Evgeny N Suspitsin

List of Publications by Year
in descending order

60	1,128	489802 18	466096 32
papers	citations	h-index	g-index

60	60	60	2057
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Identification of recurrent pathogenic alleles using exome sequencing data: Proof-of-concept study of Russian subjects. European Journal of Medical Genetics, 2022, 65, 104426.	0.7	0
2	Revisiting multiple erroneous genetic testing results and clinical misinterpretations in a patient with Li-Fraumeni syndrome: lessons for translational medicine. Hereditary Cancer in Clinical Practice, 2021, 19, 2.	0.6	1
3	ACTIVATED PHOSPHOINOSITIDE 3-KINASE Î´ SYNDROME â€“ A RARE FORM OF PRIMARY IMMUNODEFICIENCY. Pediatriia, 2021, 100, 248-253.	0.1	0
4	Systemic Lupus Erythematosus with neuropsychiatric manifestations in a child: case report and review of the international recommendations for the diagnostics and management. Rossiyskiy Vestnik Perinatologii I Pediatrii, 2021, 66, 98-105.	0.1	3
5	HEREDITARY ANGIOEDEMA â€“ A PROBLEM AT THE INTERSECTION OF IMMUNOLOGY AND ALLERGOLOGY: ANALYSIS OF LITERATURE DATA AND DESCRIPTION OF A SERIES OF 34 CASES. Pediatriia, 2021, 100, 49-57.	0.1	1
6	MEVALONATE KINASE DEFICIENCY IN AN INFANT: KEY ASPECTS OF DIAGNOSIS AND TREATMENT. Pediatriia, 2021, 100, 276-283.	0.1	1
7	ANALYSIS OF INTERFERON-I SIGNALING ACTIVITY IN CHILDREN WITH SYSTEMIC LUPUS ERYTHEMATOSUS: RESULTS OF A PILOT PROSPECTIVE STUDY. Pediatriia, 2021, 100, 77-88.	0.1	0
8	Analysis of interferon type I signature for differential diagnosis of diseases of the immune system () Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50	0.2	1
9	Genetic variants identified in children with recurrent infections. Detskie Infekcii (Moskva), 2021, 20, 13-17.	0.1	0
10	ATM mutation spectrum in Russian children with ataxia-telangiectasia. European Journal of Medical Genetics, 2020, 63, 103630.	0.7	13
11	The spectrum of Lynch syndrome-associated germ-line mutations in Russia. European Journal of Medical Genetics, 2020, 63, 103753.	0.7	9
12	Exome sequencing study of Russian breast cancer patients suggests a predisposing role for USP39. Breast Cancer Research and Treatment, 2020, 179, 731-742.	1.1	9
13	Gene rearrangements in consecutive series of pediatric inflammatory myofibroblastic tumors. Pediatric Blood and Cancer, 2020, 67, e28220.	0.8	24
14	Next generation sequencing analysis of consecutive Russian patients with clinical suspicion of inborn errors of immunity. Clinical Genetics, 2020, 98, 231-239.	1.0	10
15	PREDICTIVE RESPONSE MARKERS FOR IMMUNE RESPONSE BLOCKS. Siberian Journal of Oncology, 2020, 19, 123-131.	0.1	4
16	RECURRENT STREPTOCOCCAL INFECTION IN A CHILD WITH A CONGENITAL DEFICIENCY OF MANNOSE-BINDING LECTIN. Pediatriia, 2020, 99, 266-270.	0.1	0
17	Exome-based search for recurrent disease-causing alleles in Russian population. European Journal of Medical Genetics, 2019, 62, 103656.	0.7	7
18	GP134â€…Confirmation of pathogenetic heterogeneity of diabetes mellitus in children using whole-exome sequencing. , 2019, , .		0

#	Article	IF	Citations
19	Wholeâ€'exome sequencing in Russian children with nonâ€'type 1 diabetes mellitus reveals a wide spectrum of genetic variants in MODYâ€'related and unrelated genes. Molecular Medicine Reports, 2019, 20, 4905-4914.	1.1	18
20	Primary tumor (tumoral) calcification is a rare disease in the practice of a rheumatologist and orthopedist: experience with the use of an interleukin-1 inhibitor in combination with surgical correction. Pediatric Traumatology, Orthopaedics and Reconstructive Surgery, 2019, 7, 85-92.	0.1	1
21	Pattern of TSC1 and TSC2 germline mutations in Russian patients with tuberous sclerosis. Journal of Human Genetics, 2018, 63, 597-604.	1.1	7
22	Multigene sequencing reveals heterogeneity of NLRP12-related autoinflammatory disorders. Rheumatology International, 2018, 38, 887-893.	1.5	42
23	Spectrum of APC and MUTYH germâ€line mutations in Russian patients with colorectal malignancies. Clinical Genetics, 2018, 93, 1015-1021.	1.0	16
24	Diagnosis of carcinoma of unknown primary site with the aid of simple PCR tests: a single-center experience. Neoplasma, 2018, 65, 461-468.	0.7	2
25	Case report of the inflammatory myofibroblastic tumor of the liver in infant. Pediatric Hematology/Oncology and Immunopathology, 2018, 17, 93-102.	0.1	0
26	First Two Cases of Bloom Syndrome in Russia: Lack of Skin Manifestations in a BLM c.1642C>T (p.Q548X) Homozygote as a Likely Cause of Underdiagnosis. Molecular Syndromology, 2017, 8, 103-106.	0.3	6
27	Detection of BRCA1 gross rearrangements by droplet digital PCR. Breast Cancer Research and Treatment, 2017, 165, 765-770.	1.1	16
28	Bardet-Biedl Syndrome. Molecular Syndromology, 2016, 7, 62-71.	0.3	103
29	Evidence for a pathogenic role of BRCA1 L1705P and W1837X germ-line mutations. Molecular Biology Reports, 2016, 43, 335-338.	1.0	1
30	Exome Sequencing of a Family with Bardet-Biedl Syndrome Identifies the Common Russian Mutation c.1967_1968delTAinsC in <i>BBS7</i>. Molecular Syndromology, 2015, 6, 96-98.	0.3	8
31	Identification of novel hereditary cancer genes by whole exome sequencing. Cancer Letters, 2015, 369, 274-288.	3.2	31
32	High response rates to neoadjuvant platinum-based therapy in ovarian cancer patients carrying germ-line BRCA mutation. Cancer Letters, 2015, 369, 363-367.	3.2	82
33	Development of breast tumors in CHEK2, NBN/NBS1 and BLM mutation carriers does not commonly involve somatic inactivation of the wild-type allele. Medical Oncology, 2014, 31, 828.	1.2	20
34	High prevalence ofGPRC5Agermline mutations inBRCA1-mutant breast cancer patients. International Journal of Cancer, 2014, 134, 2352-2358.	2.3	31
35	Evidence for angiogenesis-independent contribution of VEGFR1 (FLT1) in gastric cancer recurrence. Medical Oncology, 2013, 30, 644.	1.2	4
36	Pattern of clinically relevant mutations in consecutive series of Russian colorectal cancer patients. Medical Oncology, 2013, 30, 686.	1.2	43

