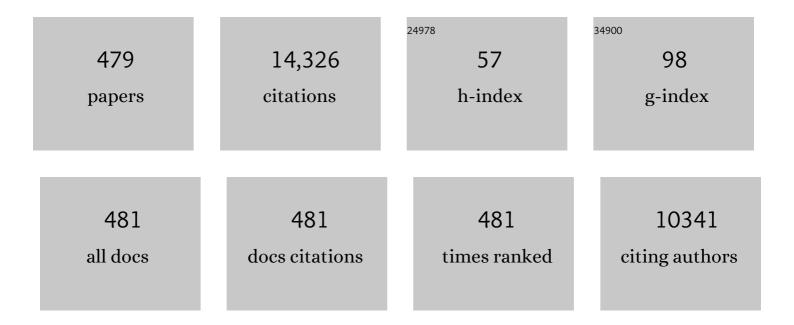
List of Publications by Year in descending order

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ΔΙΙΤΤΛΗΕΡ

#	Article	IF	CITATIONS
1	Morbidity-free survival and hemoglobin level in non-transfusion-dependent Î ² -thalassemia: a 10-year cohort study. Annals of Hematology, 2022, 101, 203-204.	0.8	21
2	Psychoâ€oncology in the Arab world: The time is now. Psycho-Oncology, 2022, 31, 148-151.	1.0	3
3	Thalassemia and autoimmune diseases: Absence of evidence or evidence of absence?. Blood Reviews, 2022, 52, 100874.	2.8	6
4	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.	1.2	8
5	Manifestation of paroxysmal nocturnal hemoglobinuria after COVID-19 mRNA vaccination. Blood Cells, Molecules, and Diseases, 2022, 93, 102641.	0.6	3
6	Clinical burden of hemophilia in older adults: Beyond bleeding risk. Blood Reviews, 2022, 53, 100912.	2.8	3
7	Deep venous thrombosis and pulmonary embolism after COVID-19 mRNA vaccination. Annals of Hematology, 2022, 101, 1111-1113.	0.8	11
8	Anhidrosis associated with long-term use of hydroxyurea in a patient with myeloproliferative neoplasm. Annals of Hematology, 2022, , 1.	0.8	2
9	Inflammatory Bowel Disease: An Indication to Screen for Thrombophilia?. Diseases (Basel,) Tj ETQq1 1 0.784314	rgBT/Ove	erlock 10 Tf 5
10	Random Forest Clustering Identifies Three Subgroups of β-Thalassemia with Distinct Clinical Severity. Thalassemia Reports, 2022, 12, 14-23.	0.1	3
11	Coronavirus Disease 2019 (COVID-19)- Associated central retinal vein occlusion: A case report and literature review. Archives of Hematology Case Reports and Reviews, 2022, 7, 009-012.	0.3	1
12	Risk of mortality from anemia and iron overload in nontransfusionâ€dependent βâ€ŧhalassemia. American Journal of Hematology, 2022, 97, .	2.0	19
13	Redox Balance in β-Thalassemia and Sickle Cell Disease: A Love and Hate Relationship. Antioxidants, 2022, 11, 967.	2.2	5
14	An openâ€label, multicenter, efficacy, and safety study of deferasirox in ironâ€overloaded patients with nonâ€transfusionâ€dependent thalassemia (<scp>THETIS</scp>): 5â€year results. American Journal of Hematology, 2022, 97, .	2.0	2
15	Crushed deferasirox <scp>filmâ€coated</scp> tablets in pediatric patients with transfusional hemosiderosis: Results from a <scp>singleâ€arm</scp> , interventional phase 4 study (<scp>MIMAS</scp>). American Journal of Hematology, 2022, 97, .	2.0	1
16	Pharmacological Induction of Fetal Hemoglobin in β-Thalassemia and Sickle Cell Disease: An Updated Perspective. Pharmaceuticals, 2022, 15, 753.	1.7	14
17	Could Preoperative Unintended Weight Loss Predispose to Postoperative Thrombosis in Patients Undergoing Colorectal Cancer Surgery? An Analysis of the NSQIP Data. Journal of the American College of Nutrition, 2021, 40, 141-147.	1.1	3
18	A land in agony: COVIDâ€19, economic collapse, political corruption, and a deadly blast. American Journal of Hematology, 2021, 96, E1-E2.	2.0	9

#	Article	IF	CITATIONS
19	Management of nonâ€transfusionâ€dependent βâ€thalassemia (<scp>NTDT</scp>): The next 5 years. Ameri Journal of Hematology, 2021, 96, E57-E59.	can 2.0	11
20	Revisiting the nonâ€transfusionâ€dependent (NTDT) vs. transfusionâ€dependent (TDT) thalassemia classification 10 years later. American Journal of Hematology, 2021, 96, E54-E56.	2.0	28
21	β-Thalassemias. New England Journal of Medicine, 2021, 384, 727-743.	13.9	183
22	CYP450 Mediates Reactive Oxygen Species Production in a Mouse Model of β-Thalassemia through an Increase in 20-HETE Activity. International Journal of Molecular Sciences, 2021, 22, 1106.	1.8	6
