Radoslava Vasileva Vazharova

List of Publications by Year in descending order

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15	96	1684188	1372567
papers	citations	h-index	g-index
16	16	16	184
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Spinal muscular atrophy with congenital bone fractures 2 caused by a rare lossâ€ofâ€function ⟨i⟩ASCC1⟨ i⟩ gene mutation in two Bulgarian Roma patients. Clinical Genetics, 2022, 102, 78-79.	2.0	1
2	Rare genetic variants prioritize molecular pathways for semaphorin interactions in Alzheimer's disease patients. Biotechnology and Biotechnological Equipment, 2021, 35, 1256-1262.	1.3	3
3	Long-term follow-up of a female patient with non-classical $11 < b > \hat{l}^2 < /b >$ -hydroxylase deficiency and two novel mutations in <i>CYP11B1</i> . Gynecological Endocrinology, 2019, 35, 23-27.	1.7	8
4	Prioritization of genetic variants predisposing to coronary heart disease in the Bulgarian population using centenarian exomes. Biotechnology and Biotechnological Equipment, 2019, 33, 1757-1765.	1.3	1
5	SCN8A p.Arg1872Gln mutation in early infantile epileptic encephalopathy type 13: Review and case report. Biotechnology and Biotechnological Equipment, 2018, 32, 1345-1351.	1.3	3
6	Double heterozygosity of novel variants found in patients with severe clinical phenotype of cardiovascular disorders. Biotechnology and Biotechnological Equipment, 2018, 32, 679-685.	1.3	0
7	New Territory for an Old Disease: 5-Alpha-Reductase Type 2 Deficiency in Bulgaria. Sexual Development, 2017, 11, 21-28.	2.0	12
8	Screening of pharmacogenetic variants associated with drug sensitivity in patients with papillary thyroid carcinoma using next generation sequencing. Biotechnology and Biotechnological Equipment, 2017, , 1-5.	1.3	1
9	Novel Hypomorphic Mutation in FANCD2Gene Observed in a Fetus with Multiple Congenital Anomalies. Case Reports in Obstetrics and Gynecology, 2016, 2016, 1-4.	0.3	1
10	Individual capacity for DNA repair and maintenance of genomic integrity: a fertile ground for studies in the field of assisted reproduction. Biotechnology and Biotechnological Equipment, 2016, 30, 419-433.	1.3	5
11	Clinical and molecular studies of EXT1/EXT2 in Bulgaria. Journal of Inherited Metabolic Disease, 2011, 34, 917-921.	3.6	8
12	Ready, Steady, Goâ€"The Current State of Carriership Status Determination and Prenatal Diagnosis of Haemophilia a in Bulgaria. Biotechnology and Biotechnological Equipment, 2011, 25, 2566-2571.	1.3	1
13	Case-control association study of 65 candidate genes revealed a possible association of a SNP of HTR5A to be a factor susceptible to bipolar disease in Bulgarian population. Journal of Affective Disorders, 2009, 117, 87-97.	4.1	37
14	Introduction of the QF-PCR analysis for the purposes of prenatal diagnosis in Bulgaria—estimation of applicability of 6 STR markers on chromosomes 21 and 18. Prenatal Diagnosis, 2004, 24, 202-208.	2.3	15
15	Alterations of gut bacteria Akkermansia muciniphila and Faecalibacterium prausnitzii in late post-transplant period after liver transplantation. Iberoamerican Journal of Medicine, 0, , 45-51.	0.2	0