#	Article	IF	Citations
37	High prevalence and breast cancer predisposing role of the BLM c.1642 C>T (Q548X) mutation in Russia. International Journal of Cancer, 2012, 130, 2867-2873.	2.3	58
38	Abstract 2597: Identification of new genes for hereditary breast cancer. , 2012, , .		0
39	Paired distribution of molecular subtypes in bilateral breast carcinomas. Cancer Genetics, 2011, 204, 96-102.	0.2	14
40	Mixed epithelial/mesenchymal metaplastic carcinoma (carcinosarcoma) of the breast in BRCA1 carrier. Breast Cancer, 2011, 18, 137-140.	1.3	18
41	Rapid Symptomatic Improvement in Gefitinib-Treated Patients with EGFR-Mutated Lung Cancer: Possible Role of Downregulation of Inflammatory Molecules?. Onkologie, 2011, 34, 559-560.	1.1	3
42	High sensitivity of BRCA1-associated tumors to cisplatin monotherapy: report of two cases. Cancer Genetics and Cytogenetics, 2010, 197, 91-94.	1.0	18
43	Non-founder BRCA1 mutations in Russian breast cancer patients. Cancer Letters, 2010, 298, 258-263.	3.2	34
44	High frequency of BRCA1, but not CHEK2 or NBS1 (NBN), founder mutations in Russian ovarian cancer patients. Hereditary Cancer in Clinical Practice, 2009, 7, 5.	0.6	31
45	Large family with both parents affected by distinct BRCA1 mutations: implications for genetic testing. Hereditary Cancer in Clinical Practice, 2009, 7, 2.	0.6	3
46	TP53 mutations in synchronous and metachronous bilateral breast carcinomas. Cancer Genetics and Cytogenetics, 2008, 184, 119-121.	1.0	6
47	CHEK2 1100 delC mutation in Russian ovarian cancer patients. Hereditary Cancer in Clinical Practice, 2007, 5, 153.	0.6	7
48	Nonrandom distribution of oncogene amplifications in bilateral breast carcinomas: Possible role of host factors and survival bias. International Journal of Cancer, 2007, 120, 297-302.	2.3	10
49	High frequency of BRCA1 5382insC mutation in Russian breast cancer patients. European Journal of Cancer, 2006, 42, 1380-1384.	1.3	70
50	Microsatellite instability analysis of bilateral breast tumors suggests treatment-related origin of some contralateral malignancies. Journal of Cancer Research and Clinical Oncology, 2006, 133, 57-64.	1.2	18
51	NBS1 657del5 mutation may contribute only to a limited fraction of breast cancer cases in Russia. International Journal of Cancer, 2005, 114, 585-589.	2.3	59
52	Apoptosis-deficient Pro allele of gene is associated with the resistance of psoriasis to the UV-based therapy. Journal of Dermatological Science, 2005, 37, 185-187.	1.0	7
53	Evidence against involvement of p53 polymorphism in breast cancer predisposition. International Journal of Cancer, 2003, 103, 431-433.	2.3	65
54	Distinct prevalence of the CYP19 Î”3(TTTA)7 allele in premenopausal versus postmenopausal breast cancer patients, but not in control individuals. European Journal of Cancer, 2002, 38, 1911-1916.	1.3	19

#	Article	IF	Citations
55	Concordance of allelic imbalance profiles in synchronous and metachronous bilateral breast carcinomas. International Journal of Cancer, 2002, 100, 557-564.	2.3	64
56	Improved Reliability of Allele-Specific PCR. BioTechniques, 2002, 33, 484-490.	0.8	4
57	CYP19 gene polymorphism in endometrial cancer patients. Journal of Cancer Research and Clinical Oncology, 2001, 127, 135-138.	1.2	51
58	L-myc polymorphism in cancer patients, healthy blood donors and elderly, tumor-free individuals in Russia. , 2000, 85, 747-750.		7
59	CYP17 polymorphism in the groups of distinct breast cancer susceptibility: comparison of patients with the bilateral disease vs. monolateral breast cancer patients vs. middle-aged female controls vs. elderly tumor-free women. Cancer Letters, 2000, 156, 45-50.	3.2	28
60	Evidence for microsatellite instability in bilateral breast carcinomas. Cancer Letters, 2000, 154, 9-17.	3.2	20