23	Antiphospholipid Antibodies in Inflammatory and Autoimmune Rheumatic and Musculoskeletal Diseases Beyond Lupus: A Systematic Review of the Available Evidence. Rheumatology and Therapy, 2021, 8, 81-94.	1.1	11
24	The use of luspatercept for thalassemia in adults. Blood Advances, 2021, 5, 326-333.	2.5	28
25	An Overview of Hepatocellular Carcinoma (HCC) in Lebanon: A Focus on Hepatitis B- and Thalassemia-Related HCC. , 2021, , 375-386.		0
26	Hepatocellular Carcinoma in Lebanon and Its Association with Thalassemia. , 2021, , 371-374.		0
27	Variations in hemoglobin level and morbidity burden in non-transfusion-dependent Î ² -thalassemia. Annals of Hematology, 2021, 100, 1903-1905.	0.8	20
28	COVID-19 pandemic and transfusion medicine: the worldwide challenge and its implications. Annals of Hematology, 2021, 100, 1115-1122.	0.8	20
29	Association between Radiotherapy and Risk of Cancer Associated Venous Thromboembolism: A Sub-Analysis of the COMPASS—CAT Study. Cancers, 2021, 13, 1033.	1.7	12
30	Antiphospholipid syndrome: the need for new international classification criteria. Expert Review of Clinical Immunology, 2021, 17, 385-394.	1.3	6
31	Intravenous immunoglobulin, neonatal alloimmune thrombocytopenia, and Sjogren's syndrome: A case report. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 258, 476-477.	0.5	4
32	Recommendations for pregnancy in Fanconi anemia. Expert Opinion on Biological Therapy, 2021, 21, 1-7.	1.4	2
33	Survival and causes of death in 2,033 patients with non-transfusion-dependent β-thalassemia. Haematologica, 2021, 106, 2489-2492.	1.7	25
34	Haematological effects of oral administration of bitopertin, a glycine transport inhibitor, in patients with nonâ€transfusionâ€dependent βâ€thalassaemia. British Journal of Haematology, 2021, 194, 474-477.	1.2	10
35	<scp>SARSâ€CoV</scp> â€2 infection in patients with βâ€thalassemia: Experience from <scp>Lebanon</scp> . American Journal of Hematology, 2021, 96, E285-E288.	2.0	3
36	Polymyalgia rheumatica-like presentation in a case of diffuse large B-cell lymphoma: a diagnostic pitfall. Journal of International Medical Research, 2021, 49, 030006052110185.	0.4	0

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37	The COVID-19 Pandemic and the Need for an Integrated and Equitable Approach: An International Expert Consensus Paper. Thrombosis and Haemostasis, 2021, 121, 992-1007.	1.8	21
38	Antiphospholipid antibodies and cerebrovascular thrombosis in the pediatric population: Few answers to many questions. Lupus, 2021, 30, 1365-1377.	0.8	0
39	The EHA Research Roadmap: Anemias. HemaSphere, 2021, 5, e607.	1.2	7
40	Implemented Interventions at the Naef K. Basile Cancer Institute to Protect Patients and Medical Personnel From COVID Infections: Effectiveness and Patient Satisfaction. Frontiers in Oncology, 2021, 11, 685107.	1.3	1
41	Abstract 56: Landscape and Challenges of Cancer Registration in the Middle East, North Africa and Turkey (MENAT). , 2021, , .		0
42	Digital thermography and vascular involvement in β-thalassemia intermedia. Annals of Hematology, 2021, 100, 2471-2477.	0.8	1
43	Cancer Registration in the Middle East, North Africa, and Turkey: Scope and Challenges. JCO Global Oncology, 2021, 7, 1101-1109.	0.8	18
44	Henochâ€Schönlein purpura: Another COVIDâ€19 complication. Pediatric Dermatology, 2021, 38, 1359-1360.	0.5	19
45	Should we screen Eastern Mediterranean COVID-19 patients for inherited thrombophilia?. Medical Hypotheses, 2021, 152, 110621.	0.8	1
