## Radoslava Vasileva Vazharova

## List of Publications by Year

 in descending orderSource: https:/|exaly.com/author-pdf/2573931/publications.pdf
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Rare genetic variants prioritize molecular pathways for semaphorin interactions in Alzheimerâ $€^{T M_{S}}$ disease patients. Biotechnology and Biotechnological Equipment, 2021, 35, 1256-1262.

5 SCN8A p.Arg1872GIn mutation in early infantile epileptic encephalopathy type 13: Review and case
report. Biotechnology and Biotechnological Equipment, 2018, 32, 1345-1351.

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

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New Territory for an Old Disease: 5-Alpha-Reductase Type 2 Deficiency in Bulgaria. Sexual Development, 2017, 11, 21-28.
2.0

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Screening of pharmacogenetic variants associated with drug sensitivity in patients with papillary 8 thyroid carcinoma using next generation sequencing. Biotechnology and Biotechnological Equipment, 2017, , 1-5.
$9 \quad$ Novel Hypomorphic Mutation inFANCD2Gene Observed in a Fetus with Multiple Congenital Anomalies.
$9 \quad$ Case Reports in Obstetrics and Gynecology, 2016, 2016, 1-4.

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.
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.
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Clinical and molecular studies of EXT1/EXT2 in Bulgaria. Journal of Inherited Metabolic Disease, 2011,
$11 \quad \begin{aligned} & \text { Clinical and m } \\ & 34,917-921 .\end{aligned}$
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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.
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Case-control association study of 65 candidate genes revealed a possible association of a SNP of
13 HTR5A to be a factor susceptible to bipolar disease in Bulgarian population. Journal of Affective
4.1

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Disorders, 2009, 117, 87-97.

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.

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