Radoslava Vasileva Vazharova

List of Publications by Year
in descending order

15 papers	96 citations	1684188 5 h-index	1372567 10 g-index
16 all docs	16 docs citations	16 times ranked	184 citing authors

#	Article	IF	Citations
1	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
2	Introduction of the QF-PCR analysis for the purposes of prenatal diagnosis in Bulgariaâ€"estimation of applicability of 6 STR markers on chromosomes 21 and 18. Prenatal Diagnosis, 2004, 24, 202-208.	2.3	15
3	New Territory for an Old Disease: 5-Alpha-Reductase Type 2 Deficiency in Bulgaria. Sexual Development, 2017, 11, 21-28.	2.0	12
4	Clinical and molecular studies of EXT1/EXT2 in Bulgaria. Journal of Inherited Metabolic Disease, 2011, 34, 917-921.	3.6	8
5	Long-term follow-up of a female patient with non-classical 11Î²-hydroxylase deficiency and two novel mutations in CYP11B1. Gynecological Endocrinology, 2019, 35, 23-27.	1.7	8
6	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
7	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
8	Rare genetic variants prioritize molecular pathways for semaphorin interactions in Alzheimerâ€™s disease patients. Biotechnology and Biotechnological Equipment, 2021, 35, 1256-1262.	1.3	3
9	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
10	Novel Hypomorphic Mutation inFANCD2Gene Observed in a Fetus with Multiple Congenital Anomalies. Case Reports in Obstetrics and Gynecology, 2016, 2016, 1-4.	0.3	1
11	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
12	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
13	Spinal muscular atrophy with congenital bone fractures 2 caused by a rare lossâ€ofâ€function ASCC1 gene mutation in two Bulgarian Roma patients. Clinical Genetics, 2022, 102, 78-79.	2.0	1
14	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
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