Jaroslav A Hubacek

List of Publications by Year in descending order

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252 papers

9,581 citations

76326 40 h-index 90 g-index

253 all docs

253 docs citations

times ranked

253

14034 citing authors

#	Article	IF	CITATIONS
1	The interleukin-6 receptor as a target for prevention of coronary heart disease: a mendelian randomisation analysis. Lancet, The, 2012, 379, 1214-1224.	13.7	886
2	An Apolipoprotein Influencing Triglycerides in Humans and Mice Revealed by Comparative Sequencing. Science, 2001, 294, 169-173.	12.6	842
3	Triglyceride-mediated pathways and coronary disease: collaborative analysis of 101 studies. Lancet, The, 2010, 375, 1634-1639.	13.7	606
4	HMG-coenzyme A reductase inhibition, type 2 diabetes, and bodyweight: evidence from genetic analysis and randomised trials. Lancet, The, 2015, 385, 351-361.	13.7	562
5	Association between alcohol and cardiovascular disease: Mendelian randomisation analysis based on individual participant data. BMJ, The, 2014, 349, g4164-g4164.	6.0	528
6	Two independent apolipoprotein A5 haplotypes influence human plasma triglyceride levels. Human Molecular Genetics, 2002, 11, 3031-3038.	2.9	352
7	C(â^'260)â†'T Polymorphism in the Promoter of the CD14 Monocyte Receptor Gene as a Risk Factor for Myocardial Infarction. Circulation, 1999, 99, 3218-3220.	1.6	314
8	Genetic association study of QT interval highlights role for calcium signaling pathways in myocardial repolarization. Nature Genetics, 2014, 46, 826-836.	21.4	281
9	Effect modification by population dietary folate on the association between MTHFR genotype, homocysteine, and stroke risk: a meta-analysis of genetic studies and randomised trials. Lancet, The, 2011, 378, 584-594.	13.7	273
10	Gene variants of the bactericidal/permeability increasing protein and lipopolysaccharide binding protein in sepsis patients. Critical Care Medicine, 2001, 29, 557-561.	0.9	194
11	Mutations in ATP-cassette binding proteins G5 (ABCG5) and G8 (ABCG8) causing sitosterolemia. Human Mutation, 2001, 18, 359-360.	2.5	149
12	Apolipoprotein E genotype, cardiovascular biomarkers and risk of stroke: Systematic review and meta-analysis of 14 015 stroke cases and pooled analysis of primary biomarker data from up to 60 883 individuals. International Journal of Epidemiology, 2013, 42, 475-492.	1.9	145
13	Secretory Phospholipase A2-IIA and Cardiovascular Disease. Journal of the American College of Cardiology, 2013, 62, 1966-1976.	2.8	115
14	Dietary polyphenols are inversely associated with metabolic syndrome in Polish adults of the HAPIEE study. European Journal of Nutrition, 2017, 56, 1409-1420.	3.9	111
15	Czech mass methanol outbreak 2012: Epidemiology, challenges and clinical features. Clinical Toxicology, 2014, 52, 1013-1024.	1.9	108
16	Essential Role of AT _{1A} Receptor in the Development of 2K1C Hypertension. Hypertension, 2002, 40, 735-741.	2.7	101
17	The influence of <i>APOAV</i> polymorphisms (Tâ€1131>C and S19>W) on plasma triglyceride levels and risk of myocardial infarction. Clinical Genetics, 2004, 65, 126-130.	2.0	96
18	The common functional $C(\hat{a}^*.159)$ T polymorphism within the promoter region of the lipopolysaccharide receptor CD14 is not associated with sepsis development or mortality. Genes and Immunity, 2000, 1, 405-407.	4.1	88

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19	Mediterranean diet score and total and cardiovascular mortality in Eastern Europe: the HAPIEE study. European Journal of Nutrition, 2017, 56, 421-429.	4.6	87
20	Long-term visual damage after acute methanol poisonings: Longitudinal cross-sectional study in 50 patients. Clinical Toxicology, 2015, 53, 884-892.	1.9	78
21	Education and wealth inequalities in healthy ageing in eight harmonised cohorts in the ATHLOS consortium: a population-based study. Lancet Public Health, The, 2020, 5, e386-e394.	10.0	77
22	Interleukin-6 gene variants and the risk of sepsis development in children. Human Immunology, 2007, 68, 756-760.	2.4	71
23	Association of serum bilirubin and promoter variations in <i>HMOX1</i> and <i>UGT1A1</i> genes with sporadic colorectal cancer. International Journal of Cancer, 2012, 131, 1549-1555.	5.1	70
24	Intermittent hemodialysis is superior to continuous veno-venous hemodialysis/hemodiafiltration to eliminate methanol and formate during treatment for methanol poisoning. Kidney International, 2014, 86, 199-207.	5.2	70
25	The Genomic Organization of the Genes for Human Lipopolysaccharide Binding Protein (LBP) and Bactericidal Permeability Increasing Protein (BPI) Is Highly Conserved. Biochemical and Biophysical Research Communications, 1997, 236, 427-430.	2.1	69
26	APOE polymorphism and its effect on plasma C-reactive protein levels in a large general population sample. Human Immunology, 2010, 71, 304-308.	2.4	63
27	Meta-Analysis of the INSIG2 Association with Obesity Including 74,345 Individuals: Does Heterogeneity of Estimates Relate to Study Design?. PLoS Genetics, 2009, 5, e1000694.	3.5	62
28	Dietary polyphenol intake and risk of type 2 diabetes in the Polish arm of the Health, Alcohol and Psychosocial factors in Eastern Europe (HAPIEE) study. British Journal of Nutrition, 2017, 118, 60-68.	2.3	62
29	The <i>FTO</i> Gene and Obesity in a Large Eastern European Population Sample: The HAPIEE Study. Obesity, 2008, 16, 2764-2766.	3.0	61
30	Effects of Six <i>APOA5</i> Variants, Identified in Patients With Severe Hypertriglyceridemia, on In Vitro Lipoprotein Lipase Activity and Receptor Binding. Arteriosclerosis, Thrombosis, and Vascular Biology, 2008, 28, 1866-1871.	2.4	59
31	Apolipoprotein A5 and triglyceridemia. Focus on the effects of the common variants. Clinical Chemistry and Laboratory Medicine, 2005, 43, 897-902.	2.3	55
32	A Possible Role of Apolipoprotein E Polymorphism in Predisposition to Higher Education. Neuropsychobiology, 2001, 43, 200-203.	1.9	54
33	Genetics of Familial Hypercholesterolemia: New Insights. Frontiers in Genetics, 2020, 11, 574474.	2.3	53
34	A FTO variant and risk of acute coronary syndrome. Clinica Chimica Acta, 2010, 411, 1069-1072.	1,1	49
35	Fruit and vegetable consumption and mortality in Eastern Europe: Longitudinal results from the Health, Alcohol and Psychosocial Factors in Eastern Europe study. European Journal of Preventive Cardiology, 2016, 23, 493-501.	1.8	49
36	T-1131â†'C polymorphism within the apolipoprotein AV gene in hypertriglyceridemic individuals. Atherosclerosis, 2003, 167, 369-370.	0.8	48

