Csaba Szalai

List of Publications by Year in descending order

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172207 2,653 96 29 citations h-index papers

47 g-index 100 100 100 3898 times ranked docs citations citing authors all docs

214527

#	Article	IF	CITATIONS
1	Co-Detection of VEGF-A and Its Regulator, microRNA-181a, May Indicate Central Nervous System Involvement in Pediatric Leukemia. Pathology and Oncology Research, 2022, 28, 1610096.	0.9	2
2	Two tagging singleâ€nucleotide polymorphisms to capture HLA ―DRB1*07:01–DQA1*02:01–DQB1*02:02 haplotype associated with asparaginase hypersensitivity. British Journal of Clinical Pharmacology, 2021, 87, 2542-2548.	1.1	5
3	Pharmacogenetics of the Central Nervous Systemâ€"Toxicity and Relapse Affecting the CNS in Pediatric Acute Lymphoblastic Leukemia. Cancers, 2021, 13, 2333.	1.7	1
4	Prevalence and characterization of severe asthma in Hungary. Scientific Reports, 2020, 10, 9274.	1.6	8
5	Investigation of circulating IncRNAs as potential biomarkers in chronic respiratory diseases. Journal of Translational Medicine, 2020, 18, 422.	1.8	18
6	MicroRNA-181a as novel liquid biopsy marker of central nervous system involvement in pediatric acute lymphoblastic leukemia. Journal of Translational Medicine, 2020, 18, 250.	1.8	19
7	Investigation of the Possible Role of Tie2 Pathway and TEK Gene in Asthma and Allergic Conjunctivitis. Frontiers in Genetics, 2020, $11,128.$	1.1	3
8	Plasma neutrophil extracellular trap level is modified by disease severity and inhaled corticosteroids in chronic inflammatory lung diseases. Scientific Reports, 2020, 10, 4320.	1.6	19
9	Pharmacogenetic Study of the Central Nervous System in Pediatric Acute Lymphoblastic Leukemia. Clinical Lymphoma, Myeloma and Leukemia, 2019, 19, S189-S190.	0.2	0
10	New Liquid Biopsy Markers for the Detection of Central Nervous System Involvement in Childhood Acute Leukemia. Clinical Lymphoma, Myeloma and Leukemia, 2019, 19, S187-S188.	0.2	0
11	Circulating microRNAs as Potential Minimal Residual Disease Biomarkers in Pediatric Acute Lymphoblastic Leukemia. Clinical Lymphoma, Myeloma and Leukemia, 2019, 19, S193.	0.2	0
12	Circulating microRNAs as minimal residual disease biomarkers in childhood acute lymphoblastic leukemia. Journal of Translational Medicine, 2019, 17, 372.	1.8	19
13	Variation in the <i><scp>TEK</scp></i> gene is not associated with asthma but with allergic conjunctivitis. International Journal of Immunogenetics, 2018, 45, 102-108.	0.8	13
14	Possible roles of genetic variations in chemotherapy related cardiotoxicity in pediatric acute lymphoblastic leukemia and osteosarcoma. BMC Cancer, 2018, 18, 704.	1.1	30
15	HLA-DRB1*07:01–HLA-DQA1*02:01–HLA-DQB1*02:02 haplotype is associated with a high risk of asparaginase hypersensitivity in acute lymphoblastic leukemia. Haematologica, 2017, 102, 1578-1586.	1.7	33
16	From genomes to diaries: a 3-year prospective, real-life study of ragweed-specific sublingual immunotherapy. Immunotherapy, 2017, 9, 1279-1294.	1.0	12
17	Rationale and design of the multiethnic Pharmacogenomics in Childhood Asthma consortium. Pharmacogenomics, 2017, 18, 931-943.	0.6	30
18	Investigation of the Possible Role of the Hippo/YAP1 Pathway in Asthma and Allergy. Allergy, Asthma and Immunology Research, 2017, 9, 247.	1.1	30

