

Svetlana Nikulina

List of Publications by Year in descending order

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54
papers

182
citations

1478505

6
h-index

1199594

12
g-index

60
all docs

60
docs citations

60
times ranked

334
citing authors

#	ARTICLE	IF	CITATIONS
1	May Measurement Month 2019: adherence to treatment and hypertension control in Russia. Russian Journal of Cardiology, 2020, 25, 3745.	1.4	14
2	Association of rs12204590 polymorphism with the development of stroke in hypertensive patients. Arterial Hypertension (Russian Federation), 2020, 25, 549-556.	0.4	0
3	Association of rs10507391 polymorphism with the development of acute cerebrovascular accident in patients with cardiovascular pathology. Bulletin of Siberian Medicine, 2020, 19, 85-93.	0.3	0
4	Relationship of matrix metalloproteinase-3 -11715A/6A polymorphism (rs35068180) and dilated cardiomyopathy. Russian Journal of Cardiology, 2020, 25, 3960.	1.4	0
5	Reply to the Letter - "The development of Brugada syndrome phenotype is multifactorial, combining genetic and environmental factors". European Review for Medical and Pharmacological Sciences, 2020, 24, 3446-3447.	0.7	0
6	Clinical application of chromosome 9p21.3 genotyping in patients with coronary artery disease. Experimental and Therapeutic Medicine, 2019, 18, 3100-3108.	1.8	2
7	Research Article Glu298Asp polymorphism in the β -NOS3 gene is not associated with susceptibility to chronic heart failure in a Russian population. Genetics and Molecular Research, 2019, 18, .	0.2	1
8	Hepcidin protects pulmonary artery hypertension in rats by activating NF- κ B/TNF- α pathway. European Review for Medical and Pharmacological Sciences, 2019, 23, 7573-7581.	0.7	6
9	Association of SOCS5 gene polymorphism with allergic bronchial asthma. Terapevticheskii Arkhiv, 2019, 91, 27-30.	0.8	1
10	Simultaneous Presentation of Ankylosing Spondylitis and Pancreatic Cancer: A Case Report. International Journal of Biomedicine, 2019, 9, 69-71.	0.2	0
11	Angiographic dynamics of coronary flow state after percutaneous coronary intervention in carriers of polymorphic RS1800470 variants of the TGF- β 1 gene. Russian Journal of Cardiology, 2019, , 77-82.	1.4	0
12	Association of rs556621 Polymorphism with Development of Stroke in Patients with Cardiovascular Pathology. Rational Pharmacotherapy in Cardiology, 2019, 15, 634-640.	0.8	0
13	Association of rs2230806 polymorphism with the development of acute cerebrovascular accident in patients with cardiovascular disease. Russian Journal of Cardiology, 2019, , 29-34.	1.4	0
14	Role of Single Nucleotide Polymorphism of β -Gene in the Development of Stroke. Rational Pharmacotherapy in Cardiology, 2018, 14, 488-493.	0.8	1
15	Differences in the Incidence of Major Risk Factors in Patients with Ischemic or Hemorrhagic Stroke in the Siberian Population. Rational Pharmacotherapy in Cardiology, 2018, 14, 509-514.	0.8	0
16	Prediction of cardiac conduction disorders using the methods of mathematical analysis. Russian Journal of Cardiology, 2018, , 53-58.	1.4	3
17	Polymorphisms of the SCN10A Gene in Patients With Sick Sinus Syndrome. Kardiologiya, 2018, 17, 53-59.	0.7	0
18	GIANT CELL MYOCARDITIS IN HIV INFECTION: A FATAL TANDEM. Russian Journal of Cardiology, 2018, , 129-133.	1.4	0

#	ARTICLE	IF	CITATIONS
19	Association of RS1800470 polymorphic variants of the transforming growth factor β 1 (TGF- β 1) gene with the severity of coronary atherosclerosis. Russian Journal of Cardiology, 2018, , 43-47.	1.4	5
20	Assessment of Association of rs2200733 SNP on Chromosome 4q25 with the Risk of the Development of Atrial Fibrillation in the Russian Population. International Journal of Biomedicine, 2018, 8, 280-283.	0.2	1
21	A Multiple Logistic Regression Model as an Additional Mathematical Method for Predicting the Development of Ischemic Stroke in Patients with Atrial Fibrillation. International Journal of Biomedicine, 2018, 8, 284-287.	0.2	0
22	POLYMORPHISMS OF ENDOTHELIAL NITRIC OXIDE SYNTHASE GENE AS PREDICTORS OF WOLFF-PARKINSON-WHITE SYNDROME. Rational Pharmacotherapy in Cardiology, 2017, 13, 597-601.	0.8	2
23	Gene-Targeted Analysis of Clinically Diagnosed Long QT Russian Families. International Heart Journal, 2017, 58, 81-87.	1.0	20
24	Genetic tests for low- and middle-income countries: a literature review. Genetics and Molecular Research, 2017, 16, .	0.2	12
25	TNFR1 -383 A \rightarrow C polymorphism and ankylosing spondylitis in a Russian Caucasian population: a preliminary study. Genetics and Molecular Research, 2017, 16, .	0.2	0
26	Research Article AluYb8 insertion in the WNK1 gene is not associated with hypertension in a Russian Caucasian population. Genetics and Molecular Research, 2017, 16, .	0.2	0
27	Organisation of the system of personal and professional development for medical academic staff: the problem and its solutions. Novosibirsk State Pedagogical University Bulletin, 2017, 7, 19-36.	0.2	0
28	GENETIC PREDICTORS OF ATRIAL FIBRILLATION. Rational Pharmacotherapy in Cardiology, 2016, 12, 331-336.	0.8	3
29	Genetic evaluation of AMPD1, CPT2, and PGYM metabolic enzymes in patients with chronic fatigue syndrome. Genetics and Molecular Research, 2016, 15, .	0.2	4
30	FIRST RUSSIA-BASED STUDY OF POLYMORPHISM rs2200733 CHROMOSOME 4q25 ASSOCIATION WITH DEVELOPMENT OF THE LONE ATRIAL FIBRILLATION. Russian Journal of Cardiology, 2016, , 28-31.	1.4	2
31	ORGANIZATIONAL ISSUES OF CARDIOREHABILITATION SERVICE IN RUSSIA. THE RESULTS OF PILOT PROJECT "DEVELOPMENT OF REHABILITATION SYSTEM OF CARDIOVASCULAR PATIENTS IN MEDICAL INSTITUTIONS OF RUSSIAN FEDERATION ENTITIES". Cardiovascular Therapy and Prevention (Russian Federation), 2016, 15, 4-12.	1.4	5
32	THE ROLE OF INTERLEUKIN-6 GENE IN DEVELOPMENT OF IDIOPATHIC SICK SINUS SYNDROME. Russian Journal of Cardiology, 2016, , 32-36.	1.4	1
33	Genetic polymorphisms in asthmatic patients living at Krasnoyarsk. Pulmonologiya, 2016, 26, 293-302.	0.8	1
34	An Investigation of the Association of the H558R Polymorphism of the SCN5A Gene with Idiopathic Cardiac Conduction Disorders. Genetic Testing and Molecular Biomarkers, 2015, 19, 288-294.	0.7	8
35	THE ROLE OF ROS1 GENE IN DEVELOPMENT OF STROKE. Russian Journal of Cardiology, 2015, , 46.	1.4	2
36	GENETIC PREDICTORS FOR CARDIOEMBOLIC STROKE IN PATIENTS WITH ATRIAL FIBRILLATION. Russian Journal of Cardiology, 2015, , 29.	1.4	1

