

Emanuela Boveri

List of Publications by Year in descending order

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Version: 2024-02-01

80
papers

6,300
citations

94269

37
h-index

74018

75
g-index

80
all docs

80
docs citations

80
times ranked

6024
citing authors

#	ARTICLE	IF	CITATIONS
1	JAK2 or CALR mutation status defines subtypes of essential thrombocythemia with substantially different clinical course and outcomes. <i>Blood</i> , 2014, 123, 1544-1551.	0.6	507
2	Survival and Disease Progression in Essential Thrombocythemia Are Significantly Influenced by Accurate Morphologic Diagnosis: An International Study. <i>Journal of Clinical Oncology</i> , 2011, 29, 3179-3184.	0.8	441
3	Clinical significance of somatic mutation in unexplained blood cytopenia. <i>Blood</i> , 2017, 129, 3371-3378.	0.6	379
4	SF3B1 mutation identifies a distinct subset of myelodysplastic syndrome with ring sideroblasts. <i>Blood</i> , 2015, 126, 233-241.	0.6	361
5	Clinical effect of driver mutations of JAK2, CALR, or MPL in primary myelofibrosis. <i>Blood</i> , 2014, 124, 1062-1069.	0.6	340
6	Prevalence and clinical significance of the MYD88 (L265P) somatic mutation in Waldenström's macroglobulinemia and related lymphoid neoplasms. <i>Blood</i> , 2013, 121, 2522-2528.	0.6	290
7	Prognostic factors for thrombosis, myelofibrosis, and leukemia in essential thrombocythemia: a study of 605 patients. <i>Haematologica</i> , 2008, 93, 1645-1651.	1.7	241
8	Relation between JAK2 (V617F) mutation status, granulocyte activation, and constitutive mobilization of CD34+ cells into peripheral blood in myeloproliferative disorders. <i>Blood</i> , 2006, 107, 3676-3682.	0.6	236
9	Clinical Relevance of Bone Marrow Fibrosis and CD34-Positive Cell Clusters in Primary Myelodysplastic Syndromes. <i>Journal of Clinical Oncology</i> , 2009, 27, 754-762.	0.8	225
10	A prognostic model to predict survival in 867 World Health Organization-defined essential thrombocythemia at diagnosis: a study by the International Working Group on Myelofibrosis Research and Treatment. <i>Blood</i> , 2012, 120, 1197-1201.	0.6	222
11	Driver somatic mutations identify distinct disease entities within myeloid neoplasms with myelodysplasia. <i>Blood</i> , 2014, 124, 1513-1521.	0.6	222
12	Splenic marginal zone lymphoma: a prognostic model for clinical use. <i>Blood</i> , 2006, 107, 4643-4649.	0.6	217
13	The BRAF V600E mutation in hairy cell leukemia and other mature B-cell neoplasms. <i>Blood</i> , 2012, 119, 188-191.	0.6	150
14	Twenty-one cases of blastic plasmacytoid dendritic cell neoplasm: focus on biallelic locus 9p21.3 deletion. <i>Blood</i> , 2011, 118, 4591-4594.	0.6	140
15	Molecular and clinical features of refractory anemia with ringed sideroblasts associated with marked thrombocytosis. <i>Blood</i> , 2009, 114, 3538-3545.	0.6	135
16	Familial Chronic Myeloproliferative Disorders: Clinical Phenotype and Evidence of Disease Anticipation. <i>Journal of Clinical Oncology</i> , 2007, 25, 5630-5635.	0.8	130
17	Splenic and nodal marginal zone lymphomas are indolent disorders at high hepatitis C virus seroprevalence with distinct presenting features but similar morphologic and phenotypic profiles. <i>Cancer</i> , 2004, 100, 107-115.	2.0	121
18	A dynamic prognostic model to predict survival in post-polycythemia vera myelofibrosis. <i>Blood</i> , 2008, 111, 3383-3387.	0.6	108