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19	Pharmacogenetic analysis of high-dose methotrexate treatment in children with osteosarcoma. Oncotarget, 2017, 8, 9388-9398.	0.8	33
20	Pharmacogenetics of anthracyclines. Pharmacogenomics, 2016, 17, 1075-1087.	0.6	16
21	Defining uncontrolled childhood asthma in the global PiCA consortium. , 2016, , .		О
22	VariantMetaCaller: automated fusion of variant calling pipelines for quantitative, precision-based filtering. BMC Genomics, 2015, 16, 875.	1.2	22
23	Subgroups of Paediatric Acute Lymphoblastic Leukaemia Might Differ Significantly in Genetic Predisposition to Asparaginase Hypersensitivity. PLoS ONE, 2015, 10, e0140136.	1.1	15
24	Impact of single nucleotide polymorphisms of cytarabine metabolic genes on drug toxicity in childhood acute lymphoblastic leukemia. Pediatric Blood and Cancer, 2015, 62, 622-628.	0.8	7
25	In interaction with gender a common CYP3A4 polymorphism may influence the survival rate of chemotherapy for childhood acute lymphoblastic leukemia. Pharmacogenomics Journal, 2015, 15, 241-247.	0.9	10
26	Novel genes in Human Asthma Based on a Mouse Model of Allergic Airway Inflammation and Human Investigations. Allergy, Asthma and Immunology Research, 2014, 6, 496.	1.1	22
27	Early repositioning through compound set enrichment analysis: a knowledge-recycling strategy. Future Medicinal Chemistry, 2014, 6, 563-575.	1.1	8
28	Complex analysis of multiple single nucleotide polymorphisms as putative risk factors of tooth agenesis in the Hungarian population. Acta Odontologica Scandinavica, 2014, 72, 216-227.	0.9	5
29	Associations of novel genetic variations in the folateâ€related and <i><scp>ARID</scp>5<scp>B</scp></i> genes with the pharmacokinetics and toxicity of highâ€dose methotrexate in paediatric acute lymphoblastic leukaemia. British Journal of Haematology, 2014, 166, 410-420.	1.2	34
30	Bayesian, Systems-based, Multilevel Analysis of Associations for Complex Phenotypes: from Interpretation to Decision., 2014, , 318-360.		4
31	Elevated Complement Factor H Levels in Asthmatic Sputa. Journal of Clinical Immunology, 2013, 33, 496-505.	2.0	8
32	Severe asthma database in Hungary, initial steps. Clinical and Translational Allergy, 2013, 3, P33.	1.4	1
33	Roles of Genetic Polymorphisms in the Folate Pathway in Childhood Acute Lymphoblastic Leukemia Evaluated by Bayesian Relevance and Effect Size Analysis. PLoS ONE, 2013, 8, e69843.	1.1	32
34	Implication of BIRC5 in asthma pathogenesis. International Immunology, 2012, 24, 293-301.	1.8	39
35	Asthma Endophenotypes and Polymorphisms in the Histamine Receptor <i>HRH4</i> Gene. International Archives of Allergy and Immunology, 2012, 159, 109-120.	0.9	23
36	980 Genetic Risk Factors of Neurotoxicity During Chemotherapy. European Journal of Cancer, 2012, 48, S236.	1.3	0

