

Nadica Matevska-Geshkovska

List of Publications by Year in descending order

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21
papers

496
citations

759233

12
h-index

677142

22
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22
all docs

22
docs citations

22
times ranked

1150
citing authors

#	ARTICLE	IF	CITATIONS
1	The impact of molecular tumor profiling on the design strategies for targeting myeloid leukemia and EGFR/CD44-positive solid tumors. <i>Beilstein Journal of Nanotechnology</i> , 2021, 12, 375-401.	2.8	1
2	AKR1D1*36 C>T (rs1872930) allelic variant is associated with variability of the CYP2C9 genotype predicted pharmacokinetics of ibuprofen enantiomers – a pilot study in healthy volunteers. <i>Acta Pharmaceutica</i> , 2019, 69, 399-412.	2.0	3
3	<p><p>The AKR1D1*36 (rs1872930) Allelic Variant Is Independently Associated With Clopidogrel Treatment Outcome</p>. <i>Pharmacogenomics and Personalized Medicine</i>, 2019, Volume 12, 287-295.</p>	0.7	2
4	Influence of MSI and 18q LOH markers on capecitabine adjuvant monotherapy in colon cancer patients. <i>Pharmacogenomics and Personalized Medicine</i> , 2018, Volume 11, 193-203.	0.7	3
5	Efficacy assessment of self-assembled PLGA-PEG-PLGA nanoparticles: Correlation of nano-bio interface interactions, biodistribution, internalization and gene expression studies. <i>International Journal of Pharmaceutics</i> , 2017, 533, 389-401.	5.2	27
6	Loss of Y Chromosome in Peripheral Blood of Colorectal and Prostate Cancer Patients. <i>PLoS ONE</i> , 2016, 11, e0146264.	2.5	79
7	Association of Single-Nucleotide Polymorphism C3435T in the ABCB1 Gene with Opioid Sensitivity in Treatment of Postoperative Pain. <i>Prilozi - Makedonska Akademija Na Naukite I Umetnostite Oddelenie Za Medicinski Nauki</i> , 2016, 37, 73-80.	0.5	12
8	Search for the presence of occult hepatitis C in patients with treatment-induced viral clearance using an ultrasensitive assay. <i>Srpski Arhiv Za Celokupno Lekarstvo</i> , 2016, 144, 418-423.	0.2	1
9	Single nucleotide polymorphisms near IL28B gene and response to treatment of chronic hepatitis C in hemodialysis patients. <i>Renal Failure</i> , 2015, 37, 1180-1184.	2.1	2
10	Genetic predictors of the response to the treatment of hepatitis C virus infection. <i>Bosnian Journal of Basic Medical Sciences</i> , 2015, 15, 55-9.	1.0	4
11	Distribution of the Most Common Genetic Variants Associated with a Variable Drug Response in the Population of the Republic of Macedonia. <i>Balkan Journal of Medical Genetics</i> , 2014, 17, 5-14.	0.5	14
12	Review: Occult hepatitis C virus infection: Still remains a controversy. <i>Journal of Medical Virology</i> , 2014, 86, 1491-1498.	5.0	21
13	Association between Gene Polymorphism of Manganese Superoxide Dismutase and Prostate Cancer Risk. <i>Journal of Biochemical and Molecular Toxicology</i> , 2013, 27, 213-218.	3.0	10
14	Influence of the SCN1A IVS5N + 5 G>A Polymorphism on Therapy with Carbamazepine for Epilepsy. <i>Balkan Journal of Medical Genetics</i> , 2012, 15, 19-24.	0.5	15
15	Association of <i>GPX1</i> polymorphism, GPX activity and prostate cancer risk. <i>Human and Experimental Toxicology</i> , 2012, 31, 24-31.	2.2	21
16	Promoter length polymorphism in UGT1A1 and the risk of sporadic colorectal cancer. <i>Cancer Genetics</i> , 2012, 205, 163-167.	0.4	28
17	The association of C3435T single-nucleotide polymorphism, Pgp-glycoprotein gene expression levels and carbamazepine maintenance dose in patients with epilepsy. <i>Neuropsychiatric Disease and Treatment</i> , 2012, 8, 191.	2.2	13
18	Analysis of TSHZ2 and TSHZ3 genes in congenital pelvi-ureteric junction obstruction. <i>Nephrology Dialysis Transplantation</i> , 2010, 25, 54-60.	0.7	28

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19	Glutathione peroxidase 1 (GPX1) genetic polymorphism, erythrocyte GPX activity, and prostate cancer risk. <i>International Urology and Nephrology</i> , 2009, 41, 63-70.	1.4	100
20	Increased oxidative/nitrosative stress and decreased antioxidant enzyme activities in prostate cancer. <i>Clinical Biochemistry</i> , 2009, 42, 1228-1235.	1.9	108
21	Methylenetetrahydrofolate Reductase C677T Polymorphism and Risk of Colorectal Cancer in the Macedonian Population. <i>Balkan Journal of Medical Genetics</i> , 2008, 11, 17-24.	0.5	1