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37	Role of Cholesterol 7α-Hydroxylase (CYP7A1) in Nutrigenetics and Pharmacogenetics of Cholesterol Lowering. Molecular Diagnosis and Therapy, 2006, 10, 93-100.	3.8	48
38	FTO Variant, Energy Intake, Physical Activity and Basal Metabolic Rate in Caucasians. The HAPIEE Study. Physiological Research, 2011, 60, 175-183.	0.9	45
39	Dietary polyphenol intake and risk of hypertension in the Polish arm of the HAPIEE study. European Journal of Nutrition, 2018, 57, 1535-1544.	3.9	41
40	Polymorphisms in CYP-7A1, not APOE, influence the change in plasma lipids in response to population dietary change in an 8 year follow-up; results from the Czech MONICA study. Clinical Biochemistry, 2003, 36, 263-267.	1.9	40
41	The effect of rapid privatisation on mortality in mono-industrial towns in post-Soviet Russia: a retrospective cohort study. Lancet Public Health, The, 2017, 2, e231-e238.	10.0	40
42	The FTO gene polymorphism is associated with end-stage renal disease: two large independent case-control studies in a general population. Nephrology Dialysis Transplantation, 2012, 27, 1030-1035.	0.7	39
43	FTO and MC4R gene variants determine BMI changes in children after intensive lifestyle intervention. Clinical Biochemistry, 2013, 46, 313-316.	1.9	39
44	Bactericidal permeability increasing protein gene variants in children with sepsis. Intensive Care Medicine, 2007, 33, 2158-2164.	8.2	38
45	A common variant in the FTO gene is associated with body mass index in males and postmenopausal females but not in premenopausal females. Czech post-MONICA and 3PMFs studies. Clinical Chemistry and Laboratory Medicine, 2009, 47, 387-90.	2.3	38
46	Impact of apolipoprotein A5 variants on statin treatment efficacy. Pharmacogenomics, 2009, 10, 945-950.	1.3	38
47	Healthy diet indicator and mortality in Eastern European populations: prospective evidence from the HAPIEE cohort. European Journal of Clinical Nutrition, 2014, 68, 1346-1352.	2.9	38
48	A Mediterranean-type diet is associated with better metabolic profile in urban Polish adults: Results from the HAPIEE study. Metabolism: Clinical and Experimental, 2015, 64, 738-746.	3.4	38
49	Multiple gene-to-gene interactions in children with sepsis: a combination of five gene variants predicts outcome of life-threatening sepsis. Critical Care, 2014, 18, R1.	5.8	37
50	ACE I/D polymorphism in Czech first-wave SARS-CoV-2-positive survivors. Clinica Chimica Acta, 2021, 519, 206-209.	1.1	36
51	Statins and Inflammation. Current Atherosclerosis Reports, 2021, 23, 80.	4.8	34
52	CCR5î"32 Deletion as a Protective Factor in Czech First-Wave COVID-19 Subjects. Physiological Research, 2021, 70, 111-115.	0.9	33
53	Plasma levels of remnant particles are determined in part by variation in the APOC3 gene insulin response element and the APOCl–APOE cluster. Journal of Lipid Research, 2000, 41, 1103-1109.	4.2	33
54	Regulatory RNAs and Cardiovascular Disease – With a Special Focus on Circulating MicroRNAs. Physiological Research, 2017, 66, S21-S38.	0.9	33

