Zoltan Prohaszka

List of Publications by Year
in descending order

281	9,255	47006 47	76900 74
papers	citations	h-index	g-index

304	304	304	11257
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Eculizumab use in a tertiary care nephrology center: data from the Vienna TMA cohort. Journal of Nephrology, 2022, 35, 451-461.	2.0	3
2	COVID-19: a trigger for severe thrombotic microangiopathy in a patient with complement gene variant. Romanian Journal of Internal Medicine = Revue Roumaine De Medecine Interne, 2022, .	0.6	2
3	Associations between the von Willebrand Factorâ€"ADAMTS13 Axis, Complement Activation, and COVID-19 Severity and Mortality. Thrombosis and Haemostasis, 2022, 122, 240-256.	3.4	15
4	Complement Genetics for the Practicing Allergist Immunologist: Focus on Complement Deficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2022, , .	3.8	1
5	Exploring red cell distribution width as a biomarker for treatment efficacy in home mechanical ventilation. BMC Pulmonary Medicine, 2022, 22, 115.	2.0	0
6	Complement Factor H-Related Proteins FHR1 and FHR5 Interact With Extracellular Matrix Ligands, Reduce Factor H Regulatory Activity and Enhance Complement Activation. Frontiers in Immunology, 2022, 13, 845953.	4.8	11
7	Decreased circulating dipeptidyl peptidase-4 enzyme activity is prognostic for severe outcomes in COVID-19 inpatients. Biomarkers in Medicine, 2022, 16, 317-330.	1.4	13
8	Serum fetuin-A level is independent of Helicobacter pylori postinfection status in systemic lupus erythematosus. Acta Microbiologica Et Immunologica Hungarica, 2022, , .	0.8	0
9	Complement Levels at Admission Reflecting Progression to Severe Acute Kidney Injury (AKI) in Coronavirus Disease 2019 (COVID-19): A Multicenter Prospective Cohort Study. Frontiers in Medicine, 2022, 9, 796109.	2.6	5
10	A Limited Course of Eculizumab in a Child with the Atypical Hemolytic Uremic Syndrome and Pre-B Acute Lymphoblastic Leukemia on Maintenance Therapy: Case Report and Literature Review. Journal of Clinical Medicine, 2022, 11, 2779.	2.4	5
11	Pregnancies in kidney transplant recipients with complement gene variant-mediated thrombotic microangiopathy. CKJ: Clinical Kidney Journal, 2021, 14, 1255-1260.	2.9	2
12	Immunogenic hotspots in the spacer domain of ADAMTS13 in immuneâ€mediated thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2021, 19, 478-488.	3.8	16
13	Soluble Vascular Biomarkers in Rheumatoid Arthritis and Ankylosing Spondylitis: Effects of 1-year Antitumor Necrosis Factor-Î± Therapy. Journal of Rheumatology, 2021, 48, 821-828.	2.0	5
14	Circulating Levels of Tissue Plasminogen Activator and Plasminogen Activator Inhibitor-1 Are Independent Predictors of Coronavirus Disease 2019 Severity: A Prospective, Observational Study. Seminars in Thrombosis and Hemostasis, 2021, 47, 451-455.	2.7	19
15	Acute heart transplantation from mechanical circulatory support in a human immunodeficiency virusâ€positive patient with fulminant myocarditis. ESC Heart Failure, 2021, 8, 1643-1648.	3.1	2
16	Complement Genetic Variants and FH Desialylation in S. pneumoniae-Haemolytic Uraemic Syndrome. Frontiers in Immunology, 2021, 12, 641656.	4.8	14
17	Complement Overactivation and Consumption Predicts In-Hospital Mortality in SARS-CoV-2 Infection. Frontiers in Immunology, 2021, 12, 663187.	4.8	87
18	Autoantibodies Against the Complement Regulator Factor H in the Serum of Patients With Neuromyelitis Optica Spectrum Disorder. Frontiers in Immunology, 2021, 12, 660382.	4.8	7

