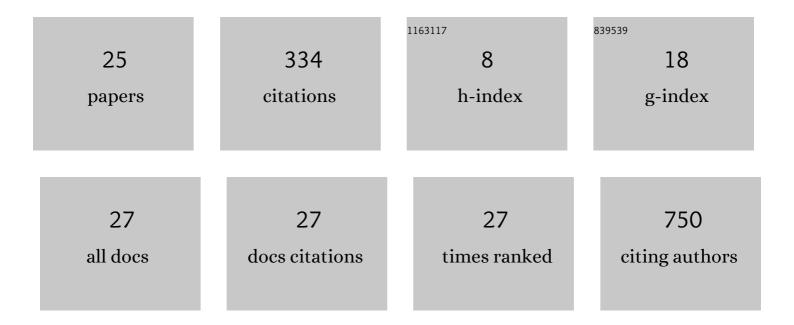
Nikolay Skryabin

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Analysis of mutations spectrum in the ATP7B gene in patients with Wilson disease using massively parallel sequencing. Klinichescheskaya Laboratornaya Diagnostika, 2022, 67, 250-256.	0.5	1
2	NLRP7 variants in spontaneous abortions with multilocus imprinting disturbances from women with recurrent pregnancy loss. Journal of Assisted Reproduction and Genetics, 2021, 38, 2893-2908.	2.5	4
3	Gene Expression Profiling Revealed 2 Types of Bronchial Basal Cell Hyperplasia and Squamous Metaplasia With Different Progression Potentials. Applied Immunohistochemistry and Molecular Morphology, 2020, 28, 477-483.	1.2	6
4	Delineation of Clinical Manifestations of the Inherited Xq24 Microdeletion Segregating with sXCI in Mothers: Two Novel Cases with Distinct Phenotypes Ranging from UBE2A Deficiency Syndrome to Recurrent Pregnancy Loss. Cytogenetic and Genome Research, 2020, 160, 245-254.	1.1	8
5	Runs of homozygosity in spontaneous abortions from families with recurrent pregnancy loss. Vavilovskii Zhurnal Genetiki I Selektsii, 2019, 23, 244-249.	1.1	Ο
6	Allele-Specific Biased Expression of the CNTN6 Gene in iPS Cell-Derived Neurons from a Patient with Intellectual Disability and 3p26.3 Microduplication Involving the CNTN6 Gene. Molecular Neurobiology, 2018, 55, 6533-6546.	4.0	18
7	Compound phenotype in a girl with r(22), concomitant microdeletion 22q13.32-q13.33 and mosaic monosomy 22. Molecular Cytogenetics, 2018, 11, 26.	0.9	15
8	A mosaic intragenic microduplication of <i>LAMA1</i> and a constitutional 18p11.32 microduplication in a patient with <i>keratosis pilaris</i> and intellectual disability. American Journal of Medical Genetics, Part A, 2018, 176, 2395-2403.	1.2	9
9	Karyotype of the blastocoel fluid demonstrates low concordance with both trophectoderm and inner cell mass. Fertility and Sterility, 2018, 109, 1127-1134.e1.	1.0	38
10	Genomic structural variations for cardiovascular and metabolic comorbidity. Scientific Reports, 2017, 7, 41268.	3.3	29
11	Epigenetic status of imprinted genes in placenta during recurrent pregnancy loss. Russian Journal of Genetics, 2017, 53, 376-387.	0.6	8
12	Epigenetic silencing of genomic structural variations. Russian Journal of Genetics, 2017, 53, 1072-1079.	0.6	3
13	Clinically relevant morphological structures in breast cancer represent transcriptionally distinct tumor cell populations with varied degrees of epithelial-mesenchymal transition and CD44+CD24-stemness. Oncotarget, 2017, 8, 61163-61180.	1.8	22
14	A de novo microtriplication at 4q21.21â€q21.22 in a patient with a vascular malignant hemangioma, elongated sigmoid colon, developmental delay, and absence of speech. American Journal of Medical Genetics, Part A, 2016, 170, 2089-2096.	1.2	7
15	Preimplantation genetic diagnosis by blastocentesis: Problems and perspectives. Russian Journal of Genetics, 2016, 52, 1-7.	0.6	5
16	Molecular karyotyping of cell-free DNA from blastocoele fluid as a basis for noninvasive preimplantation genetic screening of aneuploidy. Russian Journal of Genetics, 2015, 51, 1123-1128.	0.6	4
17	Relationship between morphological and cytogenetic heterogeneity in invasive micropapillary carcinoma of the breast: a report of one case. Journal of Clinical Pathology, 2015, 68, 758-762.	2.0	8
18	Single gene microdeletions and microduplication of 3p26.3 in three unrelated families: CNTN6 as a new candidate gene for intellectual disability. Molecular Cytogenetics, 2014, 7, 97.	0.9	51

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#	Article	IF	CITATIONS
19	Array CGH analysis of a cohort of Russian patients with intellectual disability. Gene, 2014, 536, 145-150.	2.2	40
20	Somatic genome variations in vascular tissues and peripheral blood leukocytes in patients with atherosclerosis. Russian Journal of Genetics, 2014, 50, 870-878.	0.6	6
21	Epigenetic effects of trisomy 16 in human placenta. Molecular Biology, 2013, 47, 373-381.	1.3	8
22	Dynamics of aberrant methylation of functional groups of genes in progression of breast cancer. Molecular Biology, 2013, 47, 267-274.	1.3	5
23	Phenotypic Drift as a Cause for Intratumoral Morphological Heterogeneity of Invasive Ductal Breast Carcinoma Not Otherwise Specified. BioResearch Open Access, 2013, 2, 148-154.	2.6	26
24	Multilocus epimutations of imprintome in the pathology of human embryo development. Molecular Biology, 2012, 46, 183-191.	1.3	4
25	DNA methylation profile in human placental tissues. Molecular Biology, 2011, 45, 493-499.	1.3	6