46	Oral ferroportin inhibitor vamifeport for improving iron homeostasis and erythropoiesis in β-thalassemia: current evidence and future clinical development. Expert Review of Hematology, 2021, 14, 633-644.	1.0	13
47	2021 update on clinical trials in βâ€ŧhalassemia. American Journal of Hematology, 2021, 96, 1518-1531.	2.0	38
48	Luspatercept for β-thalassemia: beyond red blood cell transfusions. Expert Opinion on Biological Therapy, 2021, 21, 1363-1371.	1.4	14
49	Improving outcomes and quality of life for patients with transfusion-dependent β-thalassemia: recommendations for best clinical practice and the use of novel treatment strategies. Expert Review of Hematology, 2021, 14, 897-909.	1.0	13
50	The Lebanese healthcare sector: The point of exsanguination. American Journal of Hematology, 2021, 96, E403-E404.	2.0	1
51	COVID-19 vaccination immune response in patients with solid organ and haematologic malignancies: call for active monitoring. Ecancermedicalscience, 2021, 15, 1284.	0.6	1
52	Recommendations for diagnosis and treatment of methemoglobinemia. American Journal of Hematology, 2021, 96, 1666-1678.	2.0	56
53	Prevalence of thrombophilia-associated mutations and their clinical significance in a large cohort of Lebanese patients. Meta Gene, 2021, 29, 100936.	0.3	0
54	Assessment of Lebanese healthcare professionals' awareness on acquired haemophilia: a cross-sectional study. Hematology, 2021, 26, 83-87.	0.7	0

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55	Summary of Joint European Hematology Association (EHA) and EuroBloodNet Recommendations on Diagnosis and Treatment of Methemoglobinemia. HemaSphere, 2021, 5, e660.	1.2	1
56	Luspatercept Improves Quality of Life and Reduces Red Blood Cell Transfusion Burden in Patients with Non-Transfusion-Dependent Î ² -Thalassemia in the BEYOND Trial. Blood, 2021, 138, 3081-3081.	0.6	4
57	Quality of life measures in Systemic Lupus Erythematosus: A systematic review. Reumatismo, 2021, 73, .	0.4	2
58	Effect of Perioperative Transfusion on Postoperative Morbidity Following Minimally Invasive Hysterectomy for Benign Indications. Journal of Minimally Invasive Gynecology, 2020, 27, 200-205.	0.3	7
59	The effect of anemia severity on postoperative morbidity among patients undergoing laparoscopic hysterectomy for benign indications. Acta Obstetricia Et Gynecologica Scandinavica, 2020, 99, 112-118.	1.3	12
60	No child with a transfusion-dependent haemoglobinopathy left unchelated: are we there yet?. Lancet Haematology,the, 2020, 7, e429-e430.	2.2	0
61	Overview of risk assessment models for venous thromboembolism in ambulatory patients with cancer. Thrombosis Research, 2020, 191, S50-S57.	0.8	13
62	A Report on the Education, Employment and Marital Status of Thalassemia Patients from a Tertiary Care Center in the Middle East. Hemoglobin, 2020, 44, 278-283.	0.4	4
63	Recommendations for Pregnancy in Rare Inherited Anemias. HemaSphere, 2020, 4, e446.	1.2	8
64	COVID-19 in benign hematology: emerging challenges and special considerations for healthcare professionals. Expert Review of Hematology, 2020, 13, 1081-1092.	1.0	11
65	COVID-19, Antiphospholipid Antibodies, and Catastrophic Antiphospholipid Syndrome: A Possible Association?. Clinical Medicine Insights: Arthritis and Musculoskeletal Disorders, 2020, 13, 117954412097866.	0.3	32
66	Care of patients with hemoglobin disorders during the <scp>COVID</scp> â€19 pandemic: An overview of recommendations. American Journal of Hematology, 2020, 95, E208-E210.	2.0	24
