

Firoz Ahmad

List of Publications by Year in descending order

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28
papers

283
citations

932766

10
h-index

996533

15
g-index

28
all docs

28
docs citations

28
times ranked

473
citing authors

#	ARTICLE	IF	CITATIONS
1	A novel coexistence of tetrasomy 8 and FLT3-ITD along with variant 3 way translocation t(4;17;15) in acute promyelocytic leukemia: Case study and literature review. <i>Cancer Genetics</i> , 2022, 262-263, 111-117.	0.2	0
2	Aberrant epigenetic inactivation of RASSF1A and MGMT gene and genetic mutations of KRAS, cKIT and BRAF in Indian testicular germ cell tumours. <i>Cancer Genetics</i> , 2020, 241, 42-50.	0.2	7
3	Molecular evaluation of BRAF V600 mutation and its association with clinicopathological characteristics: First findings from Indian malignant melanoma patients. <i>Cancer Genetics</i> , 2019, 231-232, 46-53.	0.2	6
4	Molecular evaluation of BRAF gene mutation in thyroid tumors: Significant association with papillary tumors and extra thyroidal extension indicating its role as a biomarker of aggressive disease. <i>Experimental and Molecular Pathology</i> , 2018, 105, 380-386.	0.9	10
5	Molecular profiling of gene copy number abnormalities in key regulatory genes in high-risk B-lineage acute lymphoblastic leukemia: frequency and their association with clinicopathological findings in Indian patients. <i>Medical Oncology</i> , 2017, 34, 92.	1.2	5
6	Somatic mutation analysis of KRAS, BRAF, HER2 and PTEN in EGFR mutation-negative non-small cell lung carcinoma: determination of frequency, distribution pattern and identification of novel deletion in HER2 gene from Indian patients. <i>Medical Oncology</i> , 2016, 33, 117.	1.2	13
7	Molecular evaluation of PIK3CA gene mutation in breast cancer: determination of frequency, distribution pattern and its association with clinicopathological findings in Indian patients. <i>Medical Oncology</i> , 2016, 33, 74.	1.2	17
8	Association of Genetic Variants in ARID5B, IKZF1 and CEBPE with Risk of Childhood de novo B-Lineage Acute Lymphoblastic Leukemia in India. <i>Asian Pacific Journal of Cancer Prevention</i> , 2016, 17, 3989-95.	0.5	16
9	Molecular Spectrum of Somatic EGFR and KRAS Gene Mutations in non Small Cell Lung Carcinoma: Determination of Frequency, Distribution Pattern and Identification of Novel Variations in Indian Patients. <i>Pathology and Oncology Research</i> , 2015, 21, 675-687.	0.9	11
10	Molecular spectrum of c-KIT and PDGFRA gene mutations in gastro intestinal stromal tumor: determination of frequency, distribution pattern and identification of novel mutations in Indian patients. <i>Medical Oncology</i> , 2015, 32, 424.	1.2	12
11	Cytogenetic Profile of De Novo B lineage Acute Lymphoblastic Leukemia: Determination of Frequency, Distribution Pattern and Identification of Rare and Novel Chromosomal Aberrations in Indian Patients. <i>Asian Pacific Journal of Cancer Prevention</i> , 2015, 16, 7219-7229.	0.5	4
12	Molecular characterization of complex chromosomal rearrangement: First report of novel t(7;12)(q11;q22) as part of a complex karyotype in de novo AML-M2 case. <i>Pathology Research and Practice</i> , 2014, 210, 1090-1094.	1.0	1
13	Molecular spectrum of KRAS, BRAF, and PIK3CA gene mutation: determination of frequency, distribution pattern in Indian colorectal carcinoma. <i>Medical Oncology</i> , 2014, 31, 124.	1.2	25
14	Molecular Evaluation of DNMT3A and IDH1/2 Gene Mutation: Frequency, Distribution Pattern and Associations with Additional Molecular Markers in Normal Karyotype Indian Acute Myeloid Leukemia Patients. <i>Asian Pacific Journal of Cancer Prevention</i> , 2014, 15, 1247-1253.	0.5	30
15	Molecular Cytogenetic Investigations In A Novel Chromosomal Abnormality Of T(10;15)(q22;q22) In A Pediatric Precursor-B-Acute Lymphoblastic Leukemia Patient. <i>Journal of Hematology and Oncology Research</i> , 2014, 1, 28-33.	1.8	1
16	Molecular Investigation of Isocitrate Dehydrogenase Gene (IDH) Mutations in Gliomas: First Report of IDH2 Mutations in Indian Patients. <i>Asian Pacific Journal of Cancer Prevention</i> , 2013, 14, 7261-7264.	0.5	16
17	Molecular Evaluation of CEBPA Gene Mutation in Normal Karyotype Acute Myeloid Leukemia: A Comparison of Two Methods and Report of Novel CEBPA Mutations from Indian Acute Myeloid Leukemia Patients. <i>Genetic Testing and Molecular Biomarkers</i> , 2012, 16, 707-715.	0.3	6
18	Molecular analysis of WT1 and KIT mutations in patients from an Indian population with de novo acute myeloid leukemia: determination of incidence, distribution patterns, and report of a novel KIT mutation. <i>Leukemia and Lymphoma</i> , 2011, 52, 865-876.	0.6	8

