Agnieszka BoroÅ„

List of Publications by Year
in descending order

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50	414	687220	794469
papers	citations	13	19
		h-index	g-index

50	50	50	826
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Consequences of the rs6265 (Val66Met) polymorphism in the BDNF gene in selected mental disorders and sport. Current Problems of Psychiatry, 2022, 23, 24-33.	0.1	0
2	An association of ABCG8: rs11887534 polymorphism and HDL-cholesterol response to statin treatment in the Polish population. Pharmacological Reports, 2021, 73, 1781-1786.	1.5	1
3	Influence of DAT1 Promotor Methylation on Sports Performance. Genes, 2021, 12, 1425.	1.0	6
4	Epigenetic impact of the parents' physical activity on the health of their children. Baltic Journal of Health and Physical Activity, 2021, 13, 87-95.	0.2	4
5	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.	0.3	0
6	Assessment of nursesâ€™ knowledge of nosocomial infections transmitted through contact. Zdrowie Publiczne, 2020, 130, 26-29.	0.2	0
7	Measurement of the dimensions of personality traits in patients addicted to psychoactive substances in context of relapses. Current Problems of Psychiatry, 2020, 21, 203-209.	0.1	0
8	Association of DRD2 (rs 1799732), ANKK1 (rs1800497), DAT (rs28363170), DRD4 (exon 3 - VNTR) gene polymorphisms in the context of relapses in therapy.. Current Problems of Psychiatry, 2020, 21, 193-202.	0.1	1
9	Burnout syndrome among nurses of hospitals in WrocÅ‚aw, including seniority assessed with Maslach Burnout Inventory (MBI) and Work Satisfaction Scale (WSS). Zdrowie Publiczne, 2020, 130, 22-25.	0.2	0
10	Two Functional TP53 Genetic Variants and Predisposition to Keloid Scarring in Caucasians. Dermatology Research and Practice, 2019, 2019, 1-5.	0.3	1
11	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
12	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
13	Association Between RET (rs1800860) and GFRA1 (rs45568534, rs8192663, rs181595401,) Tj ETQq1 1 0.784314 rgBT /O Molecular Biomarkers, 2016, 20, 624-628.	0.3	4
14	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
15	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.	0.3	0
16	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.	0.6	3
17	Prevalence and Clinical Outcome of CYP21A2 Gene Mutations in Patients with Nonfunctional Adrenal Incidentalomas. Hormone and Metabolic Research, 2015, 47, 662-667.	0.7	8
18	Ocena czÄ™stoÅ›ci wystÄ™powania mutacji genÃ³w BRAF, KRas oraz metylacji genu RASSF1A w wolu guzkowym na podstawie badania materiaÅ‚u cytologicznego uzyskanego drogÄ… biopsji aspiracyjnej cienkoigÅ‚owej. Endokrynologia Polska, 2015, 66, 384-393.	0.3	3

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19	THORACIC SURGERY Genetic basis of keloid formation in wounds after cardiac surgery. Kardiochirurgia I Torakochirurgia Polska, 2014, 3, 273-277.	0.1	3
20	New insights into the diagnosis of nodular goiter. Thyroid Research, 2014, 7, 6.	0.7	11
21	PTPN22 1858C>T gene polymorphism in patients with SLE: association with serological and clinical results. Molecular Biology Reports, 2014, 41, 6195-6200.	1.0	18
22	Possible association of ABCB1:c.3435T>C polymorphism with high-density-lipoprotein-cholesterol response to statin treatment - a pilot study.. Bosnian Journal of Basic Medical Sciences, 2014, 14, 144-149.	0.6	15
23	Thymidylate Synthase Gene Polymorphism and Survival of Colorectal Cancer Patients Receiving Adjuvant 5-Fluorouracil. Genetic Testing and Molecular Biomarkers, 2013, 17, 799-806.	0.3	12
24	Association of BMPR1A polymorphism, but not BMP4, with kidney size in full-term newborns. Pediatric Nephrology, 2013, 28, 433-438.	0.9	7
25	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.	1.0	3
26	Association of 1936A > G inAKAP10(A-kinase anchoring protein 10) and blood pressure in Polish full-term newborns. Blood Pressure, 2013, 22, 51-56.	0.7	4
27	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.	0.2	1
28	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.	0.9	5
29	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.	0.6	2
30	The Common C49620T Polymorphism in the Sulfonylurea Receptor Gene SUR1 (ABCC8) in Patients with Gestational Diabetes and Subsequent Glucose Metabolism Abnormalities. Experimental Diabetes Research, 2012, 2012, 1-7.	3.8	2
31	The influence of polymorphism of the MUC7 gene on the teeth and dental hygiene of students at a faculty of dentistry in Poland. Postepy Higieny I Medycyny Doswiadczalnej, 2012, 66, 204-209.	0.1	9
32	Spanish study of anticoagulation in haemodialysis. Nefrologia, 2012, 32, 143-52.	0.2	13
33	The associations between G972R polymorphism of the IRS-1 gene, insulin resistance, salt sensitivity and non-dipper hypertension. Hypertension Research, 2011, 34, 1082-1086.	1.5	17
34	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.	0.3	1
35	AMPD1 gene mutations are associated with obesity and diabetes in Polish patients with cardiovascular diseases. Journal of Applied Genetics, 2011, 52, 67-76.	1.0	16
36	The distribution of human endogenous retrovirus K-113 in health and autoimmune diseases in Poland. Rheumatology, 2011, 50, 1310-1314.	0.9	24

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37	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.	0.5	1
38	The association of â€“262C/T polymorphism in the catalase gene and delayed graft function of kidney allografts. Nephrology, 2010, 15, 587-591.	0.7	4
39	Polymorphisms of Superoxide Dismutase, Glutathione Peroxidase andÂCatalase Genes in Patients with Post-transplant Diabetes Mellitus. Archives of Medical Research, 2010, 41, 350-355.	1.5	23
40	The association between eNOS intron 4 VNTR polymorphism and delayed graft function of kidney allografts. Clinical Transplantation, 2010, 24, E130-6.	0.8	4
41	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.	0.6	28
42	Metabolic Syndrome Components in Patients with Autosomal-Dominant Polycystic Kidney Disease. Kidney and Blood Pressure Research, 2009, 32, 405-410.	0.9	20
43	Sequence variants of chemokine receptor genes and susceptibility to HIV-1 infection. Journal of Applied Genetics, 2009, 50, 159-166.	1.0	16
44	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
45	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.	0.3	5
46	â€œTreasure your exceptionsâ€• recent advances in molecular genetics of glomerular disease. Journal of Applied Genetics, 2008, 49, 93-99.	1.0	2
47	ADA*2 Allele of the Adenosine Deaminase Gene May Protect against Coronary Artery Disease. Cardiology, 2007, 108, 275-281.	0.6	31
48	DNA microsatellite analysis in families with autosomal dominant polycystic kidney disease (ADPKD): the first Polish study. Journal of Applied Genetics, 2006, 47, 383-389.	1.0	5
49	Report on the D32 CCR5 variant in the Sudanese Shagia tribe. Anthropological Review, 0, 71, 71-76.	0.2	2
50	ADPKD protects against diabetogenic effects associated with genetically-predicted lactase persistence.. Archives of Medical Science, 0, , .	0.4	0