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CCR5 Δ32 and CTLA-4 +49 A/G Gene Polymorphisms and Interferon-β Treatment Response in Croatian and Slovenian Multiple Sclerosis Patients		Nekić, Jasna; Stanković Matić, Ivana; Rački, Valentino; Janko Labinac, Dolores; Vuletić, Vladimira; Kapović, Miljenko; Ristić, Smiljana; Peterlin, Borut; Starčević Čizmarević, Nada	
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Refining the Global Phylogeny of Mitochondrial N1a, X, and HV2 Haplogroups Based on Rare Mitogenomes from Croatian Isolates		Havaš Auguštin, Dubravka; Šarac, Jelena; Reidla, Maere; Tamm, Erika; Grahovac, Blaženka; Kapović, Miljenko; Novokmet, Natalija; Rudan, Pavao; Missoni, Saša; Marjanović, Damir; Korolija, Marina	
The effects of microbiota abundance on symptom severity in Parkinson's disease: A systematic review		Papić, Eliša; Rački, Valentino; Hero, Mario; Tomić, Zoran; Starčević-Čizmarević, Nada; Kovanda, Anja; Kapović, Miljenko; Hauser, Goran; Peterlin, Borut; Vuletić, Vladimira	
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Could the CCR5-Δ32 Mutation be Protective in SARS-CoV-2 Infection?		Starčević Čizmarević, Nada; Kapović, Miljenko; Rončević, Dobrica; Ristić, Smiljana	
Association between the ACE-I/D polymorphism and nicotine dependence amongst patients with lung cancer		Nadalin, Sergej; Flego, Veljko; Pavlić, Sanja; Volarić, Darian; Badovinac, Anđelka; Kapović, Miljenko; Ristić, Smiljana	
Association of circadian rhythm genes ARNTL/BMAL1 and CLOCK with multiple sclerosis		Lavtar, Polona; Rudolf, Gorazd; Maver, Ales; Hodžić, Alenka; Čizmarević, Nada Starčević; Živkovic, Maja; Jazbec, Sasa Segal; Ketis, Zalika Klemenc; Kapović, Miljenko; Dinčić, Evica; Raičević, Ranko; Sepčić, Juraj; Lovrečić, Luca; Stanković, Aleksandra; Ristić, Smiljana; Peterlin, Borut	
Etiopatogeneza metaboličkog sindroma u shizofreniji – najnovije spoznaje		Nadalin, Sergej; Gudeljević, Marija; Severec, Josipa; Rebić, Jelena; Kapović, Miljenko; Buretić-Tomljanović, Alena	

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Angiotensin-converting enzyme insertion/deletion gene polymorphism and interferon- β treatment response in multiple sclerosis patients: a preliminary report.		Ristić, Smiljana; Starčević Čizmarević, Nada; Lavtar, Polona; Lovrečić, Luca; Perković, Olivio; Sepčić, Juraj; Šega Jazbec, Saša; Kapović, Miljenko; Peterlin, Borut.	
Functional single nucleotide polymorphisms of matrix metalloproteinase 7 and 12 genes in idiopathic recurrent spontaneous abortion		Barišić, Anita; Pereza, Nina; Hodžić, Alenka; Kapović, Miljenko; Peterlin, Borut; Ostojić, Saša	
Identification of rare genetic variation of NLRP1 gene in familial multiple sclerosis		Maver, Ales; Lavtar, Polona; Ristić, Smiljana; Stopinšek, Sanja; Simčić, Saša; Hočevar, Keli; Sepčić, Juraj; Drulović, Jelena; Pekmezović, Tatjana; Novaković, Ivana; Hodžić, Alenka; Rudolf, Gorazd; Šega, Saša; Starčević-Čizmarević, Nada; Palandačić, Anja; Zamolo, Gordana; Kapović, Miljenko; Likar, Tina; Peterlin, Borut	
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HLA-DQA1 i HLADQB1 geni u pacijenata s celjakijom		Mijandrušić Sinčić, Brankica; Starčević Čizmarević, Nada; Licul, Vanja; Crnić-Martinović, Marija; Ristić, Smiljana; Kapović, Miljenko	

Insertion/deletion polymorphism in intron 16 of ACE gene in idiopathic recurrent spontaneous abortion: case-control study, systematic review and meta-analysis		Pereza, Nina; Ostojić, Saša; Zdravčević, Matea; Volk, Marija; Kapović, Miljenko; Peterlin, Borut	
A critical update on endothelial nitric oxide synthase gene variations in women with idiopathic recurrent spontaneous abortion: genetic association study, systematic review and meta-analyse		Pereza, Nina; Peterlin, Borut; Volk, Marija; Kapović, Miljenko; Ostojić, Saša	
Altered LINE-1 methylation in Mothers of Children with Down Syndrome		Babić Božović, Ivana; Stanković, Aleksandra; Živković, Maja; Vraneković, Jadranka; Kapović, Miljenko; Brajenović-Milić, Bojana	
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The role of TPA I/D and PAI-1 4G/5G polymorphisms in multiple sclerosis.		Živković, Maja; Starčević Čizmarević, Nada; Lovrečić, Luca; Klupka-Sarić, Inge; Stanković, Aleksandra; Gašparović, Iva; Lavtar, Polona; Dinčić, Evica; Stojković, Ljiljana; Rudolf, Gorazd; Jazbec, Šega; Perković, Olivio; Sinanović, Osman; Sepčić, Juraj; Kapović, Miljenko; Peterlin, Borut; Ristić, Smiljana	
Genetic variation in tissue inhibitors of metalloproteinases as a risk factor for idiopathic recurrent spontaneous abortion		Pereza, Nina; Volk, Marija; Zrakić, Nikolina; Kapović, Miljenko; Peterlin, Borut; Ostojić, Saša	
Genetika neurodegenerativnih bolesti		Gašparović, Iva; Starčević-Čizmarević, Nada; Perković, Olivio; Antončić, Igor; Kapović, Miljenko; Ristić, Smiljana	
Insulin-like Growth Factor 2 and Insulin-like Growth Factor 2 Receptor Gene Polymorphisms in Idiopathic Male Infertility		Pereza, Nina; Ostojić, Saša; Kapović, Miljenko; Buretić-Tomljanović, Alena	
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Connection of Depression and Bone Loss in Perimenopausal and Postmenopausal Women		Ljubičić-Bistrović, Ivana; Rončević-Gržeta, Ika; Crnčević-Orlić, Željka; Frančišković, Tanja; Ljubičić, Rudolf; Orlić, Anamarija; Kapović, Miljenko	
Angiotensin-Converting Enzyme Insertion/Deletion Gene Polymorphism in Lung Cancer Patients		Dević Pavlić, Sanja; Ristić, Smiljana; Flego, Veljko; Kapović, Miljenko; Radojičić Badovinac, Anđelka	
Body Mass Index, Waist Circumference and Waist-to-Hip Ratio: Which Anthropometric Indicator is Better Predictor for the Hypertension Development in Women Population of the Island Cres		Kabalin, Milena; Kolarić, Branko; Vasiljev Marchesi, Vanja; Pereza, Nina; Ostojić, Saša; Rukavina, Tomislav; Kapović, Miljenko	
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Elevated Second-trimester Free beta-hCG as an Isolated Finding and Pregnancy Outcomes	Brajenović-Milić, Bojana; Tišlarić, Dubravka; Žuvić-Butorac, Marta; Bačić, Josip; Petrović, Oleg; Ristić, Smiljana; Mimica, Marko; Kapović, Miljenko	

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