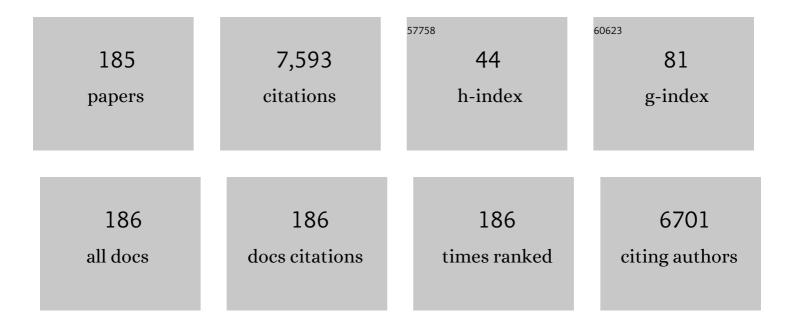
## Khaled M Musallam

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Preoperative anaemia and postoperative outcomes in non-cardiac surgery: a retrospective cohort study. Lancet, The, 2011, 378, 1396-1407.	13.7	1,007
2	Coagulation factor activity and clinical bleeding severity in rare bleeding disorders: results from the European Network of Rare Bleeding Disorders. Journal of Thrombosis and Haemostasis, 2012, 10, 615-621.	3.8	362
3	Overview on practices in thalassemia intermedia management aiming for lowering complication rates across a region of endemicity: the OPTIMAL CARE study. Blood, 2010, 115, 1886-1892.	1.4	315
4	Iron deficiency across chronic inflammatory conditions: International expert opinion on definition, diagnosis, and management. American Journal of Hematology, 2017, 92, 1068-1078.	4.1	290
5	Iron deficiency anaemia revisited. Journal of Internal Medicine, 2020, 287, 153-170.	6.0	233
6	Non-transfusion-dependent thalassemias. Haematologica, 2013, 98, 833-844.	3.5	231
7	Optimal management of β thalassaemia intermedia. British Journal of Haematology, 2011, 152, 512-523.	2.5	187
8	β-Thalassemias. New England Journal of Medicine, 2021, 384, 727-743.	27.0	183
9	Clinical experience with fetal hemoglobin induction therapy in patients with β-thalassemia. Blood, 2013, 121, 2199-2212.	1.4	154
10	Elevated liver iron concentration is a marker of increased morbidity in patients with  thalassemia intermedia. Haematologica, 2011, 96, 1605-1612.	3.5	153
11	Smoking and the Risk of Mortality and Vascular and Respiratory Events in Patients Undergoing Major Surgery. JAMA Surgery, 2013, 148, 755.	4.3	140
12	Bone disease and skeletal complications in patients with $\hat{I}^2$ thalassemia major. Bone, 2011, 48, 425-432.	2.9	127
13	Splenectomy and thrombosis: the case of thalassemia intermedia. Journal of Thrombosis and Haemostasis, 2010, 8, 2152-2158.	3.8	120
14	Residual plasmatic activity of ADAMTS13 is correlated with phenotype severity in congenital thrombotic thrombocytopenic purpura. Blood, 2012, 120, 440-448.	1.4	107
15	Iron overload in non-transfusion-dependent thalassemia: a clinical perspective. Blood Reviews, 2012, 26, S16-S19.	5.7	105
16	Â-Thalassemia Intermedia: A Clinical Perspective. Cold Spring Harbor Perspectives in Medicine, 2012, 2, a013482-a013482.	6.2	102
17	Prevalence and Risk Factors for Pulmonary Arterial Hypertension in a Large Group of β-Thalassemia Patients Using Right Heart Catheterization. Circulation, 2014, 129, 338-345.	1.6	101
18	Ageâ€related complications in treatmentâ€naÃ⁻ve patients with thalassaemia intermedia. British Journal of Haematology, 2010, 150, 486-489.	2.5	100

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19	Fetal hemoglobin levels and morbidity in untransfused patients with β-thalassemia intermedia. Blood, 2012, 119, 364-367.	1.4	85
20	Iron deficiency beyond erythropoiesis: should we be concerned?. Current Medical Research and Opinion, 2018, 34, 81-93.	1.9	83
21	Cerebral infarction in β-thalassemia intermedia: Breaking the silence. Thrombosis Research, 2012, 130, 695-702.	1.7	81
22	Impact of Preoperative Anaemia and Blood Transfusion on Postoperative Outcomes in Gynaecological Surgery. PLoS ONE, 2015, 10, e0130861.	2.5	80
23	Serum ferritin levels and endocrinopathy in medically treated patients with β thalassemia major. Annals of Hematology, 2012, 91, 1107-1114.	1.8	79
24	Redefining thalassemia as a hypercoagulable state. Annals of the New York Academy of Sciences, 2010, 1202, 231-236.	3.8	78
25	ADAMTSâ€13 activity and autoantibodies classes and subclasses as prognostic predictors in acquired thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2012, 10, 1556-1565.	3.8	74
