

مرتضی جبارپور بنیادی دانشیار دانشکده: علوم طبیعی

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دانشگاه	رشته و گرایش تحصیلی	سال اخذ مدرک	مقطع تحصيلي
دانشگاه گلاسگو	ژنتیک مولکولی پزشکی		کارشناسی ارشد
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اطلاعات استخدامي						
پایه	نوع همکاری	نوع استخدام	عنوان سمت	محل خدمت		
۲۸	تمام وقت	رسمى قطعى		دانشکده علوم طبیعی		

مقالات در نشریات

- Hasan Azizi , Mortaza Bonyadi , Abbas Rafat, A novel splice site variant of the BBS2 gene in a .1 .patient with Bardet-Biedl syndrome, Human Genome Variation, 2024 03 20
- Hossein Ahmed Hashim, Morteza Bonyadi, Seyed Abbas Rafat,IDENTIFICATION AND .2
 BIOINFORMATICS ANALYSIS OF TWO NOVEL VARIANTS IN THE SEMA4A AND SCP2 GENES IN
 A PATIENT WITH EARLY-ONSET VISUAL IMPAIRMENT AND LEUKODYSTROPHY.,Journal of
 .Population Therapeutics& Clinical Pharmacology,2024 01 20
- Sahand Mirzaei Dizaji , Amir Amandi , Morteza Bonyadi, Identification and bioinformatics .3 analysis of a novel mutation in PLA2G6 gene in a patient with neurodegenerative disorder, Gene .Reports, 2023 10 25
- Amir Reza Dalal Amandi, Neda Jabbarpour, Shadi Shiva, Mortaza Bonyadi,Identification of .4
 Two Novel Pathogenic Variants of the ATM Gene in the Iranian-Azeri Turkish Ethnic Group by
 .Applying Whole Exome Sequencing,current Genomics,2023 10 20
- Neda Jabbarpour, Morteza Bonyadi, Leyla Sadeghi, A novel loss of function mutation in the .5 PHD domain of the RAG2 gene, affecting zinc-binding affinity, Molecular Biology Reports, 2023 08

.20

- Identification of a de novo, Novel Pathogenic Variant in the Splice Region of the SOX10 Gene .6 in an Iranian Azeri Turkish Family with Waardenburg Syndrome, Molecular Syndromology, 2023 08 .20
 - Identification of a novel mutation in the HACD1 gene in an Iranian family with autosomal .7 .recessive congenital myopathy, with fibre-type disproportion, Journal of Genetics, 2023 01 20 Saba A. Nazm et al., Spectrum of MECP2 mutations in Iranian Azeri Turkish Rett syndrome .8 .patients, Neurology Asia, 2023 01 23
 - Neda Jabbarpour , Hassan Saei , Mohammad hossein Jabbrpoor bonyadi , Morteza .9 Bonyadi, Identification of novel cis-mutations in the GJA8 gene in a 3-generation Iranian family .with autosomal dominant congenital nuclear cataract, Ophthalmic Genetics, 2022 06 20 Asghari Sarfaraz A, Jabbarpour N, Bonyadi M, Khalaj ,& Kondory M., Identification and .10 bioinformatics analysis of a novel variant in the HERC2 gene in a patient with intellectual .developmental disorder., J Neurogenet., 17.06.2024