# Nikolay Skryabin

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3588444/publications.pdf

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| 25 | 334 | 1163117 8 | 839539 18 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 27 | 27 | 27 | 750 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Analysis of mutations spectrum in the ATP7B gene in patients with Wilson disease using massively parallel sequencing. Klinicheskeskaya 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 | 0 |
| 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 LAMA1 and a constitutional 18p11.32 microduplication in a patient with keratosis pilaris 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 |

| # | 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 |