26	Asymptomatic brain magnetic resonance imaging abnormalities in splenectomized adults with thalassemia intermedia. Journal of Thrombosis and Haemostasis, 2010, 8, 54-59.	3.8	72
27	Renal complications in transfusion-dependent beta thalassaemia. Blood Reviews, 2010, 24, 239-244.	5.7	70
28	Hypercoagulability in β-thalassemia: a status quo. Expert Review of Hematology, 2012, 5, 505-512.	2.2	70
29	Treating iron overload in patients with nonâ€transfusionâ€dependent thalassemia. American Journal of Hematology, 2013, 88, 409-415.	4.1	67
30	Levels of nonâ€transferrinâ€bound iron as an index of iron overload in patients with thalassaemia intermedia. British Journal of Haematology, 2009, 146, 569-572.	2.5	66
31	Discontinuing prophylactic transfusions increases the risk of silent brain infarction in children with sickle cell disease: data from STOP II. Blood, 2011, 118, 894-898.	1.4	62
32	Magnetic resonance evaluation of hepatic and myocardial iron deposition in transfusionâ€independent thalassemia intermedia compared to regularly transfused thalassemia major patients. American Journal of Hematology, 2010, 85, 288-290.	4.1	61
33	Iron Overload: Consequences, Assessment, and Monitoring. Hemoglobin, 2009, 33, S46-S57.	0.8	59
34	Mechanisms of Renal Disease in β-Thalassemia. Journal of the American Society of Nephrology: JASN, 2012, 23, 1299-1302.	6.1	58
35	Risk factors for pulmonary hypertension in patients with β thalassemia intermedia. European Journal of Internal Medicine, 2011, 22, 607-610.	2.2	56
36	Serum ferritin level and morbidity risk in transfusion-independent patients with Â-thalassemia intermedia: the ORIENT study. Haematologica, 2014, 99, e218-e221.	3.5	56

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37	Absence of cardiac siderosis despite hepatic iron overload in Italian patients with thalassemia intermedia: an MRI T2* study. Annals of Hematology, 2010, 89, 585-589.	1.8	55
38	Levels of growth differentiation factor-15 are high and correlate with clinical severity in transfusion-independent patients with β thalassemia intermedia. Blood Cells, Molecules, and Diseases, 2011, 47, 232-234.	1.4	55
39	Ineffective Erythropoiesis: Anemia and Iron Overload. Hematology/Oncology Clinics of North America, 2018, 32, 213-221.	2.2	54
40	Incidence and prophylaxis of venous thromboembolic events in multiple myeloma patients receiving immunomodulatory therapy. Thrombosis Research, 2009, 123, 679-686.	1.7	53
41	Cross-Talk between Available Guidelines for the Management of Patients with Beta-Thalassemia Major. Acta Haematologica, 2013, 130, 64-73.	1.4	49
42	Evaluation of the 5mg/g liver iron concentration threshold and its association with morbidity in patients with β-thalassemia intermedia. Blood Cells, Molecules, and Diseases, 2013, 51, 35-38.	1.4	47
43	Brain magnetic resonance angiography in splenectomized adults with β-thalassemia intermedia. European Journal of Haematology, 2011, 87, 539-546.	2.2	46
44	Iron overload and chelation therapy in myelodysplastic syndromes. Critical Reviews in Oncology/Hematology, 2014, 91, 64-73.	4.4	46
45	Hypercoagulability in non-transfusion-dependent thalassemia. Blood Reviews, 2012, 26, S20-S23.	5.7	44
46	Sustained improvements in myocardial T2* over 2 years in severely ironâ€overloaded patients with beta thalassemia major treated with deferasirox or deferoxamine. American Journal of Hematology, 2015, 90, 91-96.	4.1	43
47	Iron overload across the spectrum of nonâ€transfusionâ€dependent thalassaemias: role of erythropoiesis, splenectomy and transfusions. British Journal of Haematology, 2017, 176, 288-299.	2.5	43
48	Iron deficiency in chronic heart failure: caseâ€based practical guidance. ESC Heart Failure, 2018, 5, 764-771.	3.1	43
