

Oliver V StojkoviÄ

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

1,294
citations

759233

12
h-index

454955

30
g-index

35
all docs

35
docs citations

35
times ranked

2395
citing authors

#	ARTICLE	IF	CITATIONS
1	Cryptic Diversity of the European Blind Mole Rat <i>Nannospalax leucodon</i> Species Complex: Implications for Conservation. <i>Animals</i> , 2022, 12, 1097.	2.3	5
2	Case Report: Post-mortem Histopathological and Molecular Analyses of the Very First Documented COVID-19-Related Death in Europe. <i>Frontiers in Medicine</i> , 2021, 8, 612758.	2.6	12
3	Combinations of fibrinolytic gene polymorphisms (plasminogen activator inhibitor type 1 4G/5G, factor) Tj ETQq1 1 0.784314 rgBT /C Coagulation and Fibrinolysis, 2021, 32, 103-107.	1.0	2
4	Polymorphisms and haplotypes in VDR gene are associated with female idiopathic infertility. <i>Human Fertility</i> , 2020, 23, 101-110.	1.7	11
5	Defining screening panel of functional variants of CYP1A1, CYP2C9, CYP2C19, CYP2D6, and CYP3A4 genes in Serbian population. <i>International Journal of Legal Medicine</i> , 2020, 134, 433-439.	2.2	7
6	16S rRNA gene polymorphism supports cryptic speciation within the lesser blind mole rat <i>Nannospalax leucodon</i> superspecies (Rodentia: Spalacidae). <i>Mammalian Biology</i> , 2020, 100, 315-324.	1.5	3
7	Hemostasis-related gene polymorphisms and their epistatic relationship in women with idiopathic infertility. <i>Blood Coagulation and Fibrinolysis</i> , 2019, 30, 253-262.	1.0	1
8	Genetics of suspected thrombophilia in Serbian females with infertility, including three cases, homozygous for <i>FII</i> 20210A or <i>FV</i> 1691A mutations. <i>Human Fertility</i> , 2017, 20, 132-139.	1.7	11
9	Prediction of autosomal STR typing success in ancient and Second World War bone samples. <i>Forensic Science International: Genetics</i> , 2017, 27, 17-26.	3.1	26
10	Polymorphisms in <i>ACE</i> and <i>ACTN3</i> Genes and Blood Pressure Response to Acute Exercise in Elite Male Athletes from Serbia. <i>Tohoku Journal of Experimental Medicine</i> , 2017, 243, 311-320.	1.2	13
11	Should MTHFR 1298 A>C be tested together with MTHFR 677 C>T polymorphism in women with reproductive challenges?. <i>Genetika</i> , 2017, 49, 377-386.	0.4	0
12	Multidisciplinary investigation links backward-speech trait and working memory through genetic mutation. <i>Scientific Reports</i> , 2016, 6, 20369.	3.3	5
13	On the Bantu expansion. <i>Gene</i> , 2016, 593, 48-57.	2.2	4
14	The angiotensin converting enzyme (ace) gene polymorphism: Insight study of the renal regulation of the arterial blood pressure. <i>Medicinski Podmladak</i> , 2016, 67, 31-35.	0.0	0
15	Monitoring of Pediatric Patients With Malignant Hematological Diseases After Allogeneic HSCT. <i>Journal of Pediatric Hematology/Oncology</i> , 2012, 34, e253-e257.	0.6	0
16	Y-chromosomal microsatellite diversity in three culturally defined regions of historical Tibet. <i>Forensic Science International: Genetics</i> , 2012, 6, 437-446.	3.1	20
17	Y-STR diversity in the Himalayas. <i>International Journal of Legal Medicine</i> , 2011, 125, 367-375.	2.2	15
18	Divergent patrilineal signals in three Roma populations. <i>American Journal of Physical Anthropology</i> , 2011, 144, 80-91.	2.1	16

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19	Y-STR profiling in two Afghanistan populations. <i>Legal Medicine</i> , 2011, 13, 103-108.	1.3	26
20	Forensic DNA databases in Western Balkan region: retrospectives, perspectives, and initiatives. <i>Croatian Medical Journal</i> , 2011, 52, 235-244.	0.7	11
21	STR loci D19S216, D20S502 and D20S842 analysis in the Serbian population using dentin DNA. <i>Archives of Biological Sciences</i> , 2011, 63, 55-58.	0.5	0
22	Human Y-chromosome short tandem repeats: A tale of acculturation and migrations as mechanisms for the diffusion of agriculture in the Balkan Peninsula. <i>American Journal of Physical Anthropology</i> , 2010, 142, 380-390.	2.1	29
23	Allele frequencies and population data for 17 Y-chromosome STR loci in a Serbian population sample from Vojvodina province. <i>Forensic Science International</i> , 2008, 176, e23-e28.	2.2	31
24	DNA typing from handled items. <i>Forensic Science International: Genetics Supplement Series</i> , 2008, 1, 411-412.	0.3	26
25	Dominant-negative mutations in the DNA-binding domain of STAT3 cause hyper-IgE syndrome. <i>Nature</i> , 2007, 448, 1058-1062.	27.8	930
26	Population Data on HLA-DQA1, LDLR, GYPA, HBGG, D7S8, and GC PCR-Based Loci in Serbia. <i>Journal of Forensic Sciences</i> , 2006, 51, 699-699.	1.6	2
27	Poly(A) tailing of ancient DNA: a method for reproducible microsatellite genotyping. <i>Analytical Biochemistry</i> , 2003, 318, 124-131.	2.4	2
28	250 CTG repeats in DMPK is a threshold for correlation of expansion size and age at onset of juvenile-adult DM1. <i>Human Mutation</i> , 2002, 19, 131-139.	2.5	30
29	Is the 31 CAG repeat allele of the spinocerebellar ataxia 1 (SCA1) gene locus non-specifically associated with trinucleotide expansion diseases?. <i>Psychiatric Genetics</i> , 2001, 11, 201-205.	1.1	5
30	Yugoslav population data on nine STR loci. <i>Forensic Science International</i> , 2001, 115, 239-240.	2.2	3
31	The Status of SCA1, MJD/SCA3, FRDA, DRPLA and MD Triplet Containing Genes in Patients with Huntington Disease and Healthy Controls. <i>Journal of Neurogenetics</i> , 2000, 14, 257-263.	1.4	9
32	Correlation between triplet repeat expansion and computed tomography measures of caudate nuclei atrophy in Huntington's disease. <i>Journal of Neurology</i> , 1999, 246, 1090-1093.	3.6	13
33	Laboratory Evolution of Life-History Traits in the Bean Weevil (<i>Acanthoscelides obtectus</i>): The Effects of Density-Dependent and Age-Specific Selection. <i>Evolution; International Journal of Organic Evolution</i> , 1997, 51, 1896.	2.3	24