ValÃ©rie Soenen

List of Publications by Year in descending order

31 papers	1,602 citations	516710 16 h-index	501196 28 g-index
32 all docs	32 docs citations	32 times ranked	1982 citing authors

#	Article	IF	Citations
1	A new approach for diagnosing chronic myelomonocytic leukemia using structural parameters of Sysmex XN^{TM} analyzers in routine laboratory practice. Scandinavian Journal of Clinical and Laboratory Investigation, 2018, 78, 159-164.	1.2	14
2	BACH2 promotes indolent clinical presentation in WaldenstrÃ¶m macroglobulinemia. Oncotarget, 2017, 8, 57451-57459.	1.8	2
3	Genome wide SNP array identified multiple mechanisms of genetic changes in Waldenstrom macroglobulinemia. American Journal of Hematology, 2013, 88, 948-954.	4.1	45
4	MYD88 L265P mutation in Waldenstrom macroglobulinemia. Blood, 2013, 121, 4504-4511.	1.4	214
5	The B7-H3 Protein In Acute Myeloid Leukemia. Blood, 2013, 122, 2620-2620.	1.4	1
6	Construction of a tube-shaped tracheal substitute using fascial flap-wrapped revascularized allogenic aorta. European Journal of Cardio-thoracic Surgery, 2012, 41, 663-668.	1.4	16
7	Genome Wide SNP Array (SNPa) Analysis Reveals Clonal Evolution During Clinical Course in Waldenstrom's Macroglobulinemia (WM). Blood, 2012, 120, 297-297.	1.4	2
8	B-Cell-Specific Transcription Factor BACH2 Involved in the Clinical Behavior Heterogeneity of Waldenstrol̂m Macroglobulinemia. Blood, 2012, 120, 1288-1288.	1.4	0
9	High-Throughput Genomic Analysis in WaldenstrÃ¶m's Macroglobulinemia. Clinical Lymphoma, Myeloma and Leukemia, 2011, 11, 106-108.	0.4	19
10	Chronic myeloproliferative disorder with t(8;22)(p11;q11) can mime clonal cytogenetic evolution of authentic chronic myelogeneous leukemia. Genes Chromosomes and Cancer, 2008, 47, 915-918.	2.8	25
11	JAK2V617F-positive polycythemia vera and Philadelphia chromosome-positive chronic myeloid leukemia: one patient with two distinct myeloproliferative disorders. Leukemia, 2008, 22, 1454-1455.	7.2	45
12	Mechanisms of genesis of variant translocation in chronic myeloid leukemia are not correlated with ABL1 or BCR deletion status or response to imatinib therapy. Cancer Genetics and Cytogenetics, 2008, 182, 95-102.	1.0	46
13	Isolated tetrasomy 13: a fifth case report of a rare chromosome abnormality associated with poorly differentiated acute myeloid leukemia. Cancer Genetics and Cytogenetics, 2006, 168, 181-182.	1.0	2
14	Decellularized heart valve as a scaffold for in vivo recellularization: Deleterious effects of granulocyte colony-stimulating factor. Journal of Thoracic and Cardiovascular Surgery, 2006, 131, 843-852.	0.8	44
15	Molecular characterization of the idiopathic hypereosinophilic syndrome (HES) in 35 French patients with normal conventional cytogenetics. Leukemia, 2005, 19, 792-798.	7.2	108
16	Mesenchymal cells generated from patients with myelodysplastic syndromes are devoid of chromosomal clonal markers and support short- and long-term hematopoiesis in vitro. Oncogene, 2005, 24, 2441-2448.	5.9	71
17	Role of multiplex FISH in identifying chromosome involvement in myelodysplastic syndromes and acute myeloid leukemias with complex karyotypes: a report on 28 cases. Cancer Genetics and Cytogenetics, 2005, 157, 118-126.	1.0	30
18	Familial occurrence of thymoma and autoimmune diseases with the constitutional translocation t(14;20)(q24.1;p12.3). Genes Chromosomes and Cancer, 2005, 44, 154-160.	2.8	19

#	ARTICLE	IF	CITATIONS
19	A case of refractory anemia with 17pâ" syndrome following azathioprine treatment for heart transplantation. Leukemia, 2004, 18, 878-878.	7.2	8
20	Chromosomal insertion involving MLL in childhood acute myeloblastic leukemia (M4). Cancer Genetics and Cytogenetics, 2004, 150, 153-155.	1.0	3
21	Efficient generation of antileukemic autologous T cells by short-term culture and ?-irradiation of myeloid leukemic cells. Cancer Immunology, Immunotherapy, 2004, 53, 793-8.	4.2	8
22	Acute myeloblastic leukemia (AML) with inv (16)(p13;q22) and the rare I type CBFÎ²-MYH11 transcript: report of two new cases. Leukemia, 2002, 16, 150-151.	7.2	15
23	Several types of mutations of the Abl gene can be found in chronic myeloid leukemia patients resistant to STI571, and they can pre-exist to the onset of treatment. Blood, 2002, 100, 1014-1018.	1.4	502
24	Prognostic significance of p16INK4a immunocytochemistry in adult ALL with standard risk karyotype. Leukemia, 2001, 15, 1054-1059.	7.2	8
25	MOZ is fused top300 in an acute monocytic leukemia with t(8;22). , 2000, 28, 138-144.		157
26	FISH analysis with a YAC probe improves detection of LAZ3/BCL6 rearrangement in non-Hodgkin's lymphoma. The Hematology Journal, 2000, 1, 117-125.	1.4	14
27	Myelodysplasia during the course of myeloma. Restriction of 17p deletion and p53 overexpression to myeloid cells. Leukemia, 1998, 12, 238-241.	7.2	5
28	In SituElectrical Characterization of Magnesium Vanadate Reference Phases (meta-MgV2O6,) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 387 Journal of Catalysis, 1996, 159, 410-417.	6.2	58
29	Identification of a YAC spanning the translocation breakpoint t(8;22) associated with acute monocytic leukemia. , 1996, 15, 191-194.		20
30	Monoclonal gammopathy of undetermined significance: chromosome changes are a common finding within bone marrow plasma cells. British Journal of Haematology, 1995, 90, 693-696.	2.5	62
31	Combined immunophenotyping and in situ hybridization (FICTION): a rapid method to study cell lineage involvement in myelodysplastic syndromes. British Journal of Haematology, 1995, 90, 701-706.	2.5	39