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55	Association between heat shock protein 70s and Toll-like receptor polymorphisms with long-term renal allograft survival. Transplant International, 2006, 19, 190-196.	1.6	32
56	Antioxidant vitamin intake and mortality in three Central and Eastern European urban populations: the HAPIEE study. European Journal of Nutrition, 2016, 55, 547-560.	3.9	32
57	Apolipoprotein A5 fifteen years anniversary: Lessons from genetic epidemiology. Gene, 2016, 592, 193-199.	2.2	30
58	Homogeneous assay based on 52 primer sets to scan for mutations of the ABCA1 gene and its application in genetic analysis of a new patient with familial high-density lipoprotein deficiency syndrome. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2001, 1537, 42-48.	3.8	29
59	Depressive symptoms and levels of C-reactive protein. Social Psychiatry and Psychiatric Epidemiology, 2009, 44, 217-222.	3.1	29
60	MLXIPL variant in individuals with low and high triglyceridemia in white population in Central Europe. Human Genetics, 2008, 124, 553-555.	3.8	28
61	Impact of Selection Bias on Estimation of Subsequent Event Risk. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	28
62	Eat less and exercise more $\hat{a}\in$ is it really enough to knock down the obesity pandemia? Physiological Research, 2009, 58 Suppl 1, S1-S6.	0.9	28
63	Body Mass Index Change in Females After Short-Time Life Style Intervention Is Not Dependent on the FTO Polymorphisms. Physiological Research, 2011, 60, 199-202.	0.9	28
64	Alcohol consumption and cognitive performance: a <scp>M</scp> endelian randomization study. Addiction, 2014, 109, 1462-1471.	3.3	27
65	A systematic review and meta-analysis of 130,000 individuals shows smoking does not modify the association of APOE genotype on risk of coronary heart disease. Atherosclerosis, 2014, 237, 5-12.	0.8	27
66	DNA Methylation Profiles in a Group of Workers Occupationally Exposed to Nanoparticles. International Journal of Molecular Sciences, 2020, 21, 2420.	4.1	27
67	Does angiotensin-converting enzyme polymorphism influence the clinical manifestation and progression of heart failure in patients with dilated cardiomyopathy?. American Journal of Cardiology, 1999, 83, 461-462.	1.6	26
68	Polymorphisms in the Lipopolysaccharide-Binding Protein and Bactericidal/Permeability-Increasing Protein in Patients with Myocardial Infarction. Clinical Chemistry and Laboratory Medicine, 2002, 40, 1097-100.	2.3	25
69	Ser19->Trp polymorphism within the apolipoprotein AV gene in hypertriglyceridaemic people. Journal of Medical Genetics, 2003, 40, 105e-105.	3.2	25
70	Donor PNPLA3 rs738409 genotype is a risk factor for graft steatosis. A post-transplant biopsy-based study. Digestive and Liver Disease, 2018, 50, 490-495.	0.9	25
71	Genetics of Cardiovascular Disease: How Far Are We from Personalized CVD Risk Prediction and Management?. International Journal of Molecular Sciences, 2021, 22, 4182.	4.1	25
72	Association of Metabolic and Genetic Factors With Cholesterol Esterification Rate in HDL Plasma and Atherogenic Index of Plasma in a 40 Years Old Slovak Population. Physiological Research, 2011, 60, 785-795.	0.9	25

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73	SLCO1B1 Polymorphism is not associated with Risk of Statin-Induced Myalgia/Myopathy in a Czech Population. Medical Science Monitor, 2015, 21, 1454-1459.	1.1	24
74	Association between FTO 1st intron tagging variant and telomere length in middle aged females. 3PMFs study. Clinica Chimica Acta, 2012, 413, 1222-1225.	1.1	23
75	Association between plasma bilirubin and mortality. Annals of Hepatology, 2019, 18, 379-385.	1.5	23
76	Apolipoprotein A5 in health and disease. Physiological Research, 2009, 58 Suppl 2, S101-S110.	0.9	23
77	Imaging findings after methanol intoxication (cohort of 46 patients). Neuroendocrinology Letters, 2015, 36, 737-44.	0.2	23
78	Polymorphisms in the APOA1/C3/A4/A5 gene cluster and cholesterol responsiveness to dietary change. Clinical Chemistry and Laboratory Medicine, 2007, 45, 316-20.	2.3	22
79	Sex-specific interaction between APOE and APOA5 variants and determination of plasma lipid levels. European Journal of Human Genetics, 2008, 16, 135-138.	2.8	22
80	Association of Chromosome 9p21 With Subsequent Coronary Heart Disease Events. Circulation Genomic and Precision Medicine, 2019, 12, e002471.	3.6	22
81	Polygenic Hypercholesterolemia: Examples of GWAS Results and Their Replication in the Czech-Slavonic Population. Physiological Research, 2017, 66, S101-S111.	0.9	22
82	Rare Alleles within the <i><scp>CYP</scp>2E1</i> (<scp>MEOS</scp> System) Could be Associated with Better Shortâ€Term Health Outcome after Acute Methanol Poisoning. Basic and Clinical Pharmacology and Toxicology, 2015, 116, 168-172.	2.5	21
83	Cognitive sequelae of methanol poisoning involve executive dysfunction and memory impairment in cross-sectional and long-term perspective. Alcohol, 2017, 59, 27-35.	1.7	21
84	Coffee consumption and mortality in three Eastern European countries: results from the HAPIEE (Health, Alcohol and Psychosocial factors In Eastern Europe) study. Public Health Nutrition, 2017, 20, 82-91.	2.2	21
85	The clinical utility of structural neuroimaging in first-episode psychosis: A systematic review. Australian and New Zealand Journal of Psychiatry, 2019, 53, 1093-1104.	2.3	21
86	Trajectories of Accumulation of Health Deficits in Older Adults: Are There Variations According to Health Domains?. Journal of the American Medical Directors Association, 2019, 20, 710-717.e6.	2.5	21
87	Clinical and genetic determinants of chronic visual pathway changes after methanol - induced optic neuropathy: four-year follow-up study. Clinical Toxicology, 2019, 57, 387-397.	1.9	20
88	Genetic determination of plasma lipids and insulin in the Czech population. Clinical Biochemistry, 2001, 34, 113-118.	1.9	19
89	Effect of apolipoprotein E polymorphism on statin-induced decreases in plasma lipids and cardiovascular events. Drug Metabolism and Drug Interactions, 2011, 26, 13-20.	0.3	19
90	Alcohol, pattern of drinking and allâ€cause mortality in Russia, Belarus and Hungary: a retrospective indirect cohort study based on mortality of relatives. Addiction, 2018, 113, 1252-1263.	3.3	19

