease. <i>Scandinavian Journal of Gastroenterology</i> , 2004 , 39, 428-33	2.4	87
111	Cytokine gene polymorphisms and risks of acute rejection and delayed graft function after kidney transplantation. <i>Transplantation</i> , 2004 , 78, 1422-8	1.8	61
110	Human leukocyte antigens B8-DRB1*03 in pediatric patients with nephropathia epidemica caused by Puumala hantavirus. <i>Pediatric Infectious Disease Journal</i> , 2004 , 23, 959-61	3.4	15
109	The association between mannan-binding lectin gene alleles and celiac disease. <i>American Journal of Gastroenterology</i> , 2003 , 98, 2808-9	0.7	9
108	Additional factor in some HLA DR3/DQ2 haplotypes confers a fourfold increased genetic risk of celiac disease. <i>Tissue Antigens</i> , 2003 , 61, 308-16		30
107	Meta and pooled analysis of European coeliac disease data. <i>European Journal of Human Genetics</i> , 2003 , 11, 828-34	5.3	65
106	Genetic susceptibility to variant Creutzfeldt-Jakob disease. <i>Lancet, The</i> , 2003 , 361, 447-8	4.0	2

105	A collaborative European search for non-DQA1*05-DQB1*02 celiac disease loci on HLA-DR3 haplotypes: analysis of transmission from homozygous parents. <i>Human Immunology</i> , 2003 , 64, 350-8	2.3	20
104	HLA types in celiac disease patients not carrying the DQA1*05-DQB1*02 (DQ2) heterodimer: results from the European Genetics Cluster on Celiac Disease. <i>Human Immunology</i> , 2003 , 64, 469-77	2.3	407
103	Elevation of IgG antibodies against tissue transglutaminase as a diagnostic tool for coeliac disease in selective IgA deficiency. <i>Gut</i> , 2003 , 52, 1567-71	19.2	127
102	HLA class II associated risk and protection against multiple sclerosis-a Finnish family study. <i>Journal of Neuroimmunology</i> , 2002 , 122, 140-5	3.5	64
101	CD80 (B7-1) and CD86 (B7-2) genes and genetic susceptibility to coeliac disease. <i>International Journal of Immunogenetics</i> , 2002 , 29, 331-3		1
100	A new locus for coeliac disease mapped to chromosome 15 in a population isolate. <i>Human Genetics</i> , 2002 , 111, 40-5	6.3	23
99	Genetic dissection between coeliac disease and dermatitis herpetiformis in sib pairs. <i>Annals of Human Genetics</i> , 2002 , 66, 387-392	2.2	21
98	AIRE mutations and human leukocyte antigen genotypes as determinants of the autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy phenotype. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002 , 87, 2568-74	5.6	147
97	Not all HLA DR3 DQ2 haplotypes confer equal susceptibility to coeliac disease: transmission analysis in families. <i>Scandinavian Journal of Gastroenterology</i> , 2002 , 37, 56-61	2.4	14
96	Human leukocyte antigen-B8-DR3 is a more important risk factor for severe Puumala hantavirus infection than the tumor necrosis factor-alpha(-308) G/A polymorphism. <i>Journal of Infectious Diseases</i> , 2002 , 186, 843-6	7	86
95	Celiac disease, thyrotoxicosis, and autoimmune hepatitis in a child. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2002 , 35, 90-2	2.8	15
94	Genetic dissection between silent and clinically diagnosed symptomatic forms of coeliac disease in multiplex families. <i>Digestive and Liver Disease</i> , 2002 , 34, 842-5	3.3	12
93	Celiac disease in patients with severe liver disease: gluten-free diet may reverse hepatic failure. <i>Gastroenterology</i> , 2002 , 122, 881-8	13.3	210
92	Celiac disease and HLA DQ in patients with IgA nephropathy. <i>American Journal of Gastroenterology</i> , 2002 , 97, 2572-6	0.7	53
91	Coeliac disease among healthy members of multiple case coeliac disease families. <i>Scandinavian Journal of Gastroenterology</i> , 2002 , 37, 161-5	2.4	46
90	HLA-DQ typing in the diagnosis of celiac disease. <i>American Journal of Gastroenterology</i> , 2002 , 97, 695-9	0.7	167
89	Genomewide linkage analysis of celiac disease in Finnish families. <i>American Journal of Human Genetics</i> , 2002 , 70, 51-9	11	80
88	Novel mutations in the human CYP21 gene. <i>Prenatal Diagnosis</i> , 2001 , 21, 885-9	3.2	13