49	Longitudinal changes in serum ferritin levels correlate with measures of hepatic stiffness in transfusion-independent patients with β-thalassemia intermedia. Blood Cells, Molecules, and Diseases, 2012, 49, 136-139.	1.4	42
50	Iron overload in β-thalassemia intermedia. Current Opinion in Hematology, 2013, 20, 187-192.	2.5	42
51	Ratio Between Positive Lymph Nodes and Total Excised Axillary Lymph Nodes as an Independent Prognostic Factor for Overall Survival in Patients with Nonmetastatic Lymph Node-Positive Breast Cancer. Annals of Surgical Oncology, 2009, 16, 3388-3395.	1.5	41
52	Thrombosis in Thalassemia: Why are we so Concerned?. Hemoglobin, 2011, 35, 503-510.	0.8	40
53	Contemporary approaches to treatment of beta-thalassemia intermedia. Blood Reviews, 2012, 26, S24-S27.	5.7	40
54	The effect of maternal fasting during Ramadan on preterm delivery: a prospective cohort study. BJOC: an International Journal of Obstetrics and Gynaecology, 2012, 119, 1379-1386.	2.3	40

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55	THALASSEMIA AND VENOUS THROMBOEMBOLISM. Mediterranean Journal of Hematology and Infectious Diseases, 2011, 3, e2011025.	1.3	39
56	Health-related quality of life in adults with transfusion-independent thalassaemia intermedia compared to regularly transfused thalassaemia major: new insights. European Journal of Haematology, 2011, 87, 73-79.	2.2	39
57	2021 update on clinical trials in βâ€ŧhalassemia. American Journal of Hematology, 2021, 96, 1518-1531.	4.1	38
58	Challenges Associated With Prolonged Survival of Patients With Thalassemia: Transitioning From Childhood to Adulthood. Pediatrics, 2008, 121, e1426-e1429.	2.1	34
59	Thalassaemia Intermedia: an Update. Mediterranean Journal of Hematology and Infectious Diseases, 2009, 1, e2009004.	1.3	34
60	Absence of cardiac siderosis by MRI T2* despite transfusion burden, hepatic and serum iron overload in Lebanese patients with sickle cell disease. European Journal of Haematology, 2009, 83, 565-571.	2.2	33
61	Insight onto the Pathophysiology and Clinical Complications of Thalassemia Intermedia. Hemoglobin, 2009, 33, S145-S159.	0.8	33
62	Overview of Iron Chelation Therapy with Desferrioxamine and Deferiprone. Hemoglobin, 2009, 33, S58-S69.	0.8	33
63	Aspirin Resistance. Advances in Hematology, 2009, 2009, 1-10.	1.0	32
64	Implementation of an Emergency Department Computer System: Design Features That Users Value. Journal of Emergency Medicine, 2011, 41, 693-700.	0.7	32
65	ls VEGF a predictive biomarker to anti-angiogenic therapy?. Critical Reviews in Oncology/Hematology, 2011, 79, 103-111.	4.4	31
66	Assessment and management of iron overload in βâ€thalassaemia major patients during the 21st century: a realâ€life experience from the <scp>I</scp> talian <scp>W</scp> ebthal project. British Journal of Haematology, 2013, 161, 872-883.	2.5	31
67	Glomerular Hyperfiltration and Proteinuria in Transfusion-Independent Patients with β-Thalassemia Intermedia. Nephron, 2013, 121, c136-c143.	1.8	31
68	Postoperative Outcomes After Laparoscopic Splenectomy Compared With Open Splenectomy. Annals of Surgery, 2013, 257, 1116-1123.	4.2	29
69	The risk of local recurrence along the core-needle biopsy tract in patients with bone sarcomas. Iowa orthopaedic journal, The, 2010, 30, 80-3.	0.5	29
70	Revisiting the nonâ€transfusionâ€dependent (NTDT) vs. transfusionâ€dependent (TDT) thalassemia classification 10 years later. American Journal of Hematology, 2021, 96, E54-E56.	4.1	28
71	Iron overload indices rise linearly with transfusion rate in patients with sickle cell disease. Blood, 2010, 115, 2980-2981.	1.4	27
72	The Spine in β-Thalassemia Syndromes. Spine, 2012, 37, 334-339.	2.0	27