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91	The FTO variant is associated with chronic complications of diabetes mellitus in Czech population. Gene, 2018, 642, 220-224.	2.2	19
92	Traditional Eastern European diet and mortality: prospective evidence from the HAPIEE study. European Journal of Nutrition, 2021, 60, 1091-1100.	3.9	19
93	Apolipoprotein E4 Allele in Subjects with COVID-19. Gerontology, 2021, 67, 320-322.	2.8	19
94	TGF- \hat{i}^21 gene polymorphism is a risk factor for renal dysfunction in heart transplant recipients. Transplantation Proceedings, 2001, 33, 1567-1569.	0.6	18
95	Changes of Plasma Lipids during Weight Reduction in Females Depends on APOA5 Variants. Annals of Nutrition and Metabolism, 2008, 53, 104-108.	1.9	18
96	The gendered effects of foreign investment and prolonged state ownership on mortality in Hungary: an indirect demographic, retrospective cohort study. The Lancet Global Health, 2018, 6, e95-e102.	6.3	18
97	Leukocyte telomere length and risk of coronary heart disease and stroke mortality: prospective evidence from a Russian cohort. Scientific Reports, 2018, 8, 16627.	3.3	18
98	Rare variants in known and novel candidate genes predisposing to statin-associated myopathy. Pharmacogenomics, 2016, 17, 1405-1414.	1.3	17
99	Subsequent Event Risk in Individuals With Established Coronary Heart Disease. Circulation Genomic and Precision Medicine, 2019, 12, e002470.	3.6	17
100	Genetic variants associated with glycemic response to treatment with dipeptidylpeptidase 4 inhibitors. Pharmacogenomics, 2020, 21, 317-323.	1.3	17
101	Traditional Risk Factors of Acute Coronary Syndrome in Four Different Male Populations – Total Cholesterol Value Does Not Seem To Be Relevant Risk Factor. Physiological Research, 2017, 66, S121-S128.	0.9	17
102	Sex-specific effect of APOAV variant (Val153>Met) on plasma levels of high-density lipoprotein cholesterol. Metabolism: Clinical and Experimental, 2005, 54, 1632-1635.	3.4	16
103	Comparison of food and nutrient intakes between cohorts of the HAPIEE and Whitehall II studies. European Journal of Public Health, 2016, 26, 628-634.	0.3	16
104	Prevalence, awareness, treatment and control of dyslipidemia in older persons in urban and rural population in the Astana region, Kazakhstan. BMC Public Health, 2017, 17, 651.	2.9	16
105	Donor PNPLA3 and TM6SF2 Variant Alleles Confer Additive Risks for Graft Steatosis After Liver Transplantation. Transplantation, 2020, 104, 526-534.	1.0	16
106	ACE gene polymorphism and long-term renal graft function. Clinical Biochemistry, 2001, 34, 87-90.	1.9	15
107	Gene variants at FTO, 9p21, and 2q36.3 are age-independently associated with myocardial infarction in Czech men. Clinica Chimica Acta, 2016, 454, 119-123.	1.1	15
108	Association of methylenetetrahydrofolate reductase T677 allele with early development of chronic allograft nephropathy. Clinical Biochemistry, 2004, 37, 919-924.	1.9	14

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109	Hypertriglyceridemia: Interaction between APOE and APOAV Variants. Clinical Chemistry, 2005, 51, 1311-1313.	3.2	14
110	APOA5 variant Ser19Trp influences a decrease of the total cholesterol in a male 8 year cohort. Clinical Biochemistry, 2006, 39, 133-136.	1.9	14
111	Predictors of infection after pulmonary endarterectomy for chronic thrombo-embolic pulmonary hypertension. European Journal of Cardio-thoracic Surgery, 2011, 39, 195-200.	1.4	14
112	Does the consumption of fruits and vegetables differ between Eastern and Western European populations? Systematic review of cross-national studies. Archives of Public Health, 2015, 73, 29.	2.4	14
113	Mortality in Transition: Study Protocol of the PrivMort Project, a multilevel convenience cohort study. BMC Public Health, 2016, 16, 672.	2.9	14
114	Smoking and Mortality in Eastern Europe: Results From the PrivMort Retrospective Cohort Study of 177 376 Individuals. Nicotine and Tobacco Research, 2018, 20, 749-754.	2.6	14
115	Der (TTA)n Repeat-Polymorphismus im HMG-CoA Reduktase-Gen und der Serumcholesterinspiegel. Vasa - European Journal of Vascular Medicine, 1999, 28, 169-171.	1.4	13
116	Neuroinflammation markers and methyl alcohol induced toxic brain damage. Toxicology Letters, 2018, 298, 60-69.	0.8	13
117	Socioeconomic inequalities in physical and cognitive functioning: cross-sectional evidence from 37 cohorts across 28 countries in the ATHLOS project. Journal of Epidemiology and Community Health, 2021, 75, 980-986.	3.7	13
118	High-sensitivity C-reactive protein concentration in patients with myocardial infarction-environmental factors, and polymorphisms in interleukin-10 and CD14 genes. Clinical Chemistry and Laboratory Medicine, 2007, 45, 855-61.	2.3	12
119	APOA5 Ala315>Val, identified in patients with severe hypertriglyceridemia, is a common mutation with no major effects on plasma lipid levels. Clinical Chemistry and Laboratory Medicine, 2008, 46, 773-7.	2.3	12
120	Drug metabolising enzyme polymorphisms in Middle- and Eastern-European Slavic populations. Drug Metabolism and Drug Interactions, 2014, 29, 29-36.	0.3	12
121	Association of MTHFR genetic variants C677T and A1298C on predisposition to spontaneous abortion in Slavonic population. Clinica Chimica Acta, 2015, 440, 104-107.	1.1	12
122	Fruit, vegetable intake and blood pressure trajectories in older age. Journal of Human Hypertension, 2019, 33, 671-678.	2.2	12
123	Education and mortality in three Eastern European populations: findings from the PrivMort retrospective cohort study. European Journal of Public Health, 2019, 29, 549-554.	0.3	12
124	The impact of co-morbidities on a 6-year survival after methanol mass poisoning outbreak: possible role of metabolic formaldehyde. Clinical Toxicology, 2020, 58, 241-253.	1.9	12
125	The connexin 37 (1019C>T) gene polymorphism is associated with subclinical atherosclerosis in women with type 1 and 2 diabetes and in women with central obesity. Physiological Research, 2010, 59, 1029-1032.	0.9	12
126	Intimaâ€media Thickness of Carotid Arteries in Borderline Hypertensives. Journal of Neuroimaging, 1999, 9, 19-22.	2.0	11