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37	1158 Candidate Gene Association Study in Childhood Acute Lymphoblastic Leukemia. European Journal of Cancer, 2012, 48, S278.	1.3	O
38	<i>ABCC1</i> polymorphisms in anthracyclineâ€induced cardiotoxicity in childhood acute lymphoblastic leukaemia. Cell Biology International, 2012, 36, 79-86.	1,4	100
39	Non-synonymous single nucleotide polymorphisms in genes for immunoregulatory galectins: Association of galectin-8 (F19Y) occurrence with autoimmune diseases in a Caucasian population. Biochimica Et Biophysica Acta - General Subjects, 2012, 1820, 1512-1518.	1.1	28
40	Candidate gene association study in pediatric acute lymphoblastic leukemia evaluated by Bayesian network based Bayesian multilevel analysis of relevance. BMC Medical Genomics, 2012, 5, 42.	0.7	41
41	Evaluation of a Partial Genome Screening of Two Asthma Susceptibility Regions Using Bayesian Network Based Bayesian Multilevel Analysis of Relevance. PLoS ONE, 2012, 7, e33573.	1.1	47
42	Complement Factor H Is Elevated In Sputum And Associated With Disease Severity In Asthma. , 2012, , .		0
43	Relationship between air pollution, NFE2L2 gene polymorphisms and childhood asthma in a Hungarian population. Journal of Community Genetics, 2012, 3, 25-33.	0.5	40
44	Rationale for an international consortium to study inherited genetic susceptibility to childhood acute lymphoblastic leukemia. Haematologica, 2011, 96, 1049-1054.	1.7	36
45	Meta-analysis of adrenocortical tumour genomics data: novel pathogenic pathways revealed. Oncogene, 2010, 29, 3163-3172.	2.6	66
46	Variation in CDKN2A at 9p21.3 influences childhood acute lymphoblastic leukemia risk. Nature Genetics, 2010, 42, 492-494.	9.4	248
47	Gene expression profiling of experimental asthma reveals a possible role of paraoxonase-1 in the disease. International Immunology, 2009, 21, 967-975.	1.8	35
48	Association of some rare haplotypes and genotype combinations in the MDR1 gene with childhood acute lymphoblastic leukaemia. Leukemia Research, 2008, 32, 1214-1220.	0.4	45
49	Asthma from a pharmacogenomic point of view. British Journal of Pharmacology, 2008, 153, 1602-1614.	2.7	24
50	Synergistic interaction of ABCB1 and ABCG2 polymorphisms predicts the prevalence of toxic encephalopathy during anticancer chemotherapy. Pharmacogenomics Journal, 2008, 8, 321-327.	0.9	50
51	Chlamydophila pneumoniae infection status is dependent on the subtypes of asthma and allergy. Allergy and Asthma Proceedings, 2007, 28, 58-63.	1.0	6
52	The HLA 8.1 ancestral haplotype is strongly linked to the C allele of â^'429T>C promoter polymorphism of receptor of the advanced glycation endproduct (RAGE) gene. Haplotype-independent association of the â^'429C allele with high hemoglobinA1C levels in diabetic patients. Molecular Immunology, 2007, 44, 648-655.	1.0	26
53	Imbalance of the C4A and C4B genes dosage as a robust risk factor for premature cardiovascular disease morbidity and mortality. Molecular Immunology, 2007, 44, 173.	1.0	0
54	Frequency of Carriers of 8.1 Ancestral Haplotype and its Fragments in Two Caucasian Populations. Immunological Investigations, 2007, 36, 307-319.	1.0	10

