Natalia G Gorovenko

List of Publications by Year
in descending order

28	211	1477746 6	996533 15
papers	citations	h-index	g-index

29	29	29	279
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Beclomethasone/formoterol versus budesonide/formoterol combination therapy in asthma. European Respiratory Journal, 2007, 29, 682-689.	3.1	127
2	Mutations c.459+1G>A and p.P426L in the ARSA gene: Prevalence in metachromatic leukodystrophy patients from European countries. Molecular Genetics and Metabolism, 2005, 86, 353-359.	0.5	27
3	Modifying effects of TNF-Î±, IL-6 and VDR genes on the development risk and the course of COVID-19. Pilot study. Drug Metabolism and Personalized Therapy, 2022, 37, 133-139.	0.3	12
4	Variation in particular biochemical indicators, cytokine and adipokine profiles of the blood, and the structural and functional parameters of the liver in patients with nonalcoholic fatty liver disease and different genotypes by the polymorphic locus A313G of the GSTP1 gene. Cytology and Genetics, 2017, 51, 455-461.	0.2	7
5	Novel mutations in arylsulfatase A gene in three Ukrainian families with metachromatic leukodystrophy. Molecular Genetics and Metabolism, 2003, 80, 360-363.	0.5	6
6	Duplication of (12)(pter-q13.3) combined with deletion of (22)(pter-q11.2) in a patient with features of both chromosome aberrations. European Journal of Medical Genetics, 2007, 50, 128-132.	0.7	6
7	Application of multiplex PCR with histopathologic features for detection of familial breast cancer in formalin-fixed, paraffin-embedded histological specimens. Cytology and Genetics, 2008, 42, 120-126.	0.2	4
8	High Prevalence of c.1528G>C Rearrangement in Patients with Long Chain 3-Hydroxyacyl-CoA Dehydrogenase Deficiency from Ukraine. Cytology and Genetics, 2018, 52, 198-203.	0.2	3
9	Study on association of the polymorphic variants of ACE (I/D), AT2R1 (A1166C), TNF-α (G308A), MTHFR (C677T) genes and their combinations with the risk of development of perinatal pathology and gestation reduction. Biopolymers and Cell, 2011, 27, 206-213.	0.1	3
10	ASSOCIATION OF THE ADRB2 GENE POLYMORPHIC VARIANT C79G (rs1072714) WITH THE COURSE OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE IN OBESE AND NON-OBESE PATIENTS. CBU International Conference Proceedings, 0, 6, 960-965.	0.0	3
11	Specificities of Sanfilippo A syndrome laboratory diagnostics. Biopolymers and Cell, 2014, 30, 388-393.	0.1	2
12	The frequency of alleles in the GSTT1 and GSTM1 genes involved in phase II of xenobiotic transformation in long-lived people of subcarpathia. Advances in Gerontology, 2014, 4, 123-127.	0.1	2
13	Clinical and Prognostic Value of Calcium and Phosphorus Levels as Possible Markers of Endothelial Dysfunction in Preterm Infants. Journal of Neonatology, 2021, 35, 108-112.	0.0	2
14	Analysis of mutations in GBA gene in Ukrainian patients with Gaucher disease. Biopolymers and Cell, 2017, 33, 34-47.	0.1	2
15	Identification and characterization of six new mutations in GLB1 gene in Ukrainian patients with GM1 gangliosidosis and Morquio B disease. Biopolymers and Cell, 2016, 32, 450-460.	0.1	2
16	GENETIC POLYMORPHISM OF GLUTATHIONE-S-TRANSFERASE M1,T1 AND SUSCEPTIBILITY TO PREMENSTRUAL ASTHMA IN UKRAINIAN POPULATION. Chest, 2005, 128, 239S.	0.4	1
17	Determination of frequencies of alleles, associated with the pseudodeficiency of lysosomal hydrolases, in population of Ukraine. Ukrainian Biochemical Journal, 2016, 88, 96-106.	0.1	1
18	Determining the frequency of common mutations in the GBA gene in patients with Gaucher disease in Ukraine. Cytology and Genetics, 2007, 41, 230-236.	0.2	0

#	Article	IF	Citations
19	The role of genetic determinant in the development of severe perinatal asphyxia. Cytology and Genetics, 2010, 44, 294-299.	0.2	0
20	Clinical Presentation of a Patient with a Severe form of Pompe Disease. Clinical Therapeutics, 2011, 33, S13.	1.1	0
21	Early detection and group-specific identification of Mycobacterium tuberculosis strains by method of single-nucleotide polymorphism analysis with hairpin primers. Biopolymers and Cell, 2013, 29, 375-381.	0.1	0
22	Effect of polymorphic variants of hereditary thrombophilia genes on the risk of early pregnancy loss for married couples. Meta Gene, 2021, 29, 100902.	0.3	0
23	The reduction of two BRCA1 gene mutations frequencies in ovarian cancer patients from Ukraine. Meta Gene, 2021, 29, 100900.	0.3	0
24	Polymorphic variants of genes ADRB2, NR3C1, MDR1 in patients with chronic obstructive pulmonary disease and obesity. , 2015, , .		0
25	Efficiency of application of different DNA probes in identifying marker chromosomes. Biopolymers and Cell, 2016, 32, 49-53.	0.1	0
26	Molecular-genetic characterization of Ukrainian patients with mucopolysaccharidosis IIIA: identification of three new mutations in the heparan-N-sulfatase gene. Biopolymers and Cell, 2016, 32, 359-366.	0.1	0
27	Spectrum of mutations in patients with organic acidurias from Ukraine. Biopolymers and Cell, 2018, 34, 107-116.	0.1	0
28	Polymorphisms variants of the MTHFR C677T and PAI-1 5G/4G genes and their combinations in the group of children with arterial ischemic stroke. , 2020, 29, 27-32.		0