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127	Angiotensin converting enzyme gene â€" a candidate gene for addiction to smoking?. Atherosclerosis, 2001, 159, 237-238.	0.8	11
128	Genetic analysis of APOAV polymorphisms (T-1131/C, Ser19/Trp and Val153/Met): no effect on plasma remnant particles concentrations. Clinica Chimica Acta, 2004, 348, 171-175.	1.1	11
129	Apolipoprotein E Polymorphism in Hemodialyzed Patients and Healthy Controls. Biochemical Genetics, 2009, 47, 688-693.	1.7	11
130	Association between polymorphism within the RYR2 receptor and development of statin-associated myalgia/myopathy in the Czech population. European Journal of Internal Medicine, 2015, 26, 367-368.	2,2	11
131	Genetic variants within telomere-associated genes, leukocyte telomere length and the risk of acute coronary syndrome in Czech women. Clinica Chimica Acta, 2016, 454, 62-65.	1.1	11
132	Analysis of circulating miRNAs in patients with familial hypercholesterolaemia treated by LDL/Lp(a) apheresis. Atherosclerosis Supplements, 2017, 30, 128-134.	1.2	11
133	Lack of an association between connexin-37, stromelysin-1, plasminogen activator-inhibitor type 1 and lymphotoxin-alpha genes and acute coronary syndrome in Czech Caucasians. Experimental and Clinical Cardiology, 2010, 15, e52-6.	1.3	11
134	Dietary intake reports fidelity-fact or fiction?. Neuroendocrinology Letters, 2011, 32 Suppl 2, 29-31.	0.2	11
135	Do common genetic variants in endotoxin signaling pathway contribute to predisposition to alcoholic liver cirrhosis?. Clinical Chemistry and Laboratory Medicine, 2009, 47, 398-404.	2.3	10
136	Rs6922269 marker at the MTHFD1L gene predict cardiovascular mortality in males after acute coronary syndrome. Molecular Biology Reports, 2015, 42, 1289-1293.	2.3	10
137	Frequency of adult type-associated lactase persistence LCT-13910C/T genotypes in the Czech/Slav and Czech Roma/Gypsy populations. Genetics and Molecular Biology, 2017, 40, 450-452.	1.3	10
138	The Gene Score for Predicting Hypertriglyceridemia: New Insights from a Czech Case–Control Study. Molecular Diagnosis and Therapy, 2019, 23, 555-562.	3.8	10
139	Different prevalence of T2DM risk alleles in Roma population in comparison with the majority Czech population. Molecular Genetics & Enomic Medicine, 2020, 8, e1361.	1.2	10
140	The Impact of Physical Activity and Dietary Measures on the Biochemical and Anthropometric Parameters in Obese Children. Is There Any Genetic Predisposition?. Central European Journal of Public Health, 2015, 23, S62-S66.	1.1	10
141	<i>ADH1B</i> Polymorphism, Alcohol Consumption, and Binge Drinking in Slavic Caucasians: Results from the Czech HAPIEE Study. Alcoholism: Clinical and Experimental Research, 2012, 36, 900-905.	2.4	9
142	Lack of an association between left-handedness and <i>APOE</i> polymorphism in a large sample of adults: Results of the Czech HAPIEE study. Laterality, 2013, 18, 513-519.	1.0	9
143	Lack of an association between SNPs within the cholinergic receptor genes and smoking behavior in a Czech post-MONICA study. Genetics and Molecular Biology, 2014, 37, 625-630.	1.3	9
144	SNPs within CHRNA5-A3-B4 and CYP2A6/B6 are associated with smoking dependence but not with tobacco dependence treatment outcomes in the Czech population. Gene, 2017, 606, 35-38.	2.2	9

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145	The fat mass and obesity related gene polymorphism influences the risk of rejection in heart transplant patients. Clinical Transplantation, 2018, 32, e13443.	1.6	9
146	Analysis of the Potential Role of Apolipoprotein E Polymorphism in Genetic Predisposition to Spontaneous Abortion. American Journal of Reproductive Immunology, 2012, 67, 179-183.	1.2	8
147	The association between the FTO gene variant and alcohol consumption and binge and problem drinking in different gene-environment background: The HAPIEE study. Gene, 2019, 707, 30-35.	2.2	8
148	Alpha-1 Antitrypsin and Hepatocellular Carcinoma in Liver Cirrhosis: SERPINA1 MZ or MS Genotype Carriage Decreases the Risk. International Journal of Molecular Sciences, 2021, 22, 10560.	4.1	8
149	Serum Bilirubin in the Czech Population ― Relationship to the Risk of Myocardial Infarction in Males ―. Circulation Journal, 2020, 84, 1779-1785.	1.6	8
150	Body Composition Changes in Adult Females after Lifestyle Intervention Are Influenced by the NYD-SP18 Variant. Central European Journal of Public Health, 2015, 23, S19-S22.	1.1	8
151	Genetic determination of the prognosis in survivors of acute coronary syndromes. Study design and rationale for a multicenter study. Cor Et Vasa, 2007, 49, 134-137.	0.1	8
152	G-Protein Beta-3-Subunit and eNOS Gene Polymorphism in Transplant Recipients with Long-Term Renal Graft Function. Kidney and Blood Pressure Research, 2002, 25, 245-249.	2.0	7
153	No relation between angiotensinâ€converting enzyme gene polymorphism and smoking dependence. Scandinavian Journal of Clinical and Laboratory Investigation, 2004, 64, 575-578.	1.2	7
154	Ghrelin variants influence development of body mass index and plasma levels of total cholesterol in dialyzed patients. Clinical Chemistry and Laboratory Medicine, 2007, 45, 1121-3.	2.3	7
155	Detection of Donor DNA After Heart Transplantation: How Far Could It Be Affected by Blood Transfusion and Donor Chimerism?. Transplantation Proceedings, 2007, 39, 1593-1595.	0.6	7
156	FAT MASS AND OBESITYâ€ASSOCIATED (FTO) GENE AND ALCOHOL INTAKE. Addiction, 2012, 107, 1185-1186.	3.3	7
157	The association between APOA5 haplotypes and plasma lipids is not modified by energy or fat intake: The Czech HAPIEE study. Nutrition, Metabolism and Cardiovascular Diseases, 2014, 24, 243-247.	2.6	7
158	Effortâ€"reward imbalance at work, over-commitment personality and diet quality in Central and Eastern European populations. British Journal of Nutrition, 2016, 115, 1254-1264.	2.3	7
159	Body Adiposity Changes After Lifestyle Interventions in Children/Adolescents and the NYD-SP18 and TMEM18 Variants. Medical Science Monitor, 2018, 24, 7493-7498.	1.1	7
160	Mediterranean diet and physical functioning trajectories in Eastern Europe: Findings from the HAPIEE study. PLoS ONE, 2018, 13, e0200460.	2.5	7
161	Influence of lipoprotein apheresis on circulating plasma levels of miRNAs in patients with high Lp(a). Atherosclerosis Supplements, 2019, 40, 12-16.	1.2	7
162	Five genetic polymorphisms of cytochrome P450 enzymes in the Czech non-Roma and Czech Roma population samples. Drug Metabolism and Personalized Therapy, 2020, .	0.6	7