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19	Comparison of virus neutralization activity and results of 10 different anti-SARS-CoV-2 serological tests in COVID-19 recovered plasma donors. Practical Laboratory Medicine, 2021, 25, e00222.	1.3	10
20	Annual incidence and severity of acute episodes in hereditary thrombotic thrombocytopenic purpura. Blood, 2021, 137, 3563-3575.	1.4	31
21	A Novel Homozygous In-Frame Deletion in Complement Factor 3 Underlies Early-Onset Autosomal Recessive Atypical Hemolytic Uremic Syndrome - Case Report. Frontiers in Immunology, 2021, 12, 608604.	4.8	1
22	Atypical HUS and Crohnâ€™s diseaseâ€”interference of intestinal disease activity with complement-blocking treatment. Pediatric Nephrology, 2021, 36, 3277-3280.	1.7	0
23	Expanding Horizons in Complement Analysis and Quality Control. Frontiers in Immunology, 2021, 12, 697313.	4.8	16
24	FHR-5 Serum Levels and CFHR5 Genetic Variations in Patients With Immune Complex-Mediated Membranoproliferative Glomerulonephritis and C3-Glomerulopathy. Frontiers in Immunology, 2021, 12, 720183.	4.8	12
25	A pharmacokineticsâ€based approach to the monitoring of patient adherence to atorvastatin therapy. Pharmacology Research and Perspectives, 2021, 9, e00856.	2.4	2
26	Anti-ADAMTS13 autoantibody profiling in patients with immune-mediated thrombotic thrombocytopenic purpura. Blood Advances, 2021, 5, 3427-3435.	5.2	16
27	Complement multiplex testing: Concept, promises and pitfalls. Molecular Immunology, 2021, 140, 120-126.	2.2	2
28	The Role of Mannose-binding Lectin in Infectious Complications of Pediatric Hemato-Oncologic Diseases. Pediatric Infectious Disease Journal, 2021, 40, 154-158.	2.0	2
29	Case Report: A Case of COVID Vaccine-Induced Thrombotic Thrombocytopenia Manifested as Pulmonary Embolism and Hemorrhagia. A First Reported Case From Slovakia. Frontiers in Medicine, 2021, 8, 789972.	2.6	4
30	Case Report: Early Onset Systemic Lupus Erythematosus Due to Hereditary C1q Deficiency Treated With Fresh Frozen Plasma. Frontiers in Pediatrics, 2021, 9, 756387.	1.9	3
31	Validation of distinct pathogenic patterns in a cohort of membranoproliferative glomerulonephritis patients by cluster analysis. CKJ: Clinical Kidney Journal, 2020, 13, 225-234.	2.9	9
32	Preemptive plasma therapy prevents atypical hemolytic uremic syndrome relapse in kidney transplant recipients. European Journal of Internal Medicine, 2020, 73, 51-58.	2.2	7
33	Validation of Early Increase in Complement Activation Marker sC5b-9 as a Predictive Biomarker for the Development of Thrombotic Microangiopathy After Stem Cell Transplantation. Frontiers in Medicine, 2020, 7, 569291.	2.6	14
34	P0179ECULIZUMAB USE IN A TERTIARY CARE NEPHROLOGY CENTER. Nephrology Dialysis Transplantation, 2020, 35, .	0.7	0
35	Molecular basis and outcomes of atypical haemolytic uraemic syndrome in Czech children. European Journal of Pediatrics, 2020, 179, 1739-1750.	2.7	6
36	Successful Pregnancies During Ongoing Eculizumab Therapy in Two Patients With Complement-Mediated Thrombotic Microangiopathy. Kidney Medicine, 2020, 2, 213-217.	2.0	7

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37	Open ADAMTS13, induced by antibodies, is a biomarker for subclinical immune-mediated thrombotic thrombocytopenic purpura. Blood, 2020, 136, 353-361.	1.4	35
38	Sex Differences in Clinical Presentation and Outcomes among Patients with Complement-Gene-Variant-Mediated Thrombotic Microangiopathy. Journal of Clinical Medicine, 2020, 9, 964.	2.4	5