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19	Initial bone marrow reticulin fibrosis in polycythemia vera exerts an impact on clinical outcome. <i>Blood</i> , 2012, 119, 2239-2241.	0.6	90
20	Pattern of somatic mutations in patients with Waldenström macroglobulinemia or IgM monoclonal gammopathy of undetermined significance. <i>Haematologica</i> , 2017, 102, 2077-2085.	1.7	90
21	Littoral Cell Angiosarcoma of the Spleen Case Report with Immunohistochemical and Ultrastructural Analysis. <i>American Journal of Surgical Pathology</i> , 1995, 19, 1203-1208.	2.1	86
22	Acquired copy-neutral loss of heterozygosity of chromosome 1p as a molecular event associated with marrow fibrosis in MPL-mutated myeloproliferative neoplasms. <i>Blood</i> , 2013, 121, 4388-4395.	0.6	83
23	JAK2 (V617F) as an acquired somatic mutation and a secondary genetic event associated with disease progression in familial myeloproliferative disorders. <i>Cancer</i> , 2006, 107, 2206-2211.	2.0	82
24	Constant activation of the RAF-MEK-ERK pathway as a diagnostic and therapeutic target in hairy cell leukemia. <i>Haematologica</i> , 2013, 98, 635-639.	1.7	75
25	CALR exon 9 mutations are somatically acquired events in familial cases of essential thrombocythemia or primary myelofibrosis. <i>Blood</i> , 2014, 123, 2416-2419.	0.6	66
26	Increased risk of lymphoid neoplasm in patients with myeloproliferative neoplasm: a study of 1,915 patients. <i>Haematologica</i> , 2011, 96, 454-458.	1.7	65
27	Nongastric Marginal Zone B-Cell MALT Lymphoma: Prognostic Value of Disease Dissemination. <i>Oncologist</i> , 2006, 11, 285-291.	1.9	63
28	Primary cutaneous large B-cell lymphoma of the leg: Histogenetic analysis of a controversial clinicopathologic entity. <i>Human Pathology</i> , 2002, 33, 937-943.	1.1	62
29	Clinical, histopathological and molecular characterization of hypoplastic myelodysplastic syndrome. <i>Leukemia</i> , 2019, 33, 2495-2505.	3.3	61
30	Bone marrow microvessel density in chronic myeloproliferative disorders: a study of 115 patients with clinicopathological and molecular correlations. <i>British Journal of Haematology</i> , 2008, 140, 162-168.	1.2	60
31	Red blood cell transfusion-dependency implies a poor survival in primary myelofibrosis irrespective of IPSS and DIPSS. <i>Haematologica</i> , 2011, 96, 167-170.	1.7	60
32	Relationship between clone metrics and clinical outcome in clonal cytopenia. <i>Blood</i> , 2021, 138, 965-976.	0.6	58
33	MYD88 (L265P) mutation is an independent risk factor for progression in patients with IgM monoclonal gammopathy of undetermined significance. <i>Blood</i> , 2013, 122, 2284-2285.	0.6	56
34	Reproducibility of the WHO histological criteria for the diagnosis of Philadelphia chromosome-negative myeloproliferative neoplasms. <i>Modern Pathology</i> , 2014, 27, 814-822.	2.9	48
35	Nodal marginal zone lymphoma: current knowledge and future directions of an heterogeneous disease. <i>European Journal of Haematology</i> , 2009, 83, 165-174.	1.1	47
36	Efficacy of Ruxolitinib in Chronic Eosinophilic Leukemia Associated With a <i>PCM1-JAK2</i> Fusion Gene. <i>Journal of Clinical Oncology</i> , 2013, 31, e269-e271.	0.8	47