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163	Distribution of ADH1B genotypes predisposed to enhanced alcohol consumption in the Czech Roma/Gypsy population. Central European Journal of Public Health, 2018, 26, 284-288.	1.1	7
164	Actigenetic of ACE gene polymorphism in Czech obese sedentary females. Physiological Research, 2009, 58 Suppl 1, S47-S52.	0.9	7
165	Strong Gender-Specific Additive Effects of the NYD-SP18 and FTO Variants on BMI Values. Physiological Research, 2015, 64, S419-S426.	0.9	7
166	Apolipoprotein A-V variant (T-1131>C) affects plasma levels of non-high-density lipoprotein cholesterol in Caucasians. Experimental and Clinical Cardiology, 2008, 13, 129-32.	1.3	7
167	Genetic and biochemical characteristics in the Roma minority in the South Bohemia Region. Neuroendocrinology Letters, 2015, 36 Suppl 2, 29-34.	0.2	7
168	The Influence of Gâ€Protein β3â€Subunit Gene and Endothelial Nitric Oxide Synthase Gene in Exon 7 Polymorphisms on Progression of Autosomal Dominant Polycystic Kidney Disease. Renal Failure, 2004, 26, 119-125.	2.1	6
169	<i>Pst</i> lâ€polymorphism in the human bactericidal permeability increasing protein (BPI) gene. Clinical Genetics, 1997, 52, 249-249.	2.0	6
170	Genetic determination of triglyceridemia with special focus on apolipoprotein gene variants. Clinical Lipidology, 2010, 5, 543-554.	0.4	6
171	Aldehyde dehydrogenase 2 polymorphism affects the outcome of methanol poisoning in exposed humans. Clinical Genetics, 2018, 94, 445-449.	2.0	6
172	Baseline Leptin/Adiponectin Ratio is a Significant Predictor of BMI Changes in Children/Adolescents after Intensive Lifestyle Intervention. Experimental and Clinical Endocrinology and Diabetes, 2019, 127, 691-696.	1.2	6
173	The APOE4 allele is associated with a decreased risk of retinopathy in type 2 diabetics. Molecular Biology Reports, 2021, 48, 5873-5879.	2.3	6
174	Individual DNA Methylation Pattern Shifts in Nanoparticles-Exposed Workers Analyzed in Four Consecutive Years. International Journal of Molecular Sciences, 2021, 22, 7834.	4.1	6
175	A Polymorphism in the Cyclooxygenase 2 Gene in Type 1 Diabetic Patients With Nephropathy. Physiological Research, 2011, 60, 377-380.	0.9	6
176	Apolipoprotein AV gene polymorphisms (T-1131/C and Ser19/Trp) influence plasma triglyceride levels and risk of myocardial infarction. Experimental and Clinical Cardiology, 2003, 8, 151-4.	1.3	6
177	The Relationship between Epigenetic Age and Myocardial Infarction/Acute Coronary Syndrome in a Population-Based Nested Case-Control Study. Journal of Personalized Medicine, 2022, 12, 110.	2.5	6
178	Genetic polymorphisms of TGF-beta, PAI-1, and COL1A-1, and determination of bone mineral density in Caucasian females. Endocrine Regulations, 2006, 40, 77-81.	1.3	6
179	Possible gene-gender interaction between the SLCO1B1 polymorphism and statin treatment efficacy. Neuroendocrinology Letters, 2012, 33 Suppl 2, 22-5.	0.2	6
180	Alternative method for diagnosis of two polymorphisms in the human transforming growth factor- \hat{l}^21 by PCR-mediated double site-directed mutagenesis. Clinica Chimica Acta, 2000, 295, 187-191.	1.1	5

