Andrzej Ciechanowicz

List of Publications by Year
in descending order

123	1,762	361413	330143
papers	citations	20 h-index	37 g-index

131	131	131	3159
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Capillary Blood Recovery Variables in Young Swimmers: An Observational Case Study. International Journal of Environmental Research and Public Health, 2022, 19, 8580.	2.6	0
2	The Genetic Variants in the Renin-Angiotensin System and the Risk of Heart Failure in Polish Patients. Genes, 2022, 13, 1257.	2.4	8
3	Post-match recovery profile of leukocyte cell subsets among professional soccer players. Scientific Reports, 2021, 11, 13352.	3.3	2
4	An association of ABCG8: rs11887534 polymorphism and HDL-cholesterol response to statin treatment in the Polish population. Pharmacological Reports, 2021, 73, 1781-1786.	3.3	1
5	Key genetic variants in the renin-angiotensin system and left ventricular mass in a cohort of Polish patients with heart failure. Kardiologia Polska, 2021, 79, 765-772.	0.6	1
6	Genetic Polymorphisms ofMMP1,MMP9,COL1A1, andCOL1A2in Polish Patients with Thoracic Aortopathy. Disease Markers, 2020, 2020, 1-8.	1.3	2
7	Damage-Associated Molecular Patterns and Th-Cell-Related Cytokines Released after Progressive Effort. Journal of Clinical Medicine, 2020, 9, 876.	2.4	11
8	Replication study of four keloid-associated polymorphisms in patients of European descent â€“ a single centre study. Intractable and Rare Diseases Research, 2020, 9, 40-42.	0.9	0
9	Molecular epidemiology of SARS CoV-2: a review of current data on genetic variability of the virus. Polish Archives of Internal Medicine, 2020, 131, 63-69.	0.4	14
10	Polymorphism of Interleukin 1B May Modulate the Risk of Ischemic Stroke in Polish Patients. Medicina (Lithuania), 2019, 55, 558.	2.0	10
11	E-selectin gene haplotypes are associated with the risk of myocardial infarction. Archives of Medical Science, 2019, 15, 1223-1231.	0.9	1
12	Two Functional TP53 Genetic Variants and Predisposition to Keloid Scarring in Caucasians. Dermatology Research and Practice, 2019, 2019, 1-5.	0.8	1
13	Non-replication of an association between the Odd-skipped related 1 c.654G>A (rs12329305) polymorphism and kidney volume in newborns. Pomeranian Journal of Life Sciences, 2019, 65, 14-18.	0.1	0
14	A non-synonymous lactotransferrin gene polymorphism and dental caries in 12-year-old children from the West Pomeranian region in Poland. Pomeranian Journal of Life Sciences, 2019, 64, 5-8.	0.1	0
15	Joint Assessment of Donor and Recipient hTERT Gene Polymorphism Provides Additional Information for Early Kidney Transplantation Outcomes. Medical Science Monitor, 2017, 23, 1812-1818.	1.1	5
16	Prevalence of 1691G>A FV mutation in Poland compared with that in other Central, Eastern and South-Eastern European countries. Bosnian Journal of Basic Medical Sciences, 2017, 12, 50.	1.0	8
17	Relationship between toll-like receptor 2 R753Q and T16934A polymorphisms and Staphylococcus aureus nasal carriage. Anaesthesiology Intensive Therapy, 2017, 49, 110-115.	1.0	3
18	BICD1 and Chromosome 18 Polymorphisms Associated With Recipients' Telomere Length Affect Kidney Allograft Function After Transplantation. Transplantation Proceedings, 2016, 48, 1451-1455.	0.6	3

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19	Genetic Identification of Communist Crimesâ€™ Victims (1944â€“1956) Based on the Analysis of One of Many Mass Graves Discovered on the Powazki Military Cemetery in Warsaw, Poland. Journal of Forensic Sciences, 2016, 61, 1450-1455.	1.6	14
20	Association Between RET (rs1800860) and GFRA1 (rs45568534, rs8192663, rs181595401,) Tj ETQq0 0 0 rgBT /Overlock Molecular Biomarkers, 2016, 20, 624-628.	0.7	4