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37	Clinical course and outcome of essential thrombocythemia and prefibrotic myelofibrosis according to the revised WHO 2016 diagnostic criteria. <i>Oncotarget</i> , 2017, 8, 101735-101744.	0.8	45
38	Bone marrow stroma CD40 expression correlates with inflammatory mast cell infiltration and disease progression in splenic marginal zone lymphoma. <i>Blood</i> , 2014, 123, 1836-1849.	0.6	37
39	Splenic marginal zone cell lymphoma: Report of an indolent variant without massive splenomegaly presumably representing an early phase of the disease. <i>Human Pathology</i> , 1995, 26, 39-46.	1.1	35
40	Dyspnea secondary to pulmonary hematopoiesis as presenting symptom of myelofibrosis with myeloid metaplasia. <i>American Journal of Hematology</i> , 2006, 81, 124-127.	2.0	35
41	Cutaneous CD30+ lymphoproliferative disorders: Expression of bcl-2 and proteins of the tumor necrosis factor receptor superfamily. <i>Human Pathology</i> , 1998, 29, 1223-1230.	1.1	33
42	CD146+ bone marrow osteoprogenitors increase in the advanced stages of primary myelofibrosis. <i>Haematologica</i> , 2009, 94, 127-130.	1.7	33
43	Depletion of circulating IgM memory B cells predicts unfavourable outcome in COVID-19. <i>Scientific Reports</i> , 2020, 10, 20836.	1.6	32
44	Marginal zone-related neoplasms of splenic and nodal origin. <i>Haematologica</i> , 2003, 88, 80-93.	1.7	31
45	Splenic marginal zone lymphoma: Clinical clustering of immunoglobulin heavy chain repertoires. <i>Blood Cells, Molecules, and Diseases</i> , 2009, 42, 286-291.	0.6	30
46	Assessment of bone marrow involvement in non-Hodgkin's lymphomas: comparison between histology and flow cytometry. <i>European Journal of Haematology</i> , 2010, 85, 405-415.	1.1	30
47	Primary gastric CD30 (Ki-1)-positive large cell non-Hodgkin's lymphomas. A clinicopathologic analysis of six cases. <i>Cancer</i> , 1994, 73, 541-549.	2.0	26
48	Ruxolitinib treatment and risk of B-cell lymphomas in myeloproliferative neoplasms. <i>American Journal of Hematology</i> , 2019, 94, E185-E188.	2.0	26
49	Positive predictive value for malignancy of uncertain malignant potential (B3) breast lesions diagnosed on vacuum-assisted biopsy (VAB): is surgical excision still recommended?. <i>European Radiology</i> , 2021, 31, 920-927.	2.3	23
50	Distinctive Clinical and Histological Features of Waldenström's Macroglobulinemia and Splenic Marginal Zone Lymphoma. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2011, 11, 103-105.	0.2	22
51	High-resolution genome-wide array comparative genomic hybridization in splenic marginal zone B-cell lymphoma. <i>Human Pathology</i> , 2009, 40, 1628-1637.	1.1	21
52	Clinical and molecular characteristics of lymphoplasmacytic lymphoma not associated with an IgM monoclonal protein: A multicentric study of the Rete Ematologica Lombarda (REL) network. <i>American Journal of Hematology</i> , 2019, 94, 1193-1199.	2.0	18
53	CD5 ⁺ diffuse large B-cell lymphoma with peculiar cyclin D1+ phenotype. Pathologic and molecular characterization of a single case. <i>Human Pathology</i> , 2011, 42, 1204-1208.	1.1	15
54	European Bone Marrow Working Group trial on reproducibility of World Health Organization criteria to discriminate essential thrombocythemia from prefibrotic primary myelofibrosis. <i>Haematologica</i> 2012;97(3):360-5 - Comment. <i>Haematologica</i> , 2012, 97, e5-e6.	1.7	15