39	High-activity Classical and Alternative Complement Pathway Genotypesâ€"Association With Donor-specific Antibody-triggered Injury and Renal Allograft Survival. Transplantation Direct, 2020, 6, e534.	1.6	1
40	Dense deposit disease in an adolescent male mimicking acute post-streptococcal glomerulonephritis.. Hippokratia, 2020, 24, 191-193.	0.3	0
41	C4 nephritic factor in patients with immune-complex-mediated membranoproliferative glomerulonephritis and C3-glomerulopathy. Orphanet Journal of Rare Diseases, 2019, 14, 247.	2.7	10
42	A case report of a child with sepsis induced multiorgan failure and massive complement consumption treated with a short course of Eculizumab. Medicine (United States), 2019, 98, e14105.	1.0	11
43	Elevated Systemic Pentraxin-3 Is Associated With Complement Consumption in the Acute Phase of Thrombotic Microangiopathies. Frontiers in Immunology, 2019, 10, 240.	4.8	4
44	Decreased Ficolin-3-mediated Complement Lectin Pathway Activation and Alternative Pathway Amplification During Bacterial Infections in Patients With Type 2 Diabetes Mellitus. Frontiers in Immunology, 2019, 10, 509.	4.8	19
45	Pregnancy Outcome after Exposure to Migalastat for Fabry Disease: A Clinical Report. Case Reports in Obstetrics and Gynecology, 2019, 2019, 1-7.	0.3	0
46	Complement Markers in Blood and Urine: No Diagnostic Value in Late Silent Antibody-Mediated Rejection. Transplantation Direct, 2019, 5, e470.	1.6	4
47	Coexistence of aortic valve stenosis and cardiac amyloidosis: echocardiographic and clinical significance. Cardiovascular Ultrasound, 2019, 17, 32.	1.6	12
48	Novel Biomarkers in Cardiac Resynchronization Therapy: Hepatocyte Growth Factor Is an Independent Predictor of Clinical Outcome. Revista Espanola De Cardiologia (English Ed), 2019, 72, 48-55.	0.6	2
49	Association of Appendicitis, Helicobacter Pylori Positive Gastritis and Thrombotic Thrombocytopenic Purpura in an Adolescent. American Journal of Case Reports, 2019, 20, 131-133.	0.8	1
50	Hemolytic uremic syndrome complicating whooping cough. Srpski Arhiv Za Celokupno Lekarstvo, 2019, 147, 89-93.	0.2	0
51	P057â€...Effects of ANTI-TNF therapy on vascular biomarker levels in rheumatoid arthritis. , 2018, , .		0
52	Impact of intraoperative cytokine adsorption on outcome of patients undergoing orthotopic heart transplantationâ€"an observational study. Clinical Transplantation, 2018, 32, e13211.	1.6	53
53	Early Increase in Complement Terminal Pathway Activation Marker sC5b-9 Is Predictive for the Development of Thrombotic Microangiopathy after Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2018, 24, 989-996.	2.0	30
54	Role of complement in the pathogenesis of thrombotic microangiopathies. Memo - Magazine of European Medical Oncology, 2018, 11, 227-234.	0.5	8

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55	Maternal and Fetal Outcomes of Pregnancies in Women with Atypical Hemolytic Uremic Syndrome. Journal of the American Society of Nephrology: JASN, 2018, 29, 1020-1029.	6.1	56
56	Platelet Count, ADAMTS13 Activity, von Willebrand Factor Level and Survival in Patients with Colorectal Cancer: 5-Year Follow-up Study. Thrombosis and Haemostasis, 2018, 118, 123-131.	3.4	11
57	P824Novel biomarkers in cardiac resynchronization therapy: Hepatocyte growth factor is an independent predictor of clinical outcome. Europace, 2018, 20, i151-i151.	1.7	0
58	High serum Hsp70 level predicts poor survival in colorectal cancer: Results obtained in an independent validation cohort. Cancer Biomarkers, 2018, 23, 539-547.	1.7	16