21	Inflammation markers are associated with metabolic syndrome and ventricular arrhythmia in patients with coronary artery disease. Postepy Higieny I Medycyny Doswiadczalnej, 2016, 70, 56-66.	0.1	19
22	Trinucleotide repeat length in the first exon of the androgen receptor gene may be associated with prostate carcinogenesis and facilitate prediction of prostate cancer aggressiveness. Polish Archives of Internal Medicine, 2016, 126, 201-203.	0.4	0
23	Kidney Allograft Telomere Length Is Not Associated with Sex, Recipient Comorbid Conditions, Post-Transplant Infections, or CMV Reactivation. Annals of Transplantation, 2016, 21, 392-399.	0.9	1
24	Missense splice variant (g.20746A>G, p.Ile183Val) of interferon gamma receptor 1 (IFNGR1) coincidental with mycobacterial osteomyelitis - a screen of osteoarticular lesions. Bosnian Journal of Basic Medical Sciences, 2016, 16, 215-221.	1.0	3
25	The impact of the d3-growth hormone receptor (d3-GHR) polymorphism on the therapeutic effect of growth hormone replacement in children with idiopathic growth hormone deficiency in Poland. Neuroendocrinology Letters, 2016, 37, 282-288.	0.2	2
26	Association of functional genetic variants of A-kinase anchoring protein 10 with QT interval length in full-term Polish newborns. Archives of Medical Science, 2015, 1, 149-154.	0.9	5
27	Effect of delayed graft function, acute rejection and chronic allograft dysfunction on kidney allograft telomere length in patients after transplantation: a prospective cohort study. BMC Nephrology, 2015, 16, 23.	1.8	22
28	Epistatic interaction between common AGT G(â˜’ 6)A (rs5051) and AGTR1 A1166C (rs5186) variants contributes to variation in kidney size at birth. Gene, 2015, 572, 72-78.	2.2	3
29	<i>hTERT, BICD1</i> and Chromosome 18 Polymorphisms Associated with Telomere Length Affect Kidney Allograft Function After Transplantation. Kidney and Blood Pressure Research, 2015, 40, 111-120.	2.0	9
30	Prevalence and Clinical Outcome of CYP21A2 Gene Mutations in Patients with Nonfunctional Adrenal Incidentalomas. Hormone and Metabolic Research, 2015, 47, 662-667.	1.5	8
31	Coronary artery disease. A study of three polymorphic sites of adenosine deaminase gene. Acta Cardiologica, 2014, 69, 39-44.	0.9	7
32	Retrospective mutational analysis of NPHS1, NPHS2, WT1 and LAMB2 in children with steroid-resistant focal segmental glomerulosclerosis â€“ a single-centre experience. Bosnian Journal of Basic Medical Sciences, 2014, 14, 89.	1.0	15
33	New insights into the diagnosis of nodular goiter. Thyroid Research, 2014, 7, 6.	1.5	11
34	PTPN22 1858C>T gene polymorphism in patients with SLE: association with serological and clinical results. Molecular Biology Reports, 2014, 41, 6195-6200.	2.3	18
35	A functional IL-6 receptor (IL6R) variant is a risk factor for persistent atopic dermatitis. Journal of Allergy and Clinical Immunology, 2013, 132, 371-377.	2.9	86
36	Association of Glucocorticoid Receptor Gene NR3C1 Genetic Variants withÂAngiographically Documented Coronary Artery Disease and Its Risk Factors. Archives of Medical Research, 2013, 44, 27-33.	3.3	8

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37	Allele frequency distribution of 1691G>A F5 (which confers Factor V Leiden) across Europe, including Slavic populations. Journal of Applied Genetics, 2013, 54, 441-446.	1.9	16
38	Thymidylate Synthase Gene Polymorphism and Survival of Colorectal Cancer Patients Receiving Adjuvant 5-Fluorouracil. Genetic Testing and Molecular Biomarkers, 2013, 17, 799-806.	0.7	12
39	Association of BMPR1A polymorphism, but not BMP4, with kidney size in full-term newborns. Pediatric Nephrology, 2013, 28, 433-438.	1.7	7
40	Genetics of the renin-angiotensin system with respect to cardiac and blood pressure phenotypes in healthy newborn infants. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2013, 14, 337-347.	1.7	9
