

Miljenko KapoviÄ

List of Publications by Year in descending order

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63
papers

1,087
citations

393982

19
h-index

476904

29
g-index

63
all docs

63
docs citations

63
times ranked

1783
citing authors

#	ARTICLE	IF	CITATIONS
1	Association between the ACE ^{I/D} polymorphism and nicotine dependence amongst patients with lung cancer. <i>Biomedical Reports</i> , 2020, 13, 1-1.	0.9	8
2	Association of circadian rhythm genes ARNTL/BMAL1 and CLOCK with multiple sclerosis. <i>PLoS ONE</i> , 2018, 13, e0190601.	1.1	34
3	The insertion/deletion polymorphism in the angiotensin-converting enzyme gene and nicotine dependence in schizophrenia patients. <i>Journal of Neural Transmission</i> , 2017, 124, 511-518.	1.4	9
4	The lack of association between angiotensin ¹ -converting enzyme gene insertion/deletion polymorphism and nicotine dependence in multiple sclerosis. <i>Brain and Behavior</i> , 2017, 7, e00600.	1.0	3
5	Angiotensin-converting enzyme insertion/deletion gene polymorphism and interferon- γ treatment response in multiple sclerosis patients. <i>Pharmacogenetics and Genomics</i> , 2017, 27, 232-235.	0.7	7
6	Identification of rare genetic variation of NLRP1 gene in familial multiple sclerosis. <i>Scientific Reports</i> , 2017, 7, 3715.	1.6	53
7	Functional single nucleotide polymorphisms of matrix metalloproteinase 7 and 12 genes in idiopathic recurrent spontaneous abortion. <i>Journal of Assisted Reproduction and Genetics</i> , 2017, 34, 365-371.	1.2	2
8	Systematic review and meta-analysis of genetic association studies in idiopathic recurrent spontaneous abortion. <i>Fertility and Sterility</i> , 2017, 107, 150-159.e2.	0.5	110
9	Etiopathogenesis of metabolic syndrome in schizophrenia – recent findings. <i>Medicina</i> , 2017, 53, 27-42.	0.0	0
10	The Role of Iron and Iron Overload in Chronic Liver Disease. <i>Medical Science Monitor</i> , 2016, 22, 2144-2151.	0.5	133
11	Genetic heritage of Croatians in the Southeastern European gene pool – Y chromosome analysis of the Croatian continental and Island population. <i>American Journal of Human Biology</i> , 2016, 28, 837-845.	0.8	10
12	Angiotensin-converting enzyme insertion/deletion gene polymorphism in multiple sclerosis: a meta-analysis. <i>Neurological Sciences</i> , 2016, 37, 1955-1959.	0.9	4
13	Insertion/deletion polymorphism in intron 16 of ACE gene in idiopathic recurrent spontaneous abortion: case-control study, systematic review and meta-analysis. <i>Reproductive BioMedicine Online</i> , 2016, 32, 237-246.	1.1	14
14	The γ 2549 insertion/deletion polymorphism in the promoter region of the VEGFA gene in couples with idiopathic recurrent spontaneous abortion. <i>Journal of Assisted Reproduction and Genetics</i> , 2015, 32, 1789-1794.	1.2	9
15	MMP-2 γ 1575G/A polymorphism modifies the onset of optic neuritis as a first presenting symptom in MS?. <i>Journal of Neuroimmunology</i> , 2015, 286, 13-15.	1.1	8
16	Cornelia de Lange syndrome caused by heterozygous deletions of chromosome 8q24: Comments on the article by Perez ^a et al. [2012]. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 1426-1427.	0.7	8
17	A critical update on endothelial nitric oxide synthase gene variations in women with idiopathic recurrent spontaneous abortion: genetic association study, systematic review and meta-analyses. <i>Molecular Human Reproduction</i> , 2015, 21, 466-478.	1.3	14
18	Altered LINE-1 Methylation in Mothers of Children with Down Syndrome. <i>PLoS ONE</i> , 2015, 10, e0127423.	1.1	23