67	Beta Thalassemia: New Therapeutic Options Beyond Transfusion and Iron Chelation. Drugs, 2020, 80, 1053-1063.	4.9	49
68	A Phase 3 Trial of Luspatercept in Patients with Transfusion-Dependent β-Thalassemia. New England Journal of Medicine, 2020, 382, 1219-1231.	13.9	177
69	Acquired hemophilia A: when an overlooked autoimmune disorder causes significant bleeding. Expert Opinion on Orphan Drugs, 2020, 8, 79-89.	0.5	1
70	Thalassemia in the emergency department: special considerations for a rare disease. Annals of Hematology, 2020, 99, 1967-1977.	0.8	9
71	Less â€~reds' more â€~blues': hemoglobin level and depression in non-transfusion-dependent thalassemia. Annals of Hematology, 2020, 99, 903-904.	0.8	9
72	MERGE: A Multinational, Multicenter Observational Registry for Myeloproliferative Neoplasms in Asia, including Middle East, Turkey, and Algeria. Cancer Medicine, 2020, 9, 4512-4526.	1.3	39

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73	Emerging therapies in β-thalassemia: toward a new era in management. Expert Opinion on Emerging Drugs, 2020, 25, 113-122.	1.0	12
74	Health-Related Quality of Life Outcomes for Patients with Transfusion-Dependent Beta-Thalassemia Treated with Luspatercept in the Believe Trial. Blood, 2020, 136, 8-9.	0.6	7
75	Longitudinal Effect of Luspatercept Treatment on Iron Overload and Iron Chelation Therapy (ICT) in Adult Patients (Pts) with β-Thalassemia in the Believe Trial. Blood, 2020, 136, 47-48.	0.6	8
76	Sustained Reductions in Red Blood Cell (RBC) Transfusion Burden and Events in Î ² -Thalassemia with Luspatercept: Longitudinal Results of the Believe Trial. Blood, 2020, 136, 45-46.	0.6	8
77	Recurrent <i>Campylobacter</i> Bacteremia as the First Manifestation of Hypogammaglobulinemia: A Case Report and Literature Review. Infection and Chemotherapy, 2020, 52, 415.	1.0	7
78	Disparities in the risk of septic events in patients undergoing splenectomy for hematological malignancies (Dâ€ROSEâ€PUSH): A study based on ACSâ€NSQIP database. American Journal of Hematology, 2019, 94, E205-E207.	2.0	0
79	Quality of life in patients with βâ€thalassemia: A prospective study of transfusionâ€dependent and nonâ€transfusionâ€dependent patients in Greece, Italy, Lebanon, and Thailand. American Journal of Hematology, 2019, 94, E261-E264.	2.0	21
80	Structure and evaluation of a residency research program in a university hospital. BMC Medical Education, 2019, 19, 406.	1.0	5
81	Influence of patientâ€reported outcomes on the treatment effect of deferasirox filmâ€coated and dispersible tablet formulations in the ECLIPSE trial: A post hoc mediation analysis. American Journal of Hematology, 2019, 94, E96-E99.	2.0	5
82	Bariatric venous thromboembolism prophylaxis: an update on the literature. Expert Review of Hematology, 2019, 12, 763-771.	1.0	23
83	Thalassemia and malignancy: An emerging concern?. Blood Reviews, 2019, 37, 100585.	2.8	18
84	Development of a thalassemiaâ€related thrombosis risk scoring system. American Journal of Hematology, 2019, 94, E207-E209.	2.0	7
85	Cardiac involvement in beta-thalassaemia: current treatment strategies. Postgraduate Medicine, 2019, 131, 261-267.	0.9	13
86	EHA Research Roadmap on Hemoglobinopathies and Thalassemia: An Update. HemaSphere, 2019, 3, e208.	1.2	13
87	Development of a patientâ€reported outcomes symptom measure for patients with nontransfusionâ€dependent thalassemia (NTDTâ€PRO [©]). American Journal of Hematology, 2019, 94, 171-176.	2.0	7