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73	Brain positron emission tomography in splenectomized adults with β-thalassemia intermedia: uncovering yet another covert abnormality. Annals of Hematology, 2012, 91, 235-241.	1.8	27
74	Survival and causes of death in 2,033 patients with non-transfusion-dependent β-thalassemia. Haematologica, 2021, 106, 2489-2492.	3.5	25
75	Abdominal Manifestations of Multiple Myeloma: A Retrospective Radiologic Overview. Clinical Lymphoma and Myeloma, 2008, 8, 348-351.	1.4	24
76	Prevalence of Depression and Anxiety in Adult Patients with β-Thalassemia Major and Intermedia. International Journal of Psychiatry in Medicine, 2012, 44, 291-303.	1.8	24
77	Elevated prepartum fibrinogen levels are not associated with a reduced risk of postpartum hemorrhage. Journal of Thrombosis and Haemostasis, 2012, 10, 1451-1453.	3.8	24
78	Oneâ€year results from a prospective randomized trial comparing phlebotomy with deferasirox for the treatment of iron overload in pediatric patients with thalassemia major following curative stem cell transplantation. Pediatric Blood and Cancer, 2017, 64, 188-196.	1.5	24
79	Childhood stroke in a child with familial Mediterranean fever carrying several prothrombotic risk factors. Lupus, 2009, 18, 845-847.	1.6	23
80	A closer look at paroxysmal nocturnal hemoglobinuria. European Journal of Internal Medicine, 2010, 21, 260-267.	2.2	23
81	Left ventricular noncompaction in patients with βâ€ŧhalassemia: Uncovering a previously unrecognized abnormality. American Journal of Hematology, 2012, 87, 1079-1083.	4.1	23
82	Establishment of a bleeding score as a diagnostic tool for patients with rare bleeding disorders. Thrombosis Research, 2016, 148, 128-134.	1.7	22
83	Preoperative INR and postoperative major bleeding and mortality: A retrospective cohort study. Journal of Thrombosis and Thrombolysis, 2016, 41, 301-311.	2.1	22
84	Potential mechanisms for renal damage in beta-thalassemia. Journal of Nephrology, 2013, 26, 821-828.	2.0	22
85	Morbidity-free survival and hemoglobin level in non-transfusion-dependent β-thalassemia: a 10-year cohort study. Annals of Hematology, 2022, 101, 203-204.	1.8	21
86	Hemoglobin level and morbidity in non-transfusion-dependent thalassemia. Blood Cells, Molecules, and Diseases, 2015, 55, 108-109.	1.4	20
87	Hypercoagulability and Vascular Disease. Hematology/Oncology Clinics of North America, 2018, 32, 237-245.	2.2	20
88	Variations in hemoglobin level and morbidity burden in non-transfusion-dependent β-thalassemia. Annals of Hematology, 2021, 100, 1903-1905.	1.8	20
89	Risk of mortality from anemia and iron overload in nontransfusionâ€dependent βâ€thalassemia. American Journal of Hematology, 2022, 97, .	4.1	19
90	Primary colorectal lymphoma. Medical Oncology, 2010, 27, 249-254.	2.5	18

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91	Iron in sickleâ€cell disease: What have we learned over the years?. Pediatric Blood and Cancer, 2011, 56, 182-190.	1.5	18
92	The emerging concept of residual ADAMTS13 activity in ADAMTS13-deficient thrombotic thrombocytopenic purpura. Blood Reviews, 2013, 27, 71-76.	5.7	17
93	The Hypercoagulable State in Thalassemia Intermedia. Hemoglobin, 2009, 33, S160-S169.	0.8	16
94	Correlation of non-mass-like abnormal MR signal intensity with pathological findings surrounding pediatric osteosarcoma and Ewing's sarcoma. Skeletal Radiology, 2012, 41, 1453-1461.	2.0	16
95	Postoperative outcomes following pancreaticoduodenectomy: how should age affect clinical practice?. World Journal of Surgical Oncology, 2013, 11, 131.	1.9	16
96	Solitary mediastinal lymph node metastasis in rectosigmoid carcinoma: a case report. Cases Journal, 2008, 1, 69.	0.4	15
97	Iron chelation therapy for patients with sickle cell disease and iron overload. American Journal of Hematology, 2010, 85, 782-786.	4.1	15