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19	Genes and celiac disease. Paediatrica Croatica, 2015, 59, 88-94.	0.1	0
20	Functional Polymorphisms of Matrix Metalloproteinases 1 and 9 Genes in Women with Spontaneous Preterm Birth. Disease Markers, 2014, 2014, 1-7.	0.6	13
21	The Role of TPA I/D and PAI-1 4G/5G Polymorphisms in Multiple Sclerosis. Disease Markers, 2014, 2014, 1-8.	0.6	10
22	Genetic variation in tissue inhibitors of metalloproteinases as a risk factor for idiopathic recurrent spontaneous abortion. Fertility and Sterility, 2013, 99, 1923-1929.	0.5	7
23	Y chromosome azoospermia factor region microdeletions are not associated with idiopathic recurrent spontaneous abortion in a Slovenian population: association study and literature review. Fertility and Sterility, 2013, 99, 1663-1667.	0.5	9
24	Tumor necrosis factor-alpha gene promoter -308 and -238 polymorphisms in patients with lung cancer as a second primary tumor. Medical Science Monitor, 2013, 19, 846-851.	0.5	12
25	Angiotensin-Converting Enzyme Insertion/Deletion Gene Polymorphism in Lung Cancer Patients. Genetic Testing and Molecular Biomarkers, 2012, 16, 722-725.	0.3	24
26	Matrix metalloproteinases 1, 2, 3 and 9 functional single-nucleotide polymorphisms in idiopathic recurrent spontaneous abortion. Reproductive BioMedicine Online, 2012, 24, 567-575.	1.1	33
27	The insulin-like growth factor 2 receptor gene Gly1619Arg polymorphism and idiopathic recurrent spontaneous abortion. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 429-431.	0.7	4
28	Third case of 8q23.3-q24.13 deletion in a patient with Langer-Giedion syndrome phenotype without TRPS1 gene deletion. American Journal of Medical Genetics, Part A, 2012, 158A, 659-663.	0.7	20
29	HFE mutations and transferrin C1/C2 polymorphism among Croatian patients with schizophrenia and schizoaffective disorder. Molecular Biology Reports, 2012, 39, 2253-2258.	1.0	7
30	Genetic polymorphisms of 15 STR loci in the population of the island of Cres (Croatia). Annals of Human Biology, 2011, 38, 12-21.	0.4	3
31	CTLA-4 +49 A/G gene polymorphism in Croatian and Slovenian multiple sclerosis patients. International Journal of Immunogenetics, 2011, 38, 419-426.	0.8	7
32	Angiotensin-Converting Enzyme Gene Polymorphism in Patients with Multiple Sclerosis from Bosnia and Herzegovina. Genetic Testing and Molecular Biomarkers, 2011, 15, 835-838.	0.3	5
33	Low frequency of HFE gene mutations in Croatian patients suspected of having hereditary hemochromatosis. Medical Science Monitor, 2011, 17, CR552-CR556.	0.5	2
34	A Case of Lichen Ruber Planus in a Patient with Familial Multiple Sclerosis. Journal of International Medical Research, 2010, 38, 1856-1860.	0.4	5
35	Interleukin 7 receptor alpha polymorphism rs6897932 and susceptibility to multiple sclerosis in the Western Balkans. Multiple Sclerosis Journal, 2010, 16, 533-536.	1.4	11
36	Functional inference of methylenetetrahydrofolate reductase gene polymorphisms on enzyme stability as a potential risk factor for Down syndrome in Croatia. Disease Markers, 2010, 28, 293-8.	0.6	12