41	Skew-t Fits to Mortality Dataâ€"Can a Gaussian-Related Distribution Replace the Gompertzâ€"Makeham as the Basis for Mortality Studies?. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2013, 68, 903-913.	3.6	2
42	An insertion/deletion ACE polymorphism and kidney size in Polish full-term newborns. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2013, 14, 369-374.	1.7	3
43	Polymorphism 1936A>G in the AKAP10 gene (encoding A-kinase-anchoring protein 10) is associated with higher cholesterol cord blood concentration in Polish full-term newborns. Journal of Perinatal Medicine, 2013, 41, 205-10.	1.4	2
44	Estimation of BDNF gene polymorphism and predisposition to dependence development for selected psychoactive compounds. Human and Experimental Toxicology, 2013, 32, 236-240.	2.2	7
45	Association of Genetic Variation in Calmodulin and Left Ventricular Mass in Full-Term Newborns. International Journal of Genomics, 2013, 2013, 1-7.	1.6	1
46	Association of 1936A > G inAKAP10(A-kinase anchoring protein 10) and blood pressure in Polish full-term newborns. Blood Pressure, 2013, 22, 51-56.	1.5	4
47	The association of adenosine deaminase with coronary artery disease: Effect of gender and diabetes. Health, 2013, 05, 166-169.	0.3	3
48	Î²2-Adrenergic Receptor Gene Polymorphism and Response to Bronchodilating Treatment Evaluated by Spirometry. Advances in Experimental Medicine and Biology, 2013, 755, 169-177.	1.6	0
49	Impact of kidney donor hemostasis on risk of complications after transplantation â€“ preliminary outcomes. Medical Science Monitor, 2013, 19, 1102-1108.	1.1	3
50	The E23K polymorphism of the KCNJ11gene is associated with lower insulin release in patients with autosomal dominant polycystic kidney disease. Nefrologia, 2013, 33, 855-8.	0.4	1
51	Common Genetic Variants of the BMP4, BMPR1A, BMPR1B, and ACVR1 Genes, Left Ventricular Mass, and Other Parameters of the Heart in Newborns. Genetic Testing and Molecular Biomarkers, 2012, 16, 1309-1316.	0.7	9
52	Possible counter effect in newborns of 1936A>G (I646V) polymorphism in the AKAP10 gene encoding A-kinase-anchoring protein 10. Journal of Perinatology, 2012, 32, 230-234.	2.0	5
53	Are the Oxidoreductases (Endothelial NO Synthase, Catalase, Glutathione Peroxidase, Superoxide) Tj ETQq1 1 0.784314 rgBT /Overlo Transplantation, 2012, 94, 1047.	1.0	0
54	CYP17 and CYP19 genetic variants are not associated with age at natural menopause in Polish women. Reproductive Biology, 2012, 12, 368-373.	1.9	4

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55	1936Aâ†'G (I646ÂV) Polymorphism in theAKAP10Gene Encoding A-Kinase-Anchoring Protein 10 in Very Long-Lived Poles is Similar to that in Newborns. Experimental Aging Research, 2012, 38, 584-592.	1.2	2
56	Analysis for genotyping Duffy blood group in inhabitants of Sudan, the Fourth Cataract of the Nile. Malaria Journal, 2012, 11, 115.	2.3	22
57	Frequencies of functional caspase 12 genotypes in the North-Africa population. Russian Journal of Genetics, 2012, 48, 477-479.	0.6	1
58	Association of C49620T ABCC8 polymorphism with anthropometric and metabolic parameters in patients with autosomal dominant polycystic kidney disease: a preliminary study. Nefrologia, 2012, 32, 153-9.	0.4	3
59	The Effect of Cause of Cadaveric Kidney Donors Death on Fibrinolysis and Blood Coagulation Processes. Transplantation Proceedings, 2011, 43, 2866-2870.	0.6	3
60	Association of rs10918594 Polymorphisms of Nitric Oxide Synthase 1 Adaptor Protein (NOS1AP) with QTc Interval Prolongation During Kidney Transplantation. Transplantation Proceedings, 2011, 43, 2964-2966.	0.6	1