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55	Diagnosis and management of prefibrotic myelofibrosis. <i>Expert Review of Hematology</i> , 2018, 11, 537-545.	1.0	13
56	A risk stratification model based on the initial concentration of the serum monoclonal protein and MYD88 mutation status identifies a subset of patients with IgM monoclonal gammopathy of undetermined significance at high risk of progression to Waldenström macroglobulinaemia or other lymphoproliferative disorders. <i>British Journal of Haematology</i> , 2019, 187, 441-446.	1.2	13
57	CD56/Neural Cell Adhesion Molecule Expression in Primary Extranodal Ki-1/CD30+ Lymphoma. <i>American Journal of Dermatopathology</i> , 1997, 19, 384-390.	0.3	13
58	Parental origin of the deletion del(20q) in Shwachman-Diamond patients and loss of the paternally derived allele of the imprinted L3MBTL1 gene. <i>Genes Chromosomes and Cancer</i> , 2017, 56, 51-58.	1.5	12
59	Nijmegen Breakage Syndrome-associated T-cell-rich B-cell Lymphoma: Case Report. <i>Pediatric and Developmental Pathology</i> , 2000, 3, 264-270.	0.5	9
60	The Italian Mastocytosis Registry: 6-year experience from a hospital-based registry. <i>Future Oncology</i> , 2018, 14, 2713-2723.	1.1	9
61	CD3+ T large granular lymphocyte leukaemia in a HIV+, HCV+, HBV+ patient. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2009, 454, 349-351.	1.4	8
62	Particulate cytoplasmic structures with high concentration of ubiquitin-proteasome accumulate in myeloid neoplasms. <i>Journal of Hematology and Oncology</i> , 2015, 8, 71.	6.9	8
63	Targeted next-generation sequencing reveals molecular heterogeneity in non-chronic lymphocytic leukemia clonal B-cell lymphocytosis. <i>Hematological Oncology</i> , 2020, 38, 689-697.	0.8	7
64	Validation of cytogenetic-based risk stratification in primary myelofibrosis. <i>Blood</i> , 2010, 115, 2719-2720.	0.6	6
65	Cutaneous involvement by post-polycythemia vera myelofibrosis. <i>American Journal of Hematology</i> , 2014, 89, 448-448.	2.0	6
66	Massive mediastinal enlargement due to extramedullary haematopoiesis in a patient with MYH9-related thrombocytopenia. <i>British Journal of Haematology</i> , 2017, 178, 10-10.	1.2	5
67	First Case of an AIDS Patient With Systemic Mast Cell Disease Associated With Eosinophilia FIP1-Positive Treated With Imatinib Mesylate Therapy. <i>Journal of Clinical Oncology</i> , 2006, 24, e6-e7.	0.8	4
68	An Insidious Presentation of Splenic Marginal Zone Lymphoma. <i>Clinical Lymphoma and Myeloma</i> , 2007, 7, 432-433.	1.4	3
69	c.428T>G (p.V143A) homozygous mutation in TP53 gene as a possible mechanism of resistance to trastuzumab therapy in gastric cancer. <i>Acta Oncologica</i> , 2016, 55, 1373-1375.	0.8	3
70	Vascular endothelial growth factor overexpression in myelodysplastic syndrome bone marrow cells: biological and clinical implications. <i>Leukemia and Lymphoma</i> , 2017, 58, 1711-1720.	0.6	3
71	Telangiectasia macularis eruptiva perstans: a neglected type of mastocytosis with exclusively cutaneous involvement? A case series. <i>European Journal of Dermatology</i> , 2019, 29, 174-178.	0.3	3
72	Histopathological and immunohistochemical evaluation of bone marrow biopsy in myelodysplastic syndromes. <i>International Journal of Hematologic Oncology</i> , 2013, 2, 219-228.	0.7	1

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73	Acute megakaryoblastic leukemia with a novel GATA1 mutation in a second trimester stillborn fetus with trisomy 21. <i>Leukemia and Lymphoma</i> , 2021, 62, 2276-2279.	0.6	1
74	Splenic Marginal Zone B-Cell Lymphoma: Clinical Clustering of Immunoglobulin Heavy Chain Repertoires.. <i>Blood</i> , 2008, 112, 1775-1775.	0.6	1
75	Prevalence and Clinical Significance of the MYD88 (L265P) Somatic Mutation in Patients with Waldenström Macroglobulinemia, IgM-Monoclonal Gammopathy of Undetermined Significance or Other Mature B-Cell Neoplasms.. <i>Blood</i> , 2012, 120, 2667-2667.	0.6	1
76	Aberrant phenotype of plasmacytoid monocytes in acute myeloid leukemia. <i>American Journal of Hematology</i> , 2008, 83, 428-429.	2.0	0
77	Systemic mastocytosis and lymphoplasmacytic lymphoma: an unusual and intriguing form of SM-AHN. <i>Leukemia and Lymphoma</i> , 2021, 62, 1782-1785.	0.6	0
78	Relationship Between Granulocyte JAK2 (V617F) Mutant Allele Burden and Risk of Progression to Myelofibrosis in Polycythemia Vera: a Prospective Study of 338 Patients.. <i>Blood</i> , 2009, 114, 751-751.	0.6	0
79	Bone Marrow Biopsy Revision According to WHO Criteria in 272 Patients of the Registro Italiano Trombocitemia (RIT): Preliminary Report On Clinical and Histopathological Implications.. <i>Blood</i> , 2009, 114, 4974-4974.	0.6	0
80	Constant Activation of the RAF-MEK-ERK Pathway As a Diagnostic and Therapeutic Target in Hairy Cell Leukemia.. <i>Blood</i> , 2012, 120, 2657-2657.	0.6	0