88	Validation of a patientâ€reported outcomes symptom measure for patients with nontransfusionâ€dependent thalassemia (NTDTâ€PRO [©]). American Journal of Hematology, 2019, 94, 177-183.	2.0	7
89	Sofosbuvir/velpatasvir for chronic hepatitis C infection in patients with transfusionâ€dependent thalassemia. American Journal of Hematology, 2019, 94, E43-E45.	2.0	10
90	Sotatercept, a novel transforming growth factor β ligand trap, improves anemia in β-thalassemia: a phase II, open-label, dose-finding study. Haematologica, 2019, 104, 477-484.	1.7	61

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91	Predicting serum ferritin levels in patients with iron overload treated with the filmâ€coated tablet of deferasirox during the ECLIPSE study. American Journal of Hematology, 2019, 94, E15-E17.	2.0	2
92	Common fetal hemoglobin variants in Lebanese patients bearing the codon 29 beta gene mutation associated with different thalassemia phenotypes. Annals of Hematology, 2019, 98, 833-840.	0.8	0
93	Effects of Luspatercept on Iron Overload and Impact on Responders to Luspatercept: Results from the BELIEVE Trial. Blood, 2019, 134, 2245-2245.	0.6	7
94	Evaluating Luspatercept Responders in the Phase 3, Randomized, Double-Blind, Placebo-Controlled BELIEVE Trial of Luspatercept in Adult Beta-Thalassemia Patients (Pts) Who Require Regular Red Blood Cell (RBC) Transfusions. Blood, 2019, 134, 3545-3545.	0.6	3
95	<scp>MRI</scp> for the diagnosis of cardiac and liver iron overload in patients with transfusionâ€dependent thalassemia: An algorithm to guide clinical use when availability is limited. American Journal of Hematology, 2018, 93, E135-E137.	2.0	14
96	Hepatocellular Carcinoma in a β-Thalassemia Intermedia Patient: Yet Another Case in the Expanding Epidemic. Hemoglobin, 2018, 42, 58-60.	0.4	7
97	Microcytosis is important in screening of iron deficiency anemia. European Journal of Internal Medicine, 2018, 48, e39.	1.0	0
98	A paradigm shift on beta-thalassaemia treatment: How will we manage this old disease with new therapies?. Blood Reviews, 2018, 32, 300-311.	2.8	95
99	Hypercoagulability and Vascular Disease. Hematology/Oncology Clinics of North America, 2018, 32, 237-245.	0.9	20
100	Emerging Therapies. Hematology/Oncology Clinics of North America, 2018, 32, 343-352.	0.9	5
101	Clinical Complications and Their Management. Hematology/Oncology Clinics of North America, 2018, 32, 223-236.	0.9	14
102	Ineffective Erythropoiesis: Anemia and Iron Overload. Hematology/Oncology Clinics of North America, 2018, 32, 213-221.	0.9	54
103	Thalassemia. Hematology/Oncology Clinics of North America, 2018, 32, xv-xvi.	0.9	5
104	Rheumatological complications of beta-thalassaemia: an overview. Rheumatology, 2018, 57, 19-27.	0.9	16
105	Thalassaemia. Lancet, The, 2018, 391, 155-167.	6.3	512
106	Iron deficiency beyond erythropoiesis: should we be concerned?. Current Medical Research and Opinion, 2018, 34, 81-93.	0.9	83
107	Efficacy and safety of ruxolitinib in regularly transfused patients with thalassemia: results from a phase 2a study. Blood, 2018, 131, 263-265.	0.6	45
108	Iron overload and chelation therapy in hemoglobinopathies. Thalassemia Reports, 2018, 8, .	0.1	0

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109	DEFERASIROX: OVER A DECADE OF EXPERIENCE IN THALASSEMIA. Mediterranean Journal of Hematology and Infectious Diseases, 2018, 10, e2018066.	0.5	12
110	Ovarian teratoma: yet another cause of immune thrombocytopenia. International Journal of Women's Health, 2018, Volume 10, 567-570.	1.1	0