#	Article	IF	CITATIONS
181	Myxoma of Donor Origin in a Transplanted Heart. Journal of Heart and Lung Transplantation, 2007, 26, 865-867.	0.6	5
182	Genetic markers in hypercholesterolemic and normocholesterolemic Czech children. Clinical Genetics, 1994, 46, 88-91.	2.0	5
183	No significant association between A-501C single nucleotide polymorphism in preproghrelin and body mass index or waist-to-hip ratio in central European population. Metabolism: Clinical and Experimental, 2008, 57, 1016-1017.	3.4	5
184	Apolipoprotein E Arg136â†'Cys mutation and hyperlipidemia in a large central European population sample. Clinica Chimica Acta, 2008, 388, 217-218.	1.1	5
185	Lack of association between a new tag SNP in the FTO gene and BMI in Czech–Slavonic population. European Journal of Human Genetics, 2010, 18, 1274-1274.	2.8	5
186	Smoking impairs and circulating stem cells favour the protective effect of the T allele of the connexin37 gene in ischemic heart disease – A multinational study. Atherosclerosis, 2016, 244, 73-78.	0.8	5
187	The relationship between body mass index and 10-year trajectories of physical functioning in middle-aged and older Russians: Prospective results of the Russian HAPIEE study. Journal of Nutrition, Health and Aging, 2017, 21, 381-388.	3.3	5
188	Smoking, alcohol and cancer mortality in Eastern European men: Findings from the PrivMort retrospective cohort study. International Journal of Cancer, 2018, 143, 1128-1133.	5.1	5
189	A Newly Observed Mutation of the ABCA3 Gene Causing Lethal Respiratory Failure of a Full-Term Newborn: A Case Report. Frontiers in Genetics, 2020, 11, 568303.	2.3	5
190	Apolipoprotein E/intrauterine undernutrition interaction and hypercholesterolemia in children. Physiological Research, 2008, 57, 965-968.	0.9	5
191	An AHSG gene variant modulates basal metabolic rate and body composition development after a short-time lifestyle intervention. Neuroendocrinology Letters, 2011, 32 Suppl 2, 32-6.	0.2	5
192	IL-10 and TNF-Î ² gene polymorphisms have no major influence on lactate levels after cardiac surgeryâ [†] . European Journal of Cardio-thoracic Surgery, 2006, 30, 54-58.	1.4	4
193	Heme Oxygenase-1 Polymorphisms and Renal Transplantation Outcomes: Balancing at the Detection Limit of Allelic Association Studies. American Journal of Transplantation, 2008, 8, 1077-1078.	4.7	4
194	High Prevalence of Microchimerism in Female Patients. Transplantation Proceedings, 2008, 40, 3685-3687.	0.6	4
195	Lack of an Association Between Three Tagging SNPs Within The FTO gene and Smoking Behavior. Nicotine and Tobacco Research, 2012, 14, 998-1002.	2.6	4
196	Variant within CELSR2/PSRC1/SORT1, but not within CILP2/PBX4, PCSK9 and APOB genes, has a potential to influence statin treatment efficacy. Journal of Applied Biomedicine, 2012, 10, 19-28.	1.7	4
197	The risk of sporadic colorectal cancer development is not influenced by fat mass and obesity related gene polymorphism in Slavs. European Journal of Internal Medicine, 2012, 23, e175-e176.	2.2	4
198	APOA5 haplotypes determine triglyceride decrease after lifestyle induced weight loss in children. Nutrition, Metabolism and Cardiovascular Diseases, 2012, 22, e22-e23.	2.6	4

#	Article	IF	CITATIONS
199	COQ2 polymorphisms are not associated with increased risk of statin-induced myalgia/myopathy in the Czech population. Drug Metabolism and Personalized Therapy, 2017, 32, 177-182.	0.6	4
200	Leukocyte telomere length is not affected by long-term occupational exposure to nano metal oxides. Industrial Health, 2019, 57, 741-744.	1.0	4
201	Extending Adjuvant Endocrine Therapy for 10 Years: A Mixed-Methods Analysis of Women's Decision Making in an Online Breast Cancer Forum. Healthcare (Switzerland), 2021, 9, 688.	2.0	4
202	The effect of long-term left ventricular assist device support on flow-sensitive plasma microRNA levels. International Journal of Cardiology, 2021, 339, 138-143.	1.7	4
203	Association between apolipoprotein B promotor haplotypes and cholesterol status. Annals of Clinical Biochemistry, 2001, 38, 399-400.	1.6	3
204	Apolipoprotein CI, and not apolipoprotein E, polymorphism affects plasma levels of C-reactive protein?. Nutrition, Metabolism and Cardiovascular Diseases, 2005, 15, 450-451.	2.6	3
205	Copy Number of the Mitochondrial DNA of Leucocytes as an Aging Marker and Risk Factors for the Development of Age-Related Diseases in Humans. Advances in Gerontology, 2020, 10, 1-8.	0.4	3
206	SNPs within CHRNA5-A3-B4 and CYP2A6/B6, nicotine metabolite concentrations and nicotine dependence treatment success in smokers. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2021, 165, 84-89.	0.6	3
207	Novel mutations in TRPM6 gene associated with primary hypomagnesemia with secondary hypocalcemia. Case report. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2021, 165, 454-457.	0.6	3
208	MRAS gene marker rs9818870 is not associated with acute coronary syndrome in the Czech population and does not predict mortality in males after acute coronary syndrome. Advances in Clinical and Experimental Medicine, 2017, 26, 1213-1217.	1.4	3
209	Volunteer's willingness to genetic testing - lack of the understanding of the matter. Physiological Research, 2009, 58 Suppl 1, S53-S54.	0.9	3
210	ARH missense polymorphisms and plasma cholesterol levels. Clinical Chemistry and Laboratory Medicine, 2004, 42, 989-90.	2.3	2
211	APOAV (T-1131>C) variant has no effect on mother's height in a large population study. Lipids in Health and Disease, 2004, 3, 10.	3.0	2
212	Association of the $\hat{a} \in 159$ C $\hat{a} \in 12$ T polymorphism in the CD14 promoter with variations in serum lipoproteins in healthy subjects Blood Coagulation and Fibrinolysis, 2004, 15, 365-366.	1.0	2
213	Elevated C-reactive protein levels in individuals with low high-density lipoprotein cholesterol levels. Metabolism: Clinical and Experimental, 2005, 54, 1266-1267.	3.4	2
214	Lack of Association Between Variants within the AHSG, HCRT and NPY2R Genes and Anthropometrical Parameters in Czech Post-Monica Study. Balkan Journal of Medical Genetics, 2013, 16, 63-68.	0.5	2
215	Response to â€~CYP2E1Polymorphism and Better Outcome After Methanol Poisoning'. Basic and Clinical Pharmacology and Toxicology, 2015, 117, 3-4.	2.5	2
216	Lack of Association between NYD-SP18 Variant and Obesity. The Health Alcohol and Psychosocial Factors in Eastern Europe Study. Annals of Nutrition and Metabolism, 2016, 68, 244-248.	1.9	2