98	Development of a new disease severity scoring system for patients with non-transfusion-dependent thalassemia. European Journal of Internal Medicine, 2016, 28, 91-96.	2.2	14
99	Recent advances and treatment challenges in patients with non-transfusion-dependent thalassemia. Blood Reviews, 2012, 26, S1-S2.	5.7	13
100	Preventing Thalassemia in Lebanon: Successes and Challenges in a Developing Country. Hemoglobin, 2014, 38, 308-311.	0.8	13
101	Serum ferritin values between 300 and 800 ng/mL in nontransfusionâ€dependent thalassemia: A probability curve to guide clinical decision making when MRI is unavailable. American Journal of Hematology, 2017, 92, E35-E37.	4.1	13
102	Longâ€ŧerm improvement in cardiac magnetic resonance in βâ€ŧhalassemia major patients treated with deferasirox extends to patients with abnormal baseline cardiac function. American Journal of Hematology, 2019, 94, 312-318.	4.1	13
103	Antibiotic use and risk of gynecological cancer. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2011, 159, 388-393.	1.1	12
104	Differential effects of the type of iron chelator on the absolute number of hematopoietic peripheral progenitors in patients with Â-thalassemia major. Haematologica, 2013, 98, 555-559.	3.5	12
105	Relationship between transfusion burden, healthcare resource utilization, and complications in patients with betaâ€thalassemia in Taiwan: A realâ€world analysis. Transfusion, 2021, 61, 2906-2917.	1.6	12
106	Management of nonâ€ŧransfusionâ€dependent βâ€ŧhalassemia ( <scp>NTDT</scp> ): The next 5 years. Amer Journal of Hematology, 2021, 96, E57-E59.	rican 4.1	11
107	Serum Ferritin Levels and Morbidity in β-Thalassemia Intermedia: A 10-Year Cohort Study. Blood, 2012, 120, 1021-1021.	1.4	11
108	The impact of zoledronic acid on regenerate and native bone after consolidation and removal of the external fixator: An animal model study. Bone, 2010, 46, 363-368.	2.9	10

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109	Iron Chelation Therapy for Transfusional Iron Overload: A Swift Evolution. Hemoglobin, 2011, 35, 565-573.	0.8	10
110	Mortality in β-thalassemia patients with confirmed pulmonary arterial hypertension on right heart catheterization. Blood, 2022, 139, 2080-2083.	1.4	10
111	Sickle Cell Disease at the Dawn of the Molecular Era. Hemoglobin, 2009, 33, S93-S106.	0.8	9
112	Hepatitis C antiviral response in thalassemia: what is the role of liver iron concentration?. Annals of Hematology, 2009, 88, 1033-1034.	1.8	9
113	Preoperative Pneumonia and Postoperative Venous Thrombosis: A Cohort Study of 427,656ÂPatients Undergoing Major General Surgery. World Journal of Surgery, 2016, 40, 1288-1294.	1.6	9
114	Less â€~reds' more â€~blues': hemoglobin level and depression in non-transfusion-dependent thalassemia. Annals of Hematology, 2020, 99, 903-904.	1.8	9
115	Transfusion independence in Diamond-Blackfan anemia after deferasirox therapy. Annals of Hematology, 2009, 88, 1263-1264.	1.8	8
116	Carbamazepine-induced thrombocytopenia. Blood Cells, Molecules, and Diseases, 2012, 48, 197-198.	1.4	8
117	End stage renal disease in six patients with beta-thalassemia intermedia. Blood Cells, Molecules, and Diseases, 2013, 51, 146-148.	1.4	8
118	A liver mass in an iron-overloaded thalassaemia intermedia patient. British Journal of Haematology, 2013, 161, 1-1.	2.5	8
119	Iron chelation therapy for non-transfusion-dependent thalassemia (NTDT): A status quo. Blood Cells, Molecules, and Diseases, 2014, 52, 88-90.	1.4	8
120	Primary <i>HBB</i> gene mutation severity and longâ€ŧerm outcomes in a global cohort of βâ€ŧhalassaemia. British Journal of Haematology, 2022, 196, 414-423.	2.5	8
121	Silent Brain Abnormalities in Thalassemia Intermedia: A Combined MRI/PET Study Blood, 2009, 114, 4077-4077.	1.4	8