61	Graft Infection in Kidney Recipients and Its Relation to Transplanted Kidney Function. Transplantation Proceedings, 2011, 43, 2997-2999.	0.6	3
62	The Outcomes of Treatment and the Etiology of Lymphoceles With a Focus on Hemostasis in Kidney Recipients: A Preliminary Report. Transplantation Proceedings, 2011, 43, 3008-3012.	0.6	11
63	Donor-Recipient Gender Mismatch Affects Early Graft Loss After Kidney Transplantation. Transplantation Proceedings, 2011, 43, 2914-2916.	0.6	20
64	VAMP-8 gene variant is associated with increased risk of early myocardial infarction. Archives of Medical Science, 2011, 3, 440-443.	0.9	4
65	Risk of All-Cause Mortality in HIV Infected Patients Is Associated with Clinical, Immunologic Predictors and the CCR5 Î"32 Deletion. PLoS ONE, 2011, 6, e22215.	2.5	8
66	Lack of Association of Interleukin-1 Gene Cluster Polymorphisms with Angiographically Documented Coronary Artery Disease: Demonstration of Association with Hypertension in the Polish Population. Archives of Medical Research, 2011, 42, 426-32.	3.3	8
67	AMPD1 gene mutations are associated with obesity and diabetes in Polish patients with cardiovascular diseases. Journal of Applied Genetics, 2011, 52, 67-76.	1.9	16
68	Association Screening in the Epidermal Differentiation Complex (EDC) Identifies an SPRR3 Repeat Number Variant as a Risk Factor for Eczema. Journal of Investigative Dermatology, 2011, 131, 1644-1649.	0.7	72
69	The distribution of human endogenous retrovirus K-113 in health and autoimmune diseases in Poland. Rheumatology, 2011, 50, 1310-1314.	1.9	24
70	Prevalence of 845G>A HFE mutation in Slavic populations: an east-west linear gradient in South Slavs. Croatian Medical Journal, 2011, 52, 351-357.	0.7	4
71	Factors influencing QTc interval prolongation during kidney transplantation. Annals of Transplantation, 2011, 16, 43-49.	0.9	3
72	Association of the rs10918594 of Nitric Oxide Synthase 1 Adaptor Protein (NOS1AP) polymorphisms with the graft function after kidney transplantation. Annals of Transplantation, 2011, 16, 72-76.	0.9	1

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73	Low frequency haplotypes of E-selectin polymorphisms G2692A and C1901T give increased protection from coronary artery disease. Medical Science Monitor, 2011, 17, CR334-CR340.	1.1	13
74	The association of â€“262C/T polymorphism in the catalase gene and delayed graft function of kidney allografts. Nephrology, 2010, 15, 587-591.	1.6	4
75	HISTOMORPHOLOGICAL AND IMMUNOHISTOCHEMICAL EVALUATION OF PERFUSED AND PRESERVED KIDNEY WITH EC OR UW SOLUTION STUDIED WITH AN EXPERIMENTAL MODEL OF THE PRE-TRANSPLANTATION PERIOD. Transplantation, 2010, 90, 887.	1.0	0
76	353 The Vntr Il1Rn Gene Polymorphism in Polish Children with Diffuse Mesangial Proliferation (DMP). Pediatric Research, 2010, 68, 182-183.	2.3	0
77	An age-related decrease in factor V Leiden frequency among Polish subjects. Journal of Applied Genetics, 2010, 51, 337-341.	1.9	3
78	Polymorphisms of Superoxide Dismutase, Glutathione Peroxidase andÂCatalase Genes in Patients with Post-transplant Diabetes Mellitus. Archives of Medical Research, 2010, 41, 350-355.	3.3	23
79	Functional polymorphism of matrix metalloproteinase-9 (MMP-9) gene in alcohol dependence: Family and case control study. Brain Research, 2010, 1327, 103-106.	2.2	51
80	The association between eNOS intron 4 VNTR polymorphism and delayed graft function of kidney allografts. Clinical Transplantation, 2010, 24, E130-6.	1.6	4
81	Histopathologic Evaluation of Pretransplantation Biopsy as a Factor Influencing Graft Function After Kidney Transplantation in 3-Year Observation. Transplantation Proceedings, 2010, 42, 3375-3381.	0.6	22
82	Family-based study of brain-derived neurotrophic factor (Î’DNF) gene polymorphism in alcohol dependence. Pharmacological Reports, 2010, 62, 938-941.	3.3	33