111	Hepatocellular Carcinoma in β-Thalassemia Patients: Review of the Literature with Molecular Insight into Liver Carcinogenesis. International Journal of Molecular Sciences, 2018, 19, 4070.	1.8	25
112	Patient-reported outcomes from a randomized phase II study of the deferasirox film-coated tablet in patients with transfusion-dependent anemias. Health and Quality of Life Outcomes, 2018, 16, 216.	1.0	15
113	Renal Complications in Thalassemia. Thalassemia Reports, 2018, 8, 7481.	0.1	9
114	Triple thrombophilic simultaneous mutations in patients after bariatric surgery: is there a role for screening in the Eastern Mediterranean?. Journal of Surgical Case Reports, 2018, 2018, rjy135.	0.2	0
115	How I manage medical complications of β-thalassemia in adults. Blood, 2018, 132, 1781-1791.	0.6	78
116	Assessment and study of knowledge and practice across different settings in Lebanon based on the recommendations of the American Society of Hematology <i>Choosing Wisely</i> campaign (ASKâ€PADS) Tj I	ETQq20000r	gBÐ/Overlocl
117	The use of direct oral anticoagulants in the treatment of acute venous thromboembolism in cancer patients. Expert Review of Hematology, 2018, 11, 487-494.	1.0	1
118	Drug related bleeding as a cause of emergency department visits; a Lebanese tertiary care center experience. European Journal of Internal Medicine, 2018, 55, e10-e12.	1.0	0
119	Could sodium imbalances predispose to postoperative venous thromboembolism? An analysis of the NSQIP database. Thrombosis Journal, 2018, 16, 11.	0.9	11
120	Non-Transfusion-Dependent Thalassemia: An Update on Complications and Management. International Journal of Molecular Sciences, 2018, 19, 182.	1.8	46
121	Impact and management of iron deficiency and iron deficiency anemia in women's health. Expert Review of Hematology, 2018, 11, 727-736.	1.0	70
122	Iron overload in patients with myelodysplastic syndromes: An updated overview. Cancer, 2018, 124, 3979-3989.	2.0	24
123	Optimising management of deferasirox therapy for patients with transfusionâ€dependent thalassaemia and lowerâ€risk myelodysplastic syndromes. European Journal of Haematology, 2018, 101, 272-282.	1.1	16
124	The Believe Trial: Results of a Phase 3, Randomized, Double-Blind, Placebo-Controlled Study of Luspatercept in Adult Beta-Thalassemia Patients Who Require Regular Red Blood Cell (RBC) Transfusions. Blood, 2018, 132, 163-163.	0.6	11
125	Effect of Ljpc-401 (synthetic human hepcidin) on Iron Parameters in Healthy Adults. Blood, 2018, 132, 2336-2336.	0.6	5
126	Myeloproliferative Neoplasms in Asia, Including Middle East, Turkey, and Algeria: Epidemiological Indices and Treatment Practice Patterns from the Multinational, Multicenter, Observational MERGE Registry. Blood, 2018, 132, 5461-5461.	0.6	1

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127	GDF15 and Erythroferrone Mark Erythropoietic Response to ACE-011 (Sotatercept) in Thalassemia. Blood, 2018, 132, 3633-3633.	0.6	0
128	Impact of Myeloproliferative Neoplasms (MPNs) on Health-Related Quality of Life (HRQOL) and Medical Resource Utilization: Results from the MERGE Registry. Blood, 2018, 132, 4311-4311.	0.6	4
129	Phase 2 Clinical Trial Results for Bitopertin, an Oral Glycine Transporter 1 Inhibitor, in Patients with Non-Transfusion Dependent Beta-Thalassemia. Blood, 2018, 132, 3635-3635.	0.6	2
130	Reply to Management of hepatocellular carcinoma in thalassemia and importance of the human factor. Cancer, 2017, 123, 1073-1073.	2.0	1