122	Pulmonary Complications of Sickle Cell Disease. Hemoglobin, 2011, 35, 625-635.	0.8	7
123	Cerebral Infarction in Children with Sickle Cell Disease: A Concise Overview. Hemoglobin, 2011, 35, 618-624.	0.8	7
124	Preoperative Hematocrit Concentration and the Risk of Stroke in Patients Undergoing Isolated Coronary-Artery Bypass Grafting. Anemia, 2013, 2013, 1-7.	1.7	7
125	A killer revealed: 10-year experience with beta-thalassemia intermedia. Hematology, 2014, 19, 196-198.	1.5	7
126	Development of a thalassemiaâ€related thrombosis risk scoring system. American Journal of Hematology, 2019, 94, E207-E209.	4.1	7

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127	Burkitt's lymphoma of the colon and bronchi: three case reports. Cases Journal, 2008, 1, 15.	0.4	6
128	Deferiprone or deferasirox for cardiac siderosis in beta-thalassemia major. Haematologica, 2011, 96, e5-e6.	3.5	6
129	von Willebrand factor propeptide to antigen ratio in acquired thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2012, 10, 728-730.	3.8	6
130	Thalassemia and autoimmune diseases: Absence of evidence or evidence of absence?. Blood Reviews, 2022, 52, 100874.	5.7	6
131	Solitary Hydatid Cyst of the Thigh: A Challenging Diagnosis. Vector-Borne and Zoonotic Diseases, 2009, 9, 743-745.	1.5	5
132	High-output cardiac failure secondary to a large arteriovenous fistula: a persistent threat to the dialysis and kidney transplant patient. CKJ: Clinical Kidney Journal, 2009, 2, 147-148.	2.9	5
133	Small bowel intussusception following blunt abdominal trauma in an adult patient. Emergency Medicine Journal, 2009, 26, 752-753.	1.0	5
134	Does absolute excess of alpha chains compromise the benefit of splenectomy in patients with thalassemia intermedia?. Haematologica, 2012, 97, 151-153.	3.5	5
135	Emerging Therapies. Hematology/Oncology Clinics of North America, 2018, 32, 343-352.	2.2	5
136	Distinctive Presentation of a Diaphragmatic Hernia 15 Years After A Traumatic Insult. Annals of Thoracic Surgery, 2009, 88, 651-653.	1.3	4
137	Covert brain ischaemia in splenectomised adults with thalassemia intermedia: An emerging entity. Thrombosis and Haemostasis, 2010, 104, 652-653.	3.4	4
138	Molecular Spectra and Frequency Patterns of Somatic Mutations in Arab Women with Breast Cancer. Oncologist, 2021, 26, e2086-e2089.	3.7	4
139	Prevalence and Risk Factors of Left Ventricular Noncompaction in Patients with Î <sup>2</sup> -Thalassemia Blood, 2012, 120, 2127-2127.	1.4	4
140	Effects of divalproex sodium on hemoglobin level. Blood Cells, Molecules, and Diseases, 2009, 43, 49-52.	1.4	3
141	Ratio between positive lymph nodes and total excised axillary lymph nodes as an independent prognostic factor for overall survival in patients with nonmetastatic lymph node-positive breast cancer. Indian Journal of Surgical Oncology, 2010, 1, 68-75.	0.7	3
142	Predicting venous thromboembolism in hospitalized medical patients: are we there yet?. Expert Review of Hematology, 2011, 4, 1-3.	2.2	3
143	Sequential Therapy with Gemcitabine and Carboplatin Followed by Paclitaxel as First Line Treatment for Advanced Urothelial Cancer. Journal of Cancer, 2012, 3, 362-368.	2.5	3
144	Deferiprone-induced seizures in a patient with β-thalassemia major. Blood Cells, Molecules, and Diseases, 2013, 51, 94-95.	1.4	3

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145	Random Forest Clustering Identifies Three Subgroups of β-Thalassemia with Distinct Clinical Severity. Thalassemia Reports, 2022, 12, 14-23.	0.5	3
146	Iron and hepatitis C: What can we learn from thalassemia major?. Liver International, 2008, 28, 892-893.	3.9	2
147	Malignancy and hypercoagulability: a two-way association revisited. Journal of Thrombosis and Thrombolysis, 2010, 30, 340-341.	2.1	2