59	SP007EARLY FETAL OUTCOME OF 28 PREGNANCIES IN WOMEN WITH ATYPICAL HEMOLYTIC UREMIC SYNDROME. Nephrology Dialysis Transplantation, 2018, 33, i348-i349.	0.7	0
60	Complement analysis in the era of targeted therapeutics. Molecular Immunology, 2018, 102, 84-88.	2.2	29
61	Concentration and Subclass Distribution of Anti-ADAMTS13 IgG Autoantibodies in Different Stages of Acquired Idiopathic Thrombotic Thrombocytopenic Purpura. Frontiers in Immunology, 2018, 9, 1646.	4.8	20
62	SP730PREEMPTIVE PLASMA THERAPY AND ECULIZUMAB RESCUE FOR ATYPICAL HEMOLYTIC UREMIC SYNDROME RELAPSE FOLLOWING KIDNEY TRANSPLANTATION. Nephrology Dialysis Transplantation, 2018, 33, i592-i593.	0.7	2
63	FRI0057â€…Effects of anti-tnf therapy on vascular biomarker levels in rheumatoid arthritis. , 2018, , .		0
64	Long-Term Survival and Apolipoprotein A1 Level in Chronic Heart Failure: Interaction With Tumor Necrosis Factor Î± â^'308 G/A Polymorphism. Journal of Cardiac Failure, 2017, 23, 113-120.	1.7	19
65	A unique haplotype of RCCX copy number variation: from the clinics of congenital adrenal hyperplasia to evolutionary genetics. European Journal of Human Genetics, 2017, 25, 702-710.	2.8	10
66	Decreased Neutrophil Extracellular Trap Degradation in Shiga Toxin-Associated Haemolytic Uraemic Syndrome. Journal of Innate Immunity, 2017, 9, 12-21.	3.8	28
67	The Phenotypic Spectrum of Nephropathies Associated with Mutations in Diacylglycerol Kinase Îµ. Journal of the American Society of Nephrology: JASN, 2017, 28, 3066-3075.	6.1	50
68	FHR-1 Binds to C-Reactive Protein and Enhances Rather than Inhibits Complement Activation. Journal of Immunology, 2017, 199, 292-303.	0.8	43
69	The role of the complement system in hereditary angioedema. Molecular Immunology, 2017, 89, 59-68.	2.2	35
70	First description of a rifampicin-resistant Neisseria meningitidis serogroup Y strain causing recurrent invasive meningococcal disease in Hungary. Acta Microbiologica Et Immunologica Hungarica, 2017, 64, 1-7.	0.8	6
71	Complement activation, inflammation and relative ADAMTS13 deficiency in secondary thrombotic microangiopathies. Immunobiology, 2017, 222, 119-127.	1.9	30
72	Circulating mitochondrial stress 70 protein/mortalin and cytosolic Hsp70 in blood: Risk indicators in colorectal cancer. International Journal of Cancer, 2017, 141, 2329-2335.	5.1	23

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73	The role of human leukocyte antigen DRB1-DQB1 haplotypes in the susceptibility to acquired idiopathic thrombotic thrombocytopenic purpura. Human Immunology, 2017, 78, 80-87.	2.4	16
74	Analysis of Linear Antibody Epitopes on Factor H and CFHR1 Using Sera of Patients with Autoimmune Atypical Hemolytic Uremic Syndrome. Frontiers in Immunology, 2017, 8, 302.	4.8	18
75	Functional Characterization of the Disease-Associated N-Terminal Complement Factor H Mutation W198R. Frontiers in Immunology, 2017, 8, 1800.	4.8	4
76	Carboxiterminal pro-endothelin-1 as an endothelial cell biomarker in thrombotic thrombocytopenic purpura. Thrombosis and Haemostasis, 2016, 115, 1034-1043.	3.4	3
77	Human Fetuin-A Rs4918 Polymorphism and its Association with Obesity in Healthy Persons and in Patients with Myocardial Infarction in Two Hungarian Cohorts. Medical Science Monitor, 2016, 22, 2742-2750.	1.1	5
78	Measurement of the Red Blood Cell Distribution Width Improves the Risk Prediction in Cardiac Resynchronization Therapy. Disease Markers, 2016, 2016, 1-13.	1.3	9
