# Angelo Lonoce

List of Publications by Year
in descending order



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|---|---|---|---|
| 21 papers | 936 citations | 840776 11 h-index | 752698 20 g-index |
| 22 all docs | 22 docs citations | 22 times ranked | 1682 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Identification and molecular characterization of recurrent genomic deletions on 7p12 in the IKZF1 gene in a large cohort of BCR-ABL1â€“positive acute lymphoblastic leukemia patients: on behalf of Gruppo Italiano Malattie Ematologiche dell'Adulto Acute Leukemia Working Party (GIMEMA AL WP). Blood, 2009, 114, 2159-2167. | 1.4 | 201 |
| 2 | Gene amplification as double minutes or homogeneously staining regions in solid tumors: Origin and structure. Genome Research, 2010, 20, 1198-1206. | 5.5 | 194 |
| 3 | MYC-containing double minutes in hematologic malignancies: evidence in favor of the episome model and exclusion of MYC as the target gene. Human Molecular Genetics, 2006, 15, 933-942. | 2.9 | 116 |
| 4 | Comparative mapping of human alphoid sequences in great apes using fluorescence in situ hybridization. Genomics, 1995, 25, 477-484. | 2.9 | 110 |
| 5 | Genomic organization and evolution of double minutes/homogeneously staining regions withMYCamplification in human cancer. Nucleic Acids Research, 2014, 42, 9131-9145. | 14.5 | 91 |
| 6 | MYC-containing amplicons in acute myeloid leukemia: genomic structures, evolution, and transcriptional consequences. Leukemia, 2018, 32, 2152-2166. | 7.2 | 70 |
| 7 | Structural Organization of Multiple Alphoid Subsets Coexisting on Human Chromosomes 1, 4, 5, 7, 9, 15, 18, and 19. Genomics, 1996, 38, 325-330. | 2.9 | 45 |
| 8 | Familial adenomatous polyposis: Identification of a new frameshift mutation of the APC gene in an Italian family. Biochemical and Biophysical Research Communications, 1992, 184, 1357-1363. | 2.1 | 24 |
| 9 | Molecular cytogenetic characterization of a complex rearrangement involving chromosomes 9 and 22 in a case of Ph-negative chronic myeloid leukemia. Cancer Genetics and Cytogenetics, 2002, 136, 141-145. | 1.0 | 12 |
| 10 | t(15;21) translocations leading to the concurrent downregulation of RUNX1 and its transcription factor partner genes SIN3A and TCF12 in myeloid disorders. Molecular Cancer, 2015, 14, 211. | 19.2 | 12 |
| 11 | Two alternatively spliced 5â€²BCR/3â€²JAK2 fusion transcripts in a myeloproliferative neoplasm with a three-way t(9;18;22)(p23;p11.3;q11.2) translocation. Cancer Genetics, 2011, 204, 512-515. | 0.4 | 11 |
| 12 | Characterization of a hotspot region on chromosome 12 for amplification in ring chromosomes in atypical lipomatous tumors. Genes Chromosomes and Cancer, 2009, 48, 993-1001. | 2.8 | 10 |
| 13 | Epigenetically induced ectopic expression of UNCX impairs the proliferation and differentiation of myeloid cells. Haematologica, 2017, 102, 1204-1214. | 3.5 | 8 |
| 14 | A rare but recurrent t(8;13)(q24;q14) translocation in Bâ€cell chronic lymphocytic leukaemia causing <scp>MYC</scp> upâ€regulation and concomitant loss of <scp>PVT</scp>1, miRâ€15/16 and <scp>DLEU</scp>7. British Journal of Haematology, 2016, 172, 296-299. | 2.5 | 7 |
| 15 | Extramedullary molecular evidence of the 5â€²KIAA1509/3â€²PDGFRB fusion gene in chronic eosinophilic leukemia. Leukemia Research, 2008, 32, 347-351. | 0.8 | 5 |
| 16 | Linkage studies in Italian families with familial adenomatous polyposis. Human Genetics, 1993, 90, 545-550. | 3.8 | 4 |
| 17 | 1q23.1 homozygous deletion and downregulation of Fc receptor-like family genes confer poor prognosis in chronic lymphocytic leukemia. Clinical and Experimental Medicine, 2019, 19, 261-267. | 3.6 | 4 |
| 18 | Bone marrow ectopic expression of a non-coding RNA in childhood T-cell acute lymphoblastic leukemia with a novel t(2;11)(q11.2;p15.1) translocation. Molecular Cancer, 2008, 7, 80. | 19.2 | 3 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | MYC-containing amplicons in acute myeloid leukemia: Genomic structures, evolution, and transcriptional consequences. Leukemia, 2017, , . | 7.2 | 2 |
| 20 | RALE051: a novel established cell line of sporadic Burkitt lymphoma. Leukemia and Lymphoma, 2018, 59, 1252-1255. | 1.3 | 0 |
| 21 | A New Entity of Acute Myeloid Leukemia Driven By Epigenetic and Somatic Dis-Regulation of Uncx, a Novel Homeobox Transcription Factor Gene. Blood, 2015, 126, 1356-1356. | 1.4 | 0 |