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55	CCR5î"32 mutation, Mycoplasma pneumoniae infection, and asthma. Journal of Allergy and Clinical Immunology, 2007, 119, 1545-1547.	1.5	19
56	Pediatric Asthmatic Patients Have Low Serum Levels of Monocyte Chemoattractant Protein-1. Journal of Asthma, 2006, 43, 399-404.	0.9	9
57	Genomic Investigation of Asthma in Human and Animal Models. , 2006, , 419-441.		0
58	The role of ABC-transporter gene polymorphisms in chemotherapy induced immunosuppression, a retrospective study in childhood acute lymphoblastic leukaemia. Cellular Immunology, 2006, 244, 121-124.	1.4	27
59	Involvement of TNFα –308A Promoter Polymorphism in the Development of Asthma in Children Infected With Chlamydophila pneumoniae. Pediatric Research, 2006, 60, 543-548.	1.1	13
60	Genomic Strategies in Pharmacology of Asthma and Autoimmunity. Anti-Inflammatory and Anti-Allergy Agents in Medicinal Chemistry, 2006, 5, 383-399.	1.1	2
61	Genetic basis of tobacco smoking: strong association of a specific major histocompatibility complex haplotype on chromosome 6 with smoking behavior. International Immunology, 2004, 16, 1507-1514.	1.8	29
62	Frequencies of two common mutations (c.35delG and c.167delT) of the connexin 26 gene in different populations of Hungary. International Journal of Molecular Medicine, 2004, 14, 1105.	1.8	1
63	Polymorphism in the promoter region of the apolipoprotein A5 gene is associated with an increased susceptibility for coronary artery disease. Atherosclerosis, 2004, 173, 109-114.	0.4	120
64	Relationship between the tumor necrosis factor alpha polymorphism and the serum C-reactive protein levels in inflammatory bowel disease. Immunogenetics, 2003, 55, 247-252.	1.2	63
65	The development of asthma in children infected with Chlamydia pneumoniae is dependent on the modifying effect of mannose-binding lectin. Journal of Allergy and Clinical Immunology, 2003, 112, 729-734.	1.5	73
66	Relationship between complement components C4A and C4B diversities and two TNFA promoter polymorphisms in two healthy Caucasian populations. Human Immunology, 2003, 64, 543-552.	1.2	15
67	Evolution of the thyrotropin receptor: a G protein coupled receptor with an intrinsic capacity to dimerize. Molecular Genetics and Metabolism, 2003, 78, 275-290.	0.5	27
68	Diversity in Intrinsic Strengths of the Human Complement System: Serum C4 Protein Concentrations Correlate with $<$ i>C4 $<$ i $>$ Gene Size and Polygenic Variations, Hemolytic Activities, and Body Mass Index. Journal of Immunology, 2003, 171, 2734-2745.	0.4	108
69	Histidine decarboxylase deficiency in gene knockout mice elevates male sex steroid production. Journal of Endocrinology, 2002, 175, 193-199.	1.2	27
70	The P28T Mutation in the GALK1 Gene Accounts for Galactokinase Deficiency in Roma (Gypsy) Patients across Europe. Pediatric Research, 2002, 51, 602-606.	1.1	40
71	Bone marrowâ€derived mast cell differentiation is strongly reduced in histidine decarboxylase knockout, histamineâ€free mice. International Immunology, 2002, 14, 381-387.	1.8	29
72	Histamine Genomics In Silico. Molecular Diagnosis and Therapy, 2002, 2, 67-72.	3.3	12