79	Association of Low Ficolinâ€“Lectin Pathway Parameters with Cardiac Syndrome X. Scandinavian Journal of Immunology, 2016, 84, 174-181.	2.7	1
80	Alternative complement pathway activation during invasive coronary procedures in acute myocardial infarction and stable angina pectoris. Clinica Chimica Acta, 2016, 463, 138-144.	1.1	3
81	Monomeric C-reactive protein inhibits renal cell-directed complement activation mediated by properdin. American Journal of Physiology - Renal Physiology, 2016, 310, F1308-F1316.	2.7	24
82	Complement C3a predicts outcome in cardiac resynchronization therapy of heart failure. Inflammation Research, 2016, 65, 933-940.	4.0	7
83	Complement analysis 2016: Clinical indications, laboratory diagnostics and quality control. Immunobiology, 2016, 221, 1247-1258.	1.9	77
84	Novel Vasoregulatory Aspects of Hereditary Angioedema: the Role of Arginine Vasopressin, Adrenomedullin and Endothelin-1. Journal of Clinical Immunology, 2016, 36, 160-170.	3.8	16
85	Genetic analysis and functional characterization of novel mutations in a series of patients with atypical hemolytic uremic syndrome. Molecular Immunology, 2016, 71, 10-22.	2.2	27
86	Atrial natriuretic peptide as a novel biomarker of hereditary angioedema. Clinical Immunology, 2016, 165, 45-46.	3.2	4
87	Heterogeneity but individual constancy of epitopes, isotypes and avidity of factor H autoantibodies in atypical hemolytic uremic syndrome. Molecular Immunology, 2016, 70, 47-55.	2.2	33
88	The role of mannose binding lectin on fever episodes in pediatric oncology patients. Pathology and Oncology Research, 2016, 22, 139-143.	1.9	5
89	The ratio of the neutrophil leucocytes to the lymphocytes predicts the outcome after cardiac resynchronization therapy. Europace, 2016, 18, 747-754.	1.7	20
90	Complement-Mediated Glomerular Injury in Children. , 2016, , 927-958.		0

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91	Association of Human Fetuin-A rs4917 Polymorphism with Obesity in 2 Cohorts. Journal of Investigative Medicine, 2015, 63, 548-553.	1.6	8
92	Comprehensive study into the activation of the plasma enzyme systems during attacks of hereditary angioedema due to C1-inhibitor deficiency. Orphanet Journal of Rare Diseases, 2015, 10, 132.	2.7	39
93	The use of a rapid fluorogenic neuraminidase assay to differentiate acute Streptococcus pneumoniae-associated hemolytic uremic syndrome (HUS) from other forms of HUS. Clinical Chemistry and Laboratory Medicine, 2015, 53, e117-9.	2.3	4
94	The Major Autoantibody Epitope on Factor H in Atypical Hemolytic Uremic Syndrome Is Structurally Different from Its Homologous Site in Factor H-related Protein 1, Supporting a Novel Model for Induction of Autoimmunity in This Disease. Journal of Biological Chemistry, 2015, 290, 9500-9510.	3.4	69
95	Functional characterization of two novel non-synonymous alterations in CD46 and a Q950H change in factor H found in atypical hemolytic uremic syndrome patients. Molecular Immunology, 2015, 65, 367-376.	2.2	24
96	Copeptin (C-terminal pro Arginine-Vasopressin) is an Independent Long-Term Prognostic Marker in Heart Failure with Reduced Ejection Fraction. Heart Lung and Circulation, 2015, 24, 359-367.	0.4	39
97	Increased circulating heat shock protein 70 (HSPA1A) levels in gestational diabetes mellitus: a pilot study. Cell Stress and Chaperones, 2015, 20, 575-581.	2.9	29
98	Structural Basis for the Function of Complement Component C4 within the Classical and Lectin Pathways of Complement. Journal of Immunology, 2015, 194, 5488-5496.	0.8	69