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37	AN INHERITANCE OF IDIOPATHIC CARDIAC CONDUCTION DISORDERS. Russian Journal of Cardiology, 2015, , 82.	1.4	0
38	CLINICAL-GENETIC RISKOMETER FOR THE ISCHEMIC STROKE RISK ASSESSMENT IN ATRIAL FIBRILLATION. Russian Journal of Cardiology, 2015, , 42.	1.4	1
39	CLINICAL SIGNIFICANCE OF CORONARY ARTERY DISEASE GENETIC MARKERS: REALITY OR FAR FUTURE?. Russian Journal of Cardiology, 2015, , 7.	1.4	0
40	FORECASTING OF CCD IN FAMILIES BY MEANS OF A METHOD OF LOGISTIC REGRESSION IN KRASNOYARSK. Russian Journal of Cardiology, 2015, , 46.	1.4	0
41	A NOVEL GENETIC MARKER FOR INHERITED DISORDER OF THE HEART CONDUCTION SYSTEM. Russian Journal of Cardiology, 2015, , 30.	1.4	0
42	P577Relationship between rs1805124 polymorphism of SCN5A gene and primary left bundle branch block:. Cardiovascular Research, 2014, 103, S104.5-S104.	3.8	0
43	Relationship between polymorphism of SCN5A gene and primary cardiac conduction disorders. European Heart Journal, 2013, 34, P2309-P2309.	2.2	0
44	IMPACT OF LOCUS 9P21.3 SINGLE NUCLEOTIDE POLYMORPHISMS ON CORONARY ATHEROSCLEROSIS SEVERITY AND LONG-TERM OUTCOMES AFTER PERCUTANEOUS CORONARY INTERVENTION IN PATIENT WITH MYOCARDIAL INFARCTION. Rational Pharmacotherapy in Cardiology, 2013, 9, 241-246.	0.8	0
45	GLYCOPROTEIN INTEGRIN ALPHA POLYMORPHISM AND ACUTE CEREBROVASCULAR EVENTS IN THE FAMILIES OF PATIENTS WITH ATRIAL FIBRILLATION. Russian Journal of Cardiology, 2013, , 6-10.	1.4	0
46	GENETIC ASPECTS OF CONGENITAL LONG QT SYNDROME. Rational Pharmacotherapy in Cardiology, 2012, 8, 694-698.	0.8	1
47	GENETIC PREDICTORS OF IDIOPATHIC SICK SINUS SYNDROME. Rational Pharmacotherapy in Cardiology, 2012, 8, 804-809.	0.8	1
48	PREDICTIVE ROLE OF CONNEXIN 40 IN THE PATHOGENESIS OF HEREDITARY SICK SINUS SYNDROME. Rational Pharmacotherapy in Cardiology, 2011, 7, 174-176.	0.8	0
49	HEREDITARY INTRAVENTRICULAR CONDUCTION DISORDERS IN THE FAMILY FROM KRASNOYARSK. Rational Pharmacotherapy in Cardiology, 2011, 7, 601-604.	0.8	0
50	Genetics of Atrial Fibrillation. Circulation: Arrhythmia and Electrophysiology, 2010, 3, 291-299.	4.8	62
51	Association of ADRB1 Gene Polymorphism with Atrial Fibrillation. Genetic Testing and Molecular Biomarkers, 2010, 14, 249-253.	0.7	8
52	CLINICAL AND GENETIC PECULIARITIES OF ATRIAL FIBRILLATION. Rational Pharmacotherapy in Cardiology, 2008, 4, 13-18.	0.8	2
53	Apoptosis of leukocytes as a marker of neutrophil-endotheliocyte interaction in coronary heart disease. Bulletin of Experimental Biology and Medicine, 2007, 144, 39-41.	0.8	6
54	IDIOPATHIC SICK SINUS SYNDROME. Rational Pharmacotherapy in Cardiology, 2007, 3, 58-61.	0.8	0