87	Celiac disease without villous atrophy: revision of criteria called for. <i>Digestive Diseases and Sciences</i> , 2001 , 46, 879-87	4	133
86	Polymorphism of the cytokine genes in hospitalized patients with Puumala hantavirus infection. <i>Nephrology Dialysis Transplantation</i> , 2001 , 16, 1368-73	4.3	38
85	Candidate gene region 15q26 and genetic susceptibility to coeliac disease in Finnish families. <i>Scandinavian Journal of Gastroenterology</i> , 2001 , 36, 372-4	2.4	6
84	Candidate gene regions and genetic heterogeneity in gluten sensitivity. <i>Gut</i> , 2001 , 48, 696-701	19.2	39
83	Use of closely related affected individuals for the genetic study of complex diseases in founder populations. <i>American Journal of Human Genetics</i> , 2001 , 68, 154-159	11	38
82	Concordance of dermatitis herpetiformis and celiac disease in monozygous twins. <i>Journal of Investigative Dermatology</i> , 2000 , 115, 990-3	4.3	58
81	Autoimmunity to glutamic acid decarboxylase in patients with autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy (APECED). <i>Clinical and Experimental Immunology</i> , 2000 , 119, 419-25	6.2	9
80	Major histocompatibility complex (MHC)-linked microsatellite markers in a founder population. <i>Tissue Antigens</i> , 2000 , 56, 45-51		15
79	The HLA-DRB4 gene does not explain genetic susceptibility in HLA-DQ2-negative celiac disease. <i>Immunogenetics</i> , 2000 , 51, 249-51	3.2	5
78	Single Founder Mutation (W380G) in Type II Protein C Deficiency in Finland. <i>Thrombosis and Haemostasis</i> , 2000 , 84, 424-428	7	12
77	ss-cell autoantibodies, human leukocyte antigen II alleles, and type 1 diabetes in autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2000 , 85, 4434-40	5.6	60
76	Intolerance to cereals is not specific for coeliac disease. <i>Scandinavian Journal of Gastroenterology</i> , 2000 , 35, 942-6	2.4	97
75	Molecular characterization of two mutations in platelet glycoprotein (GP) Ib alpha in two Finnish Bernard-Soulier syndrome families. <i>European Journal of Haematology</i> , 1999 , 62, 160-8	3.8	11
74	Variant Bernard-Soulier syndrome due to homozygous Asn45Ser mutation in the platelet glycoprotein (GP) IX in seven patients of five unrelated Finnish families. <i>European Journal of Haematology</i> , 1999 , 62, 256-64	3.8	20
73	Expression of HSP-65 in jejunal epithelial cells in patients clinically suspected of coeliac disease. <i>Autoimmunity</i> , 1999 , 31, 125-32	3	14
72	HLA-DQ alleles and human papillomavirus DNA in adult-onset laryngeal papillomatosis. <i>Journal of Infectious Diseases</i> , 1999 , 179, 682-5	7	13
71	CD28/CTLA4 gene region on chromosome 2q33 confers genetic susceptibility to celiac disease. A linkage and family-based association study. <i>Tissue Antigens</i> , 1999 , 53, 470-5		110
70	Tracing past population migrations: genealogy of steroid 21-hydroxylase (CYP21) gene mutations in Finland. <i>European Journal of Human Genetics</i> , 1999 , 7, 188-96	5.3	15

69	Celiac disease and autoimmune endocrinologic disorders. <i>Digestive Diseases and Sciences</i> , 1999 , 44, 1428-33	4.3	61
68	Celiac disease and markers of celiac disease latency in patients with primary Sjögren's syndrome. <i>American Journal of Gastroenterology</i> , 1999 , 94, 1042-6	0.7	88
67	Genome scan for predisposing loci for distal interphalangeal joint osteoarthritis: evidence for a locus on 2q. <i>American Journal of Human Genetics</i> , 1999 , 65, 1060-7	11	106
66	Celiac disease and markers of celiac disease latency in patients with primary Sjögren's syndrome. <i>American Journal of Gastroenterology</i> , 1999 , 94, 1042-1046	0.7	69
65	Increased density of jejunal gammadelta+ T cells in patients having normal mucosa--marker of operative autoimmune mechanisms?. <i>Autoimmunity</i> , 1999 , 29, 179-87	3	35
64	Genetic polymorphism H131R of Fc gamma receptor type IIA (Fc gammaRIIA) in a healthy Finnish population and in patients with or without platelet-associated IgG. <i>European Journal of Haematology</i> , 1998 , 61, 183-9	3.8	11
63	TNF microsatellite alleles a2 and b3 are not primarily associated with celiac disease in the Finnish population. <i>Tissue Antigens</i> , 1998 , 51, 553-5		17
62	Genetic polymorphism in human platelet glycoprotein GP Ib/IX/V complex is enriched in GP V (CD42d). <i>Tissue Antigens</i> , 1998 , 52, 236-41		8
61	Genetic susceptibility to gluten sensitive enteropathy in Irish setter dogs is not linked to the major histocompatibility complex. <i>Tissue Antigens</i> , 1998 , 52, 543-9		33
60	The natural history of an HLA haplotype and its recombinants. <i>Immunogenetics</i> , 1998 , 48, 8-15	3.2	10
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2	HLA RNAseq reveals high allele-specific variability in mRNA expression		3
1	FinnGen: Unique genetic insights from combining isolated population and national health register data		11