148	JAK2V617F and Prothrombin G20210A Gene Mutations in a Patient With Budd-Chiari Syndrome and Essential Thrombocythemia. Clinical and Applied Thrombosis/Hemostasis, 2010, 16, 472-474.	1.7	2
149	Managing unresponsiveness or intolerance to deferasirox therapy: a tale of two doses. Expert Review of Hematology, 2011, 4, 411-414.	2.2	2
150	ANTITHROMBOTIC PROPHYLAXIS IN THE MIDDLE EAST. Mediterranean Journal of Hematology and Infectious Diseases, 2011, 3, e2011023.	1.3	2
151	Case report: use of thienopyridines in a patient with acquired idiopathic thrombotic thrombocytopenic purpura. Journal of Thrombosis and Thrombolysis, 2012, 34, 416-418.	2.1	2
152	Mondor's Disease of the Breast in the Context of Inherited Thrombophilia. Breast Journal, 2012, 18, 373-374.	1.0	2
153	Switching patients from warfarin to dabigatran therapy: To RE-LY or not to rely. International Journal of Cardiology, 2012, 154, e27-e28.	1.7	2
154	Iron overload in non-transfusion-dependent thalassemia. Thalassemia Reports, 2013, 3, 11.	0.5	2
155	On the use of substandard medicines in hematology: An emerging concern in the Middle East and North Africa region. European Journal of Internal Medicine, 2018, 48, e40-e41.	2.2	2
156	1-Year Results From A Prospective Randomized Trial Comparing Phlebotomy with Deferasirox for the Treatment of Iron Overload in Pediatric Patients with Thalassemia Major Following Curative Stem Cell Transplantation. Blood, 2011, 118, 904-904.	1.4	2
157	Elevated Hematocrit Concentration and the Risk of Mortality and Vascular Events in Patients Undergoing Major Surgery Blood, 2012, 120, 2088-2088.	1.4	2
158	A novel genotype c.1228C>G/c.1448C-1498C (L371V/Rec-Ncil) in a 3-year-old child with type 1 Gaucher disease. Journal of Applied Genetics, 2008, 49, 421-424.	1.9	1
159	Axillary Lymph Node Ratio Revisited. Journal of Clinical Oncology, 2009, 27, e67-e67.	1.6	1
160	Antiphospholipid Syndrome, Microalbuminuria, and Risk of Venous Thromboembolism. JAMA - Journal of the American Medical Association, 2009, 302, 945.	7.4	1
161	Clinical images: Severe photosensitive skin reaction secondary to an herbal treatment in a patient with systemic lupus erythematosus. Arthritis and Rheumatism, 2009, 60, 2854-2854.	6.7	1
162	The use of biological therapy in refractory rheumatic diseases other than rheumatoid arthritis: experience at a tertiary care center in Lebanon. Rheumatology International, 2009, 29, 1255-1257.	3.0	1

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163	Thromboangiitis obliterans and the inherited thrombophilias: does an association merit consideration?. Blood Coagulation and Fibrinolysis, 2009, 20, 223-224.	1.0	1
164	The role of liver iron in hepatitis C antiviral treatment. European Journal of Gastroenterology and Hepatology, 2010, 22, 769.	1.6	1
165	The wanderer. American Journal of Obstetrics and Gynecology, 2010, 202, 662.e1.	1.3	1
166	A new chelator in the house. Blood, 2012, 119, 3191-3192.	1.4	1
167	Association between cardiac T2* magnetic resonance imaging values and endocrine function tests in patients with β-thalassemia major. Blood Cells, Molecules, and Diseases, 2014, 52, 50-51.	1.4	1
168	Iron Overload Indices In Thalassemia Major Children Cured by Stem Cell Transplantation at Enrollment In a Prospective Randomized Trial Comparing Phlebotomy and Deferasirox. Blood, 2010, 116, 2082-2082.	1.4	1
169	Evaluation of the 5 Mg/g Liver Iron Concentration Threshold and Its Association with Vascular and Endocrine/Bone Morbidity in β-Thalassemia Intermedia. Blood, 2012, 120, 1024-1024.	1.4	1
170	Greater Red Blood Cell Transfusion Burden Is Associated with More Healthcare Resource Utilization in Patients with Beta-Thalassemia. Blood, 2019, 134, 5790-5790.	1.4	1
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