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Erythromelalgic, thrombotic and haemorrhagic manifestations of thrombocythaemia

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#	Paper	IF	Citations
52	Acquired von Willebrand disease as a cause of recurrent mucocutaneous bleeding in primary thrombocythemia: relationship with platelet count. <i>Annals of Hematology</i> , <b>1994</b> , 69, 81-4	3	62
51	The determination of spontaneous megakaryocyte colony formation is an unequivocal test for discrimination between essential thrombocythaemia and reactive thrombocytosis. <i>British Journal of Haematology</i> , <b>1995</b> , 90, 326-31	4.5	32
50	The myeloproliferative disorders. An historical appraisal and personal experiences. <i>Leukemia and Lymphoma</i> , <b>1996</b> , 22 Suppl 1, 1-14	1.9	17
49	Acquired von Willebrand disease in myeloproliferative disorders. <i>Leukemia and Lymphoma</i> , <b>1996</b> , 22 Suppl 1, 79-82	1.9	29
48	Treatment strategies in essential thrombocythemia. A critical appraisal of various experiences in different centers. <i>Leukemia and Lymphoma</i> , <b>1996</b> , 22 Suppl 1, 149-60	1.9	46
47	Bleeding time and platelet function in essential thrombocythemia and other myeloproliferative syndromes. <i>Leukemia and Lymphoma</i> , <b>1996</b> , 22 Suppl 1, 71-8	1.9	45
46	Interferon-alpha in the treatment of essential thrombocythemia. <i>Leukemia and Lymphoma</i> , <b>1996</b> , 22 Suppl 1, 135-42	1.9	66
45	Erythromelalgic, thrombotic and hemorrhagic manifestations in 50 cases of thrombocythemia. Leukemia and Lymphoma, <b>1996</b> , 22 Suppl 1, 47-56	1.9	59
44	The Management of Polycythaemia Vera. <i>Hematology</i> , <b>1997</b> , 2, 55-64	2.2	1
43	Portal vein thrombosis in a 17-year-old female adolescent with essential thrombocytosis. <i>Pediatric Hematology and Oncology</i> , <b>1997</b> , 14, 457-62	1.7	6
42	The use of aspirin in polycythaemia vera and primary thrombocythaemia. <i>Blood Reviews</i> , <b>1998</b> , 12, 12-2	211.1	21
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40	Acquired von Willebrand disease due to increasing platelet count can readily explain the paradox of thrombosis and bleeding in thrombocythemia. <i>Clinical and Applied Thrombosis/Hemostasis</i> , <b>1999</b> , 5, 147	7-3 <sup>3</sup>	31
39	Thrombosis and bleeding in myeloproliferative disorders: identification of at-risk patients with whole blood platelet aggregation studies. <i>British Journal of Haematology</i> , <b>1999</b> , 105, 618-25	4.5	33
38	Acquired von Willebrand syndromes: clinical features, aetiology, pathophysiology, classification and management. <i>Best Practice and Research in Clinical Haematology</i> , <b>2001</b> , 14, 401-36	4.2	111
37	Current treatment practice for essential thrombocythaemia in adults. <i>Expert Opinion on Pharmacotherapy</i> , <b>2001</b> , 2, 385-93	4	2
36	Acquired von Willebrand syndrome: its pathophysiology, laboratory features and management. <i>Journal of Thrombosis and Thrombolysis</i> , <b>2003</b> , 15, 141-9	5.1	23

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35	Acute myelogenous leukemia associated with extreme symptomatic thrombocytosis and chromosome 3q translocation: case report and review of literature. <i>American Journal of Hematology</i> , <b>2003</b> , 72, 20-6	7.1	15
34	Thrombocythaemia and pregnancy. Best Practice and Research in Clinical Haematology, 2003, 16, 227-42	4.2	48
33	Pathophysiology and treatment of platelet-mediated microvascular disturbances, major thrombosis and bleeding complications in essential thrombocythaemia and polycythaemia vera. <i>Platelets</i> , <b>2004</b> , 15, 67-84	3.6	73
32	Risk-adapted therapy in essential thrombocythemia and polycythemia vera. <i>Blood Reviews</i> , <b>2005</b> , 19, 243-52	11.1	65
31	Thrombosis and haemorrhage in polycythaemia vera and essential thrombocythaemia. <i>British Journal of Haematology</i> , <b>2005</b> , 128, 275-90	4.5	371
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29	Evidence-based management of polycythemia vera. <i>Best Practice and Research in Clinical Haematology</i> , <b>2006</b> , 19, 483-93	4.2	12
28	Thrombosis and bleeding in polycythemia vera and essential thrombocythemia: pathogenetic mechanisms and prevention. <i>Best Practice and Research in Clinical Haematology</i> , <b>2006</b> , 19, 617-33	4.2	68
27	Platelet-mediated erythromelalgic, cerebral, ocular and coronary microvascular ischemic and thrombotic manifestations in patients with essential thrombocythemia and polycythemia vera: a distinct aspirin-responsive and coumadin-resistant arterial thrombophilia. <i>Platelets</i> , <b>2006</b> , 17, 528-44	3.6	8o
26	Acquired von Willebrand syndrome: features and management. <i>American Journal of Hematology</i> , <b>2006</b> , 81, 616-23	7.1	69
25	Thrombocytosis: Essential Thrombocythemia and Reactive Causes. 2007, 295-318		
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21	Thrombohemorrhagic complications of myeloproliferative disorders. <i>Blood Reviews</i> , <b>2010</b> , 24, 227-32	11.1	39
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19	Aspirin in low-risk essential thrombocythemia, not so simple after all?. Leukemia Research, 2011, 35, 280	6- <b>-9</b> .7	13
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17	Management of myeloproliferative neoplasms: from academic guidelines to clinical practice. <i>Current Hematologic Malignancy Reports</i> , <b>2012</b> , 7, 50-6	4.4	19
16	Thrombocytosis. <b>2013</b> , 298-323		2
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14	Essential thrombocythemia-related stroke and improvement with tissue plasminogen activator. <i>American Journal of the Medical Sciences</i> , <b>2015</b> , 350, 152	2.2	2
13	Ischemic stroke with essential thrombocythemia: a case series. <i>Journal of Stroke and Cerebrovascular Diseases</i> , <b>2015</b> , 24, 890-3	2.8	20
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4	Risk Classification. <b>2012</b> , 71-84		
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