New prognostic scoring system for primary myelofibrosis International Working Group for Myelofibrosis Research

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Citation Report

#	Article	IF	CITATIONS
1	Facial Cosmetic Filler Injections as Possible Target for Systemic Sarcoidosis in Patients Treated with Interferon for Chronic Hepatitis C: Two Cases. Dermatology, 2008, 217, 81-84.	0.9	52
2	Prospect of JAK2 inhibitor therapy in myeloproliferative neoplasms. Expert Review of Anticancer Therapy, 2009, 9, 663-670.	1.1	28
3	Philadelphia Chromosome–Negative Chronic Myeloproliferative Disease. American Journal of Clinical Pathology, 2009, 132, 261-280.	0.4	33
4	Emerging drugs for the therapy of primary and post essential thrombocythemia, post polycythemia vera myelofibrosis. Expert Opinion on Emerging Drugs, 2009, 14, 471-479.	1.0	19
5	Dynamic Model for Predicting Death Within 12 Months in Patients With Primary or Post–Polycythemia Vera/Essential Thrombocythemia Myelofibrosis. Journal of Clinical Oncology, 2009, 27, 5587-5593.	0.8	117
6	Epigenetic therapy in myeloproliferative neoplasms: evidence and perspectives. Journal of Cellular and Molecular Medicine, 2009, 13, 1437-1450.	1.6	23
7	Conventional cytogenetics in myelofibrosis: literature review and discussion. European Journal of Haematology, 2009, 82, 329-338.	1.1	107
8	Prognostic relevance of cytogenetic abnormalities in primary myelofibrosis: comparison of recent reports from Japan, the Mayo Clinic and MD Anderson Cancer Center. European Journal of Haematology, 2009, 83, 290-291.	1.1	1
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16	Deferasirox treatment may be associated with reversible renal Fanconi syndrome. American Journal of Hematology, 2010, 85, 132-134.	2.0	25
17	Resolution of cerebral artery stenosis in a child with sickle cell anemia treated with hydroxyurea. American Journal of Hematology, 2010, 85, 135-137.	2.0	3
18	Leukocytosis as a risk factor for thrombosis in myeloproliferative neoplasms–biologically plausible but clinically uncertain. American Journal of Hematology, 2010, 85, 93-94.	2.0	11

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21	An NMR Study of the Bortezomib Degradation under Clinical Use Conditions. Advances in Hematology, 2009, 1-5.	0.6	14
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