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73	Association of plasma lipid levels with apolipoprotein E polymorphism in Type 2 diabetes. Diabetes Research and Clinical Practice, 2002, 56, 63-68.	1.1	16
74	Lack of association between atopic eczema/dermatitis syndrome and polymorphisms in the promoter region of RANTES and regulatory region of MCP-1. Allergy: European Journal of Allergy and Clinical Immunology, 2002, 57, 160-163.	2.7	45
75	Differences in the genetic background of latent autoimmune diabetes in adults (LADA) and type 1 diabetes mellitus. Immunology Letters, 2002, 84, 109-115.	1.1	44
76	Polymorphism in the gene regulatory region of MCP-1 is associated with asthma susceptibility and severity. Journal of Allergy and Clinical Immunology, 2001, 108, 375-381.	1.5	156
77	Frequency of the R3500Q mutation of the apolipoprotein B-100 gene in a sample screened clinically for familial hypercholesterolemia in Hungary. Atherosclerosis, 2001, 154, 247-251.	0.4	20
78	The association of serum lipoprotein(a) levels, apolipoprotein(a) size and (TTTTA)n polymorphism with coronary heart disease. Clinica Chimica Acta, 2001, 309, 45-51.	0.5	23
79	Increased Frequency of the C3*F Allele and the Leiden Mutation of Coagulation FactorÂV in Patients with Severe Coronary Heart Disease Who Survived Myocardial Infarction. Experimental and Clinical Immunogenetics, 2001, 18, 206-212.	1.4	10
80	HEPATIC REGENERATION INDUCES TRANSIENT ACUTE PHASE REACTION: SYSTEMIC ELEVATION OF ACUTE PHASE REACTANTS AND SOLUBLE CYTOKINE RECEPTORS. Cell Biology International, 2001, 25, 585-592.	1.4	30
81	GP130-SPECIFIC ANTISENSE OLIGONUCLEOTIDES INHIBIT IL-6 SIGNAL INDUCING JUNB MRNA TRANSCRIPTION IN THE HUMAN HEPATOMA CELL LINE, HEPG2. Cell Biology International, 2001, 25, 835-840.	1.4	4
82	INTERLEUKIN-6-INDUCED PRODUCTION OF TYPE II ACUTE PHASE PROTEINS AND EXPRESSION OF junB GENE ARE DOWNREGULATED BY HUMAN RECOMBINANT GROWTH HORMONE IN VITRO. Cell Biology International, 2000, 24, 109-114.	1.4	24
83	Histidine Decarboxylase Expression in Human Melanoma. Journal of Investigative Dermatology, 2000, 115, 345-352.	0.3	61
84	Soluble interleukin-6 receptor (sIL-6R) makes IL-6R negative T cell line respond to IL-6; it inhibits TNF production. Immunology Letters, 2000, 71, 143-148.	1.1	19
85	Analysis of the genetic variability of the 1st (CCC/ACC, P52T) and the 10th exons (bp 1012-1704) of the TSH receptor gene in Graves' disease. International Journal of Immunogenetics, 2000, 27, 17-23.	1.2	29
86	Biosynthesis of interleukin-6, an autocrine growth factor for melanoma, is regulated by melanoma-derived histamine. Seminars in Cancer Biology, 2000, 10, 25-28.	4.3	19
87	H1 histamine receptor antagonist inhibits constitutive growth of Jurkat T cells and antigen-specific proliferation of ovalbumin-specific murine T cells. Seminars in Cancer Biology, 2000, 10, 41-45.	4.3	32
88	Influence of apolipoprotein E genotypes on serum lipid parameters in a biracial sample of children. European Journal of Pediatrics, 2000, 159, 257-260.	1.3	9
89	Angiotensin II type 1 receptor gene polymorphism and mitral valve prolapse syndrome. American Heart Journal, 2000, 139, 101-105.	1.2	28
90	Prevalence of CCR5î"32 in allergic diseases. Lancet, The, 2000, 355, 66.	6.3	10

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91	Exon–intron organization of the human gp130 gene. Gene, 2000, 243, 161-166.	1.0	4
92	Histamine and histamine-receptor antagonists modify gene expression and biosynthesis of interferon $\tilde{A}\check{Z}\hat{A}^3$ in peripheral human blood mononuclear cells and in CD19-depleted cell subsets. Immunology Letters, 1999, 70, 95-99.	1.1	38
93	Keratin 9 Mutations in the Coil 1A Region in Epidermolytic Palmoplantarâ€∫Keratoderma. Pediatric Dermatology, 1999, 16, 430-435.	0.5	17
94	The Histidine Decarboxylase (HDC) Gene of Tetrahymena Pyriformis is Similar to the Mammalian One. A Study of HDC Expression. Bioscience Reports, 1999, 19, 73-79.	1.1	9
95	Elevated hepatic glucocorticoid receptor expression during liver regeneration in rats. Pathology and Oncology Research, 1999, 5, 107-109.	0.9	13
96	Chemokine Receptor CCR2 and CCR5 Polymorphisms in Children with Insulin-Dependent Diabetes Mellitus. Pediatric Research, 1999, 46, 82-84.	1.1	49