99	Inflammation and oxidative stress caused by nitric oxide synthase uncoupling might lead to left ventricular diastolic and systolic dysfunction in patients with hypertension. Journal of Geriatric Cardiology, 2015, 12, 1-10.	0.2	27
100	Common Genetic Variants of the Human Steroid 21-Hydroxylase Gene (CYP21A2) Are Related to Differences in Circulating Hormone Levels. PLoS ONE, 2014, 9, e107244.	2.5	12
101	AB0960â€...Autoantibodies to Heat-Shock Protein60 and Cardiovascular Disease in Patients with Mixed Connective Tissue Disease. Annals of the Rheumatic Diseases, 2014, 73, 1117.3-1118.	0.9	0
102	First-line therapy in atypical hemolytic uremic syndrome: consideration on infants with a poor prognosis. Italian Journal of Pediatrics, 2014, 40, 101.	2.6	8
103	Complement-Mediated Glomerular Injury in Children. , 2014, , 1-34.		0
104	Clinical usefulness of measuring red blood cell distribution width in patients with systemic sclerosis. Rheumatology, 2014, 53, 1439-1445.	1.9	31
105	Complement Activation and its Prognostic role in Postâ€cardiac Arrest Patients. Scandinavian Journal of Immunology, 2014, 79, 404-409.	2.7	12
106	Activation of the ficolin-lectin pathway during attacks ofÂhereditary angioedema. Journal of Allergy and Clinical Immunology, 2014, 134, 1388-1393.e1.	2.9	13
107	Efficacy of Eculizumab in a Patient With Immunoadsorption-Dependent Catastrophic Antiphospholipid Syndrome. Medicine (United States), 2014, 93, e143.	1.0	91
108	Elevated plasma neutrophil elastase concentration is associated with disease activity in patients with thrombotic thrombocytopenic purpura. Thrombosis Research, 2014, 133, 616-621.	1.7	26

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109	Evidence-based hydro- and balneotherapy in Hungaryâ€"a systematic review and meta-analysis. International Journal of Biometeorology, 2014, 58, 311-323.	3.0	68
110	Endothelial cell activation during edematous attacks of hereditary angioedema types I and II. Journal of Allergy and Clinical Immunology, 2014, 133, 1686-1691.	2.9	35
111	Autoantibodies to complement components in C3 glomerulopathy and atypical hemolytic uremic syndrome. Immunology Letters, 2014, 160, 163-171.	2.5	50
112	Increased levels of anti-heat-shock protein 60 (anti-Hsp60) indicate endothelial dysfunction, atherosclerosis and cardiovascular diseases in patients with mixed connective tissue disease. Immunologic Research, 2014, 60, 50-59.	2.9	12
113	Red cell distribution width as predictive marker in CHF: Testing of model performance by reclassification methods. International Journal of Cardiology, 2014, 174, 783-785.	1.7	7
114	A systematic analysis of the complement pathways in patients with neuromyelitis optica indicates alteration but no activation during remission. Molecular Immunology, 2014, 57, 200-209.	2.2	19
115	Update on the role of the complement system in the pathogenesis of thrombotic microangiopathies. Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki, 2014, 35, 115-22.	0.5	0
116	A rare case: childhood-onset C3 glomerulonephritis due to homozygous factor H deficiency. CEN Case Reports, 2013, 2, 234-238.	0.9	4
117	Elevated extracellular HSP70 (HSPA1A) level as an independent prognostic marker of mortality in patients with heart failure. Cell Stress and Chaperones, 2013, 18, 809-813.	2.9	34
118	Persistently elevated extracellular HSP70 (HSPA1A) level as an independent prognostic marker in post-cardiac-arrest patients. Cell Stress and Chaperones, 2013, 18, 447-454.	2.9	18