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19	Cytogenetic and molecular characterization of a hepatosplenic T-cell lymphoma: report of a novel chromosomal aberration. <i>Cancer Genetics</i> , 2011, 204, 103-107.	0.2	6
20	Analysis of FLT3-ITD and FLT3-Asp835 Mutations in de novo Acute Myeloid Leukemia: Evaluation of Incidence, Distribution Pattern, Correlation with Cytogenetics and Characterization of Internal Tandem Duplication from Indian Population. <i>Cancer Investigation</i> , 2010, 28, 63-73.	0.6	21
21	Novel t(8;17)(q23;q24.2) and t(9;22)(p24.1;q12.2) in acute megakaryoblastic leukemia AML-M7 subtype in an adult patient. <i>Cancer Genetics and Cytogenetics</i> , 2009, 193, 112-115.	1.0	4
22	Mutations of <i>NPM1</i> gene in de novo acute myeloid leukaemia: determination of incidence, distribution pattern and identification of two novel mutations in Indian population. <i>Hematological Oncology</i> , 2009, 27, 90-97.	0.8	27
23	Molecular cytogenetic findings in a three-way novel variant of t(1;8;21)(p35;q22;q22): a unique relocation of the AML1/ETO fusion gene 1p35 in AML-M2. <i>Cancer Genetics and Cytogenetics</i> , 2008, 180, 153-157.	1.0	17
24	Molecular cytogenetic investigations in a novel complex variant of t(8;21)(q22;q22) with ins(15;21)(q15;q22.2q22.3) in a patient with AML-M2 subtype. <i>Cancer Genetics and Cytogenetics</i> , 2008, 184, 52-56.	1.0	3
25	Specific chromosomal aberrations in de novo acute myeloid leukemia: A comparative analysis of results with a report of three novel chromosomal rearrangements t(7;14)(q35;q13), t(8;18)(p11.2;q12), t(13;15) in Indian population. <i>Cancer Detection and Prevention</i> , 2008, 32, 168-177.	2.1	5
26	Cytogenetic profile of acute lymphocytic leukemia patients: report of a novel translocation t(4;13)(q21A-3; q35) from an Indian population. <i>Hematology</i> , 2008, 13, 28-33.	0.7	2
27	Acute Myelogenous Leukemia (M0/M1) with novel chromosomal abnormality of t(14;17)(q32; q11.2). <i>American Journal of Hematology</i> , 2007, 82, 676-678.	2.0	5
28	Cytogenetic investigation in chronic myeloid leukemia: study from an Indian population. <i>Asian Pacific Journal of Cancer Prevention</i> , 2006, 7, 423-6.	0.5	5