#	Article	IF	CITATIONS
217	FTOgene variant and risk of spontaneous abortion. Acta Obstetricia Et Gynecologica Scandinavica, 2016, 95, 118-118.	2.8	2
218	Hypertension and consequent mortality risk in China. Journal of Epidemiology and Community Health, 2018, 72, 1071-1072.	3.7	2
219	Association between aortic telomere length and cardiac post-transplant allograft function. International Journal of Cardiology, 2019, 290, 129-133.	1.7	2
220	Household availability of dietary fats and cardiovascular disease and mortality: prospective evidence from Russia. European Journal of Public Health, 2021, 31, 1037-1041.	0.3	2
221	An Integrative Study of Aortic mRNA/miRNA Longitudinal Changes in Long-Term LVAD Support. International Journal of Molecular Sciences, 2021, 22, 7414.	4.1	2
222	Multiplex Protein Biomarker Profiling in Patients with Familial Hypercholesterolemia. Genes, 2021, 12, 1599.	2.4	2
223	CHARACTERISTICS OF MAIN DRUG THERAPY TYPES IN SUBJECTS WITH ATRIAL FIBRILLATION IN POPULATION. Cardiovascular Therapy and Prevention (Russian Federation), 2018, 17, 43-48.	1.4	2
224	No association betweenCD14(C-260→T) variant and plasma triglycerides or body mass index in non-diabetic Caucasians. Diabetic Medicine, 2007, 24, 99-100.	2.3	1
225	Increased prevalence of the CVD-associated ANRIL allele in the Roma/Gypsy population in comparison with the majority Czech population. Genetics and Molecular Biology, 2021, 44, e20200405.	1.3	1
226	Genetic determination of an endothelial function and the size of the heart sections in juvenile hypertensives. Journal of Applied Biomedicine, 2006, 4, 59-65.	1.7	1
227	The relationship between dynamics of indicators of cognitive functions and status of economic activity in population with aging. Profilakticheskaya Meditsina, 2020, 23, 27.	0.6	1
228	Longitudinal trajectories of blood lipid levels in an ageing population sample of Russian Western-Siberian urban population. PLoS ONE, 2021, 16, e0260229.	2.5	1
229	Apolipoprotein L1 variability is associated with increased risk of renal failure in the Czech population. Gene, 2022, 818, 146248.	2.2	1
230	Five genetic polymorphisms of cytochrome P450 enzymes in the Czech non-Roma and Czech Roma population samples. Drug Metabolism and Drug Interactions, 2020, 35, .	0.3	1
231	Association between marital status and age dynamics of cognitive functions in ageing in a Russian population sample. Meditsinskiy Sovet, 2022, , 27-35.	0.5	1
232	Genetics of Myocardial Infarction. , 2011, , 103-111.		0
233	FTO First Intron rs1558902 Variant and Platelets Count in White Middle-aged Women: TABLE 1. Journal of Investigative Medicine, 2013, 61, 291-293.	1.6	0
234	Adult type associated lactase persistence genotypes in Czech and Roma/Gypsy populations. European Journal of Public Health, 2016, 26, .	0.3	0

#	Article	IF	CITATIONS
235	Telomere Length in Aortal Tissue and Heart Allograft Rejection in Patients after Transplantation. Journal of Heart and Lung Transplantation, 2016, 35, S194.	0.6	O
236	Can leukocyte telomere length predict survival time in heart transplant recipients over a minimal follow-up of 20 years?. Atherosclerosis, 2017, 263, e278-e279.	0.8	0
237	Connexin 37 gene polymorphism and atherosclerotic changes in women with diabetes type 1 and 2. Atherosclerosis, 2018, 275, e184-e185.	0.8	0
238	Short-term trajectories of exercise-induced plasma lipid changes in overweight females, with a focus on HDL-cholesterol. Advances in Clinical and Experimental Medicine, 2021, 30, 239-243.	1.4	0
239	A case of homozygous familial hypercholesterolemia with an atypical phenotype and delayed clinical symptoms. Journal of Clinical Lipidology, 2021, 15, 435-440.	1.5	0
240	Plasma levels of thiocyanate in the Czech population and their genetic determination. Cor Et Vasa, 2006, 48, 262-266.	0.1	0
241	Effect of ghrelin variants on weight loss after dietary/exercise intervention. Cor Et Vasa, 2007, 49, 213-215.	0.1	0
242	Detecting free plasma DNA in transplant medicine. Possibilities and pitfalls. Cor Et Vasa, 2008, 50, 322-327.	0.1	0
243	Oxide synthase donor gene polymorphisms and cardiac graft coronary artery disease. Cor Et Vasa, 2009, 51, 202-209.	0.1	0
244	Genes and Plasma Lipids in Czech Slavic Population. , 2011, , 149-157.		0
245	Global DNA methylation in rats´ liver is not affected by hypercholesterolemic diet. Physiological Research, 2020, 69, 347-352.	0.9	0
246	Number of children is associated with plasma CRP levels. Neuroendocrinology Letters, 2011, 32 Suppl 2, 21-3.		
	21 5.	0.2	0
247	Comparison of relative telomere length measured in aortic tissue and leukocytes in patients with end stage heart failure. Neuroendocrinology Letters, 2016, 37, 124-8.	0.2	0
247	Comparison of relative telomere length measured in aortic tissue and leukocytes in patients with end		
	Comparison of relative telomere length measured in aortic tissue and leukocytes in patients with end stage heart failure. Neuroendocrinology Letters, 2016, 37, 124-8. The associations of physical incapacity and wealth with remaining in paid employment after age 60 in	0.2	0
248	Comparison of relative telomere length measured in aortic tissue and leukocytes in patients with end stage heart failure. Neuroendocrinology Letters, 2016, 37, 124-8. The associations of physical incapacity and wealth with remaining in paid employment after age 60 in five middle-income and high-income countries. Ageing and Society, 2023, 43, 2994-3017. Lack of an Association Between a Rs9818870 Marker at the Mras Gene Locus and Acute Coronary Syndrome in Czech Males. International Journal of Systems Applications Engineering & Development,	0.2	0
248	Comparison of relative telomere length measured in aortic tissue and leukocytes in patients with end stage heart failure. Neuroendocrinology Letters, 2016, 37, 124-8. The associations of physical incapacity and wealth with remaining in paid employment after age 60 in five middle-income and high-income countries. Ageing and Society, 2023, 43, 2994-3017. Lack of an Association Between a Rs9818870 Marker at the Mras Gene Locus and Acute Coronary Syndrome in Czech Males. International Journal of Systems Applications Engineering & Development, 2022, 16, 41-43. Triglycerides, polymorphisms and the risk of acute coronary syndrome in the Czech population.	0.2	0 0