119	Elevated levels of mitochondrial mortalin and cytosolic HSP70 in blood as risk factors in patients with colorectal cancer. International Journal of Cancer, 2013, 133, 514-518.	5.1	49
120	The role of complement in Streptococcus pneumoniae-associated haemolytic uraemic syndrome. Nephrology Dialysis Transplantation, 2013, 28, 2237-2245.	0.7	70
121	Novel duplication in the F12 gene in a patient with recurrent angioedema. Clinical Immunology, 2013, 149, 142-145.	3.2	66
122	Serum concentration of immunoglobulin G-type antibodies against the whole Epsteinâ€"Barr nuclear antigen 1 and its aa35â€"58 or aa398â€"404 fragments in the sera of patients with systemic lupus erythematosus and multiple sclerosis. Clinical and Experimental Immunology, 2013, 171, 255-262.	2.6	13
123	Intraspecific Evolution of Human RCCX Copy Number Variation Traced by Haplotypes of the CYP21A2 Gene. Genome Biology and Evolution, 2013, 5, 98-112.	2.5	12
124	Both Positive and Negative Selection Pressures Contribute to the Polymorphism Pattern of the Duplicated Human CYP21A2 Gene. PLoS ONE, 2013, 8, e81977.	2.5	9
125	Association of Ficolin-3 with Severity and Outcome of Chronic Heart Failure. PLoS ONE, 2013, 8, e60976.	2.5	34
126	Serum Ghrelin Level and TNF-Î±/Ghrelin Ratio in Patients with Previous Myocardial Infarction. Archives of Medical Research, 2012, 43, 548-554.	3.3	10

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127	Consumption and dysregulation of complement classical and alternative pathways in patients with streptococcus pneumoniae-associated haemolytic uremic syndrome. Immunobiology, 2012, 217, 1219.	1.9	0
128	Atypical Hemolytic Uremic Syndrome-Associated Variants and Autoantibodies Impair Binding of Factor H and Factor H-Related Protein 1 to Pentraxin 3. Journal of Immunology, 2012, 189, 1858-1867.	0.8	62
129	OS004. A link between the complement system and angiogenic imbalance inpreeclampsia: ficolin-2 deficiency. Pregnancy Hypertension, 2012, 2, 177.	1.4	1
130	Complement anaphylatoxin C3a as a novel independent prognostic marker in heart failure. Clinical Research in Cardiology, 2012, 101, 607-615.	3.3	26
131	Anti-mutated citrullinated vimentin (anti-MCV) and anti-65kDa heat shock protein (anti-hsp65): New biomarkers in ankylosing spondylitis. Joint Bone Spine, 2012, 79, 63-66.	1.6	23
132	Complement activation in thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2012, 10, 791-798.	3.8	125
133	The use of â€˜realâ€timeâ€™ complement analysis to differentiate atypical haemolytic uraemic syndrome from other forms of thrombotic microangiopathies. British Journal of Haematology, 2012, 158, 424-425.	2.5	11
134	Circulating ficolin-2 and ficolin-3 in normal pregnancy and pre-eclampsia. Clinical and Experimental Immunology, 2012, 169, 49-56.	2.6	29
135	Comparison of epitope specificity of anti-heat shock protein 60/65 IgG type antibodies in the sera of healthy subjects, patients with coronary heart disease and inflammatory bowel disease. Cell Stress and Chaperones, 2012, 17, 215-227.	2.9	15
136	Endothelial Cell Function in Patients with Hereditary Angioedema: Elevated Soluble E-selectin Level During Inter-attack Periods. Journal of Clinical Immunology, 2012, 32, 61-69.	3.8	18
137	Circulating Chaperones in Health and Disease. Heat Shock Proteins, 2012, , 279-290.	0.2	0