83	G(â€“2548)A leptin gene polymorphism in obese subjects is associated with serum leptin concentration and bone mass. Polish Archives of Internal Medicine, 2010, 120, 175-180.	0.4	8
84	Association of C34TAMPD1gene polymorphism with features of metabolic syndrome in patients with coronary artery disease or heart failure. Scandinavian Journal of Clinical and Laboratory Investigation, 2009, 69, 102-112.	1.2	28
85	Metabolic Syndrome Components in Patients with Autosomal-Dominant Polycystic Kidney Disease. Kidney and Blood Pressure Research, 2009, 32, 405-410.	2.0	20
86	C677T polymorphism of the methylenetetrahydrofolate reductase gene and the risk of ischemic stroke in Polish subjects. Journal of Applied Genetics, 2009, 50, 63-67.	1.9	24
87	Sequence variants of chemokine receptor genes and susceptibility to HIV-1 infection. Journal of Applied Genetics, 2009, 50, 159-166.	1.9	16
88	A common variant on chromosome 11q13 is associated with atopic dermatitis. Nature Genetics, 2009, 41, 596-601.	21.4	297
89	Plasma concentrations of TNFâ€Î± and its soluble receptors sTNFR1 and sTNFR2 in patients with coronary artery disease. Tissue Antigens, 2009, 74, 386-392.	1.0	60
90	Lack of Association of Polymorphisms 239+34A/C in the SOD1 Gene and 47C/T in the SOD2 Gene With Delayed Graft Function and Acute and Chronic Rejection of Kidney Allografts. Transplantation Proceedings, 2009, 41, 3701-3703.	0.6	5

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91	Association of the A1936G (rs203462) of A-Kinase Anchoring Protein 10 Polymorphisms With QT Interval Prolongation During Kidney Transplantation. Transplantation Proceedings, 2009, 41, 3036-3038.	0.6	3
92	N7 Trinucleotide repeat length polymorphisms of the androgen receptor gene â€“ a step forward in understanding the pathogenesis of prostate cancer. European Urology Supplements, 2009, 8, 569.	0.1	0
93	Frequency of common CYP3A5 gene variants in healthy Polish newborn infants. Pharmacological Reports, 2009, 61, 947-951.	3.3	18
94	Is dyslipidemia sustained during remission of nephrotic syndrome genetically determined? Evaluation of genetic polymorphisms of proteins involved in lipoprotein metabolism in children and adolescents with nephrotic syndrome. Polish Archives of Internal Medicine, 2009, 119, 11-17.	0.4	4
95	â€œTreasure your exceptionsâ€• recent advances in molecular genetics of glomerular disease. Journal of Applied Genetics, 2008, 49, 93-99.	1.9	2
96	Expression of E-SOD, GPX5 mRNAs and immunoexpression of Cu/ZnSOD in epididymal epithelial cells of finasteride-treated rats. Andrologia, 2008, 40, 303-311.	2.1	11
97	Effect of Trimetazidine on Xanthine Oxidoreductase Expression in Rat Kidney with Ischemiaâ˜'Reperfusion Injury. Archives of Medical Research, 2008, 39, 459-462.	3.3	14
98	The Common C49620T Polymorphism in the Sulfonylurea Receptor Gene (ABCC8), Pancreatic Beta Cell Function and Long-Term Diabetic Complications in Obese Patients with Long-Lasting Type 2 Diabetes Mellitus. Experimental and Clinical Endocrinology and Diabetes, 2007, 115, 317-321.	1.2	12
99	ADA*2 Allele of the Adenosine Deaminase Gene May Protect against Coronary Artery Disease. Cardiology, 2007, 108, 275-281.	1.4	31
100	Inhibition of phospholipase A2 activity by conjugated linoleic acids in human macrophages. European Journal of Nutrition, 2007, 46, 28-33.	3.9	17
101	659 Angiotensin-converting enzyme 2 (ACE2) gene polymorphisms and long-term survival in advanced heart failure. European Journal of Heart Failure, Supplement, 2007, 6, 142-142.	0.0	0
102	Association between the Pro12Ala variant of the peroxisome proliferator-activated receptor-gamma2 gene and increased 24-h diastolic blood pressure in obese patients with type II diabetes. Journal of Human Hypertension, 2006, 20, 684-692.	2.2	28