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37	The Prevalence of Live Birth Down Syndrome in the Region of Primorsko-goranska County in Croatia, 1996â€“2005: The Impact of Screening and Amniocentesis. <i>Maternal and Child Health Journal</i> , 2008, 12, 620-623.	0.7	4
38	ORIGINAL ARTICLE: Genetic Predisposition to Idiopathic Recurrent Spontaneous Abortion: Contribution of Genetic Variations in IGFâ€“2 and H19 Imprinted Genes. <i>American Journal of Reproductive Immunology</i> , 2008, 60, 111-117.	1.2	24
39	BanI polymorphism of cytosolic phospholipase A2 gene is associated with age at onset in male patients with schizophrenia and schizoaffective disorder. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2008, 78, 351-360.	1.0	17
40	Pregnant Womenâ€™s Attitudes Toward Amniocentesis Before Receiving Down Syndrome Screening Results. <i>Women's Health Issues</i> , 2008, 18, 79-84.	0.9	13
41	Severe psychomotor retardation in a boy with a small supernumerary marker chromosome 19p. <i>Cytogenetic and Genome Research</i> , 2008, 121, 298-301.	0.6	6
42	PAI and TPA gene polymorphisms in multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2008, 14, 243-247.	1.4	13
43	Tumor Necrosis Factor-Î±-308 Gene Polymorphism in Croatian and Slovenian Multiple Sclerosis Patients. <i>European Neurology</i> , 2007, 57, 203-207.	0.6	22
44	Epidemiology of multiple sclerosis in western Herzegovina. <i>Clinical Neurology and Neurosurgery</i> , 2007, 109, 779-783.	0.6	15
45	Sex-specific differences of craniofacial traits in Croatia: The impact of environment in a small geographic area. <i>Annals of Human Biology</i> , 2007, 34, 296-314.	0.4	40
46	Prenatal diagnosis of complete trisomy 19q. <i>Prenatal Diagnosis</i> , 2007, 27, 644-647.	1.1	9
47	Polymorphisms in the Interleukin-12/18 Genes and Recurrent Spontaneous Abortion. <i>American Journal of Reproductive Immunology</i> , 2007, 58, 403-408.	1.2	20
48	Region with persistent high frequency of multiple sclerosis in Croatia and Slovenia. <i>Journal of the Neurological Sciences</i> , 2006, 247, 169-172.	0.3	21
49	Angiotensin-converting enzyme I/D gene polymorphism and risk of multiple sclerosis. <i>Acta Neurologica Scandinavica</i> , 2006, 114, 374-377.	1.0	20
50	Secular change of craniofacial measures in Croatian younger adults. <i>American Journal of Human Biology</i> , 2006, 18, 668-675.	0.8	28
51	No association of CCR5D32 gene mutation with multiple sclerosis in Croatian and Slovenian patients. <i>Multiple Sclerosis Journal</i> , 2006, 12, 360-362.	1.4	15
52	Human Y-specific STR haplotypes in the Western Croatian population sample. <i>Forensic Science International</i> , 2005, 149, 257-261.	1.3	17
53	LIVER AND MUSCLE TISSUE FATTY ACID COMPOSITION OF THE LIPID FRACTIONS OF DIPLodus VULGARIS FROM THE NORTH ADRIATIC SEA, CROATIA. <i>Journal of Food Lipids</i> , 2005, 12, 286-298.	0.9	3
54	Mutations in the hemochromatosis gene (HFE) and multiple sclerosis. <i>Neuroscience Letters</i> , 2005, 383, 301-304.	1.0	22

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55	Elevated Second-Trimester Free β -hCG as an Isolated Finding and Pregnancy Outcomes. <i>Fetal Diagnosis and Therapy</i> , 2004, 19, 483-487.	0.6	6
56	Composition and seasonal variation of fatty acids of <i>Diplodus vulgaris</i> L. from the Adriatic Sea. <i>JAOCS, Journal of the American Oil Chemists' Society</i> , 2004, 81, 759-763.	0.8	6
57	Secular change in body height and cephalic index of Croatian medical students (University of Rijeka). <i>American Journal of Physical Anthropology</i> , 2004, 123, 91-96.	2.1	34
58	Hemochromatosis gene mutations in the Croatian and Slovenian populations. <i>Clinical Genetics</i> , 2003, 64, 444-446.	1.0	17
59	The Influence of Smoking and Parity on Serum Markers for Downâ€™s Syndrome Screening. <i>Fetal Diagnosis and Therapy</i> , 2002, 17, 17-21.	0.6	24
60	Chromosome Studies in Patients with Defective Reproductive Success. <i>American Journal of Reproductive Immunology</i> , 2000, 44, 279-283.	1.2	17
61	Maternal Serum Screening for Down Syndrome: A Survey of Pregnant Womenâ€™s Views. <i>Public Health Genomics</i> , 1999, 2, 109-112.	0.6	7
62	Chromosomal Anomalies in Abnormal Human Pregnancies. <i>Fetal Diagnosis and Therapy</i> , 1998, 13, 187-191.	0.6	27
63	Screening for Downâ€™s Syndrome and Neural Tube Defect in Croatia. <i>Fetal Diagnosis and Therapy</i> , 1998, 13, 367-371.	0.6	7