138	Serum heat shock protein 70 levels in relation to circulating cytokines, chemokines, adhesion molecules and angiogenic factors in women with preeclampsia. Clinica Chimica Acta, 2011, 412, 1957-1962.	1.1	51
139	Fetuin-A serum levels in patients with aortic aneurysms of Marfan syndrome and atherosclerosis. European Journal of Clinical Investigation, 2011, 41, 176-182.	3.4	8
140	Serum fetuin-A in metabolic and inflammatory pathways in patients with myocardial infarction. European Journal of Clinical Investigation, 2011, 41, 703-709.	3.4	38
141	Diagnosis and Classification of Hemolytic Uremic Syndrome: The Hungarian Experience. Transplantation Proceedings, 2011, 43, 1247-1249.	0.6	7
142	Complement activation in animal and human pregnancies as a model for immunological recognition. Molecular Immunology, 2011, 48, 1621-1630.	2.2	71
143	Low ficolin-3 levels in early follow-up serum samples are associated with the severity and unfavorable outcome of acute ischemic stroke. Journal of Neuroinflammation, 2011, 8, 185.	7.2	39
144	Analysis of the 8.1 ancestral MHC haplotype in severe, pneumonia-related sepsis. Clinical Immunology, 2011, 139, 282-289.	3.2	6

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145	Serum soluble E-selectin and NT-proBNP levels additively predict mortality in diabetic patients with chronic heart failure. Clinical Research in Cardiology, 2011, 100, 587-594.	3.3	14
146	High levels of acute phase proteins and soluble 70ÂkDa heat shock proteins are independent and additive risk factors for mortality in colorectal cancer. Cell Stress and Chaperones, 2011, 16, 49-55.	2.9	22
147	Serum level of soluble Hsp70 is associated with vascular calcification. Cell Stress and Chaperones, 2011, 16, 257-265.	2.9	37
148	Serum leptin levels in relation to circulating cytokines, chemokines, adhesion molecules and angiogenic factors in normal pregnancy and preeclampsia. Reproductive Biology and Endocrinology, 2011, 9, 124.	3.3	109
149	Human anti-60 kD heat shock protein autoantibodies are characterized by basic features of natural autoantibodies. Acta Physiologica Hungarica, 2010, 97, 1-10.	0.9	12
150	The effect of long-term danazol prophylaxis on liver function in hereditary angioedemaâ€”a longitudinal study. European Journal of Clinical Pharmacology, 2010, 66, 419-426.	1.9	44
151	Serum level of soluble 70-kD heat shock protein is associated with high mortality in patients with colorectal cancer without distant metastasis. Cell Stress and Chaperones, 2010, 15, 143-151.	2.9	48
152	Increased circulating heat shock protein 70 levels in pregnant asthmatics. Cell Stress and Chaperones, 2010, 15, 295-300.	2.9	30
153	Circulating heat shock protein 70 (HSPA1A) in normal and pathological pregnancies. Cell Stress and Chaperones, 2010, 15, 237-247.	2.9	94
154	Functional analysis of the mannose-binding lectin complement pathway in normal pregnancy and preeclampsia. Journal of Reproductive Immunology, 2010, 87, 90-96.	1.9	18
155	Proinflammatory activation pattern of human umbilical vein endothelial cells induced by ILâ€1Î², TNFâ€Î±, and LPS. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2010, 77A, 962-970.	1.5	146
156	Plasma osteopontin concentrations in preeclampsia â€“ is there an association with endothelial injury?. Clinical Chemistry and Laboratory Medicine, 2010, 48, 181-187.	2.3	14
157	The R1141X Loss-of-Function Mutation of the ABCC6 Gene Is a Strong Genetic Risk Factor for Coronary Artery Disease. Genetic Testing and Molecular Biomarkers, 2010, 14, 75-78.	0.7	57
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