103	Optimized RT-PCR Method for Assaying Expression of Monocyte Chemotactic Protein Type 1 (MCP-1) in Rabbit Aorta. Biochemical Genetics, 2006, 44, 129-139.	1.7	10
104	DNA microsatellite analysis in families with autosomal dominant polycystic kidney disease (ADPKD): the first Polish study. Journal of Applied Genetics, 2006, 47, 383-389.	1.9	5
105	Lack of Association between the Pro12Ala Polymorphism in PPAR-Î³2 Gene and Body Weight Changes, Insulin Resistance and Chronic Diabetic Complications in Obese Patients with Type 2 Diabetes. Archives of Medical Research, 2006, 37, 736-743.	3.3	29
106	Plasma Prekallikrein as a Risk Factor for Diabetic Retinopathy. Archives of Medical Research, 2005, 36, 539-543.	3.3	14
107	Liddle syndrome caused by P616R mutation of the epithelial sodium channel Î² subunit. Pediatric Nephrology, 2005, 20, 837-838.	1.7	16
108	The in situ expression of interleukin-8 in the normal human kidney and in different morphological forms of glomerulonephritis. American Journal of Kidney Diseases, 2004, 43, 983-998.	1.9	31

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109	Effects of perindopril treatment on hemostatic function in patients with essential hypertension in relation to angiotensin converting enzyme (ACE) and plasminogen activator inhibitor-1 (PAI-1) gene polymorphisms. Nutrition, Metabolism and Cardiovascular Diseases, 2004, 14, 259-269.	2.6	12
110	THE I/D ACE GENOTYPE MAY PREDICT THE DIASTOLIC BLOOD PRESSURE RESPONSE TO LOSARTAN IN PATIENTS WITH ESSENTIAL HYPERTENSION. Journal of Hypertension, 2004, 22, S258.	0.5	0
111	The expression of telomerase componens in human gastric cancer. Gastroenterology, 2003, 124, A553.	1.3	0
112	EcoRI polymorphism of the L-myc gene in gastric cancer patients. European Journal of Gastroenterology and Hepatology, 2002, 14, 1231-1235.	1.6	8
113	Selective inhibition by probucol of vascular cell adhesion molecule-1 (VCAM-1) expression in human vascular endothelial cells. Atherosclerosis, 2001, 155, 123-130.	0.8	55
114	Lack of Association between Gly460Trp Polymorphism of Alpha-Adducin Gene and Salt Sensitivity of Blood Pressure in Polish Hypertensives. Kidney and Blood Pressure Research, 2001, 24, 201-206.	2.0	24
115	Selective Inhibition by Î±-Tocopherol of Vascular Cell Adhesion Molecule-1 Expression in Human Vascular Endothelial Cells. Biochemical and Biophysical Research Communications, 2000, 274, 609-615.	2.1	52
116	The detection of somatic mutations of thyrotropin receptor gene in fine needle biopsy samples from thyroid nodules. Endocrine Regulations, 1999, 33, 95-101.	1.3	3
117	Leptin receptor isoforms expressed in human adipose tissue. Metabolism: Clinical and Experimental, 1998, 47, 844-847.	3.4	110
118	F069 The influence of probucol on PDGF-A and PDGF-B gene expression in peropheral blood monocytes (PBM) in patients after myocardial infraction. Atherosclerosis, 1998, 136, S64.	0.8	0
119	Inducible nitric oxide synthase in the epithelial epididymal cells of the rat. Reproduction, Fertility and Development, 1997, 9, 789.	0.4	15
120	Extrahepatic Transcription of Plasma Prekallikrein Gene in Human and Rat Tissues. Biochemical and Biophysical Research Communications, 1993, 197, 1370-1376.	2.1	17
121	Plasma Prekallikrein Levels in Patients with Hepatocellular Carcinoma and Liver Cirrhosis: A Pilot Study. Annals of Clinical Biochemistry, 1993, 30, 445-448.	1.6	1
122	Coronary artery disease. A study of three polymorphic sites of adenosine deaminase gene. , 0, .		2
123	ADPKD protects against diabetogenic effects associated with genetically-predicted lactase persistence.. Archives of Medical Science, 0, , .	0.9	0