

# Rita Khusainova

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6531998/publications.pdf>

Version: 2024-02-01

23  
papers

5,201  
citations

758635

12  
h-index

752256

20  
g-index

23  
all docs

23  
docs citations

23  
times ranked

9845  
citing authors

#	ARTICLE	IF	CITATIONS
1	The Simons Genome Diversity Project: 300 genomes from 142 diverse populations. <i>Nature</i> , 2016, 538, 201-206.	13.7	1,216
2	Ancient human genomes suggest three ancestral populations for present-day Europeans. <i>Nature</i> , 2014, 513, 409-413.	13.7	1,179
3	Genome-wide meta-analysis identifies 56 bone mineral density loci and reveals 14 loci associated with risk of fracture. <i>Nature Genetics</i> , 2012, 44, 491-501.	9.4	1,100
4	Genomic analyses inform on migration events during the peopling of Eurasia. <i>Nature</i> , 2016, 538, 238-242.	13.7	360
5	A recent bottleneck of Y chromosome diversity coincides with a global change in culture. <i>Genome Research</i> , 2015, 25, 459-466.	2.4	348
6	Global diversity, population stratification, and selection of human copy-number variation. <i>Science</i> , 2015, 349, aab3761.	6.0	293
7	The Caucasus as an Asymmetric Semipermeable Barrier to Ancient Human Migrations. <i>Molecular Biology and Evolution</i> , 2012, 29, 359-365.	3.5	161
8	The Genetic Legacy of the Expansion of Turkic-Speaking Nomads across Eurasia. <i>PLoS Genetics</i> , 2015, 11, e1005068.	1.5	149
9	A counter-clockwise northern route of the Y-chromosome haplogroup N from Southeast Asia towards Europe. <i>European Journal of Human Genetics</i> , 2007, 15, 204-211.	1.4	142
10	Human Y Chromosome Haplogroup N: A Non-trivial Time-Resolved Phylogeography that Cuts across Language Families. <i>American Journal of Human Genetics</i> , 2016, 99, 163-173.	2.6	98
11	A Mitochondrial Etiology of Neurodegenerative Diseases: Evidence from Parkinson's Disease. <i>Annals of the New York Academy of Sciences</i> , 2008, 1147, 1-20.	1.8	92
12	Between Lake Baikal and the Baltic Sea: genomic history of the gateway to Europe. <i>BMC Genetics</i> , 2017, 18, 110.	2.7	34
13	Y-chromosomal analysis of clan structure of Kalmyks, the only European Mongol people, and their relationship to Oirat-Mongols of Inner Asia. <i>European Journal of Human Genetics</i> , 2019, 27, 1466-1474.	1.4	8
14	Genetic Structure of Dagestan Populations: A Study of 11 Alu Insertion Polymorphisms. <i>Human Biology</i> , 2006, 78, 465-476.	0.4	5
15	Investigating the role of osteoprotegerin gene polymorphic variants in osteoporosis. <i>Russian Open Medical Journal</i> , 2021, 10, .	0.1	4
16	Modern classification and molecular-genetic aspects of osteogenesis imperfecta. <i>Vavilovskii Zhurnal Genetiki i Seleksii</i> , 2020, 24, 219-227.	0.4	4
17	Osteogenesis Imperfecta: Search for Mutations in Patients from the Republic of Bashkortostan (Russia). <i>Genes</i> , 2022, 13, 124.	1.0	4
18	The role of DNA methylation in the disorders of bone metabolism. <i>Vavilovskii Zhurnal Genetiki i Seleksii</i> , 2019, 23, 67-74.	0.4	2

#	ARTICLE	IF	CITATIONS
19	The role of VNTR aggrecan gene polymorphism in the development of osteoarthritis in women. Vavilovskii Zhurnal Genetiki I Seleksii, 2018, 22, 865-872.	0.4	1
20	The level of microelements and heterogeneity of joint hypermobility as an endophenotype of undifferentiated connective tissue dysplasia. Russian Open Medical Journal, 2020, 9, .	0.1	1
21	The role of polymorphic variants of matrix metalloproteinases genes in the formation of disruption of connective tissue homeostasis and pathology of the musculoskeletal system.. Klinicheskaia Meditsina, 2018, 96, 754-761.	0.2	0
22	Association of vascular endothelial growth factor B (VEGF $\beta$ ) gene polymorphisms with intracranial aneurysms. Vavilovskii Zhurnal Genetiki I Seleksii, 2019, 22, 992-999.	0.4	0
23	A clinical case of a hereditary form of colorectal cancer associated with a CHEK2 gene defect. Russian Journal of Oncology, 2021, 26, 29-34.	0.1	0