131	Anemia in the elderly: a consequence of aging?. Expert Review of Hematology, 2017, 10, 327-335.	1.0	57
132	Outcomes of Laparoscopic vs Open Common Bile Duct Exploration: Analysis of the NSQIP Database. Journal of the American College of Surgeons, 2017, 224, 833-840e2.	0.2	18
133	Pregnancy in βâ€thalassemia intermedia at two tertiary care centers in Lebanon and Italy: A followâ€up report on fetal and maternal outcomes. American Journal of Hematology, 2017, 92, E96-E99.	2.0	6
134	New film oated tablet formulation of deferasirox is well tolerated in patients with thalassemia or lowerâ€risk MDS: Results of the randomized, phase II ECLIPSE study. American Journal of Hematology, 2017, 92, 420-428.	2.0	66
135	Hematologic malignancies in thalassemia: Adding new cases to the repertoire. American Journal of Hematology, 2017, 92, E68-E70.	2.0	6
136	A higher prevalence of hematologic malignancies in patients with thalassemia: Background and culprits. American Journal of Hematology, 2017, 92, 414-416.	2.0	11
137	Hepatocellular carcinoma as an emerging morbidity in the thalassemia syndromes: A comprehensive review. Cancer, 2017, 123, 751-758.	2.0	54
138	Iron overload across the spectrum of nonâ€ŧransfusionâ€dependent thalassaemias: role of erythropoiesis, splenectomy and transfusions. British Journal of Haematology, 2017, 176, 288-299.	1.2	43
139	Clinical management of iron deficiency anemia in adults: Systemic review on advances in diagnosis and treatment. European Journal of Internal Medicine, 2017, 42, 16-23.	1.0	76
140	A Predictive Score for Thrombosis Associated with Breast, Colorectal, Lung, or Ovarian Cancer: The Prospective COMPASS–Cancer-Associated Thrombosis Study. Oncologist, 2017, 22, 1222-1231.	1.9	167
141	Revisiting beta thalassemia intermedia: past, present, and future prospects. Hematology, 2017, 22, 607-616.	0.7	19
142	Iron deficiency across chronic inflammatory conditions: International expert opinion on definition, diagnosis, and management. American Journal of Hematology, 2017, 92, 1068-1078.	2.0	290
143	Circulating microparticles and the risk of thromboembolic events in Egyptian beta thalassemia patients. Annals of Hematology, 2017, 96, 597-603.	0.8	9
144	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.	2.0	13

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145	Safety and pharmacokinetics of the oral iron chelator SPâ€420 in βâ€thalassemia. American Journal of Hematology, 2017, 92, 1356-1361.	2.0	10
146	Causes of hospital admission in βâ€ŧhalassemia (CHAT) in Lebanon from 1995 to 2015: A pilot retrospective study from a tertiary care center. American Journal of Hematology, 2017, 92, E652-E653.	2.0	1
147	Novel mutation in the Transferrin receptor-2 in a patient with Hereditary Hemochromatosis type 3. Meta Gene, 2017, 14, 30-32.	0.3	1
148	Insights into the diagnosis and management of iron deficiency in inflammatory bowel disease. Expert Review of Hematology, 2017, 10, 801-808.	1.0	17
149	Low utility of lower extremity ultrasound prior to application of sequential compression device in critically ill adults. Vascular Medicine, 2017, 22, 66-68.	0.8	0
150	Clinical utility of serum ferritin thresholds for guiding iron chelation therapy when magnetic resonance imaging is unavailable in patients with nonâ€ŧransfusionâ€dependent thalassaemias – response to Ang <i>etÂal</i> . British Journal of Haematology, 2017, 176, 989-990.	1.2	1
151	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.	0.8	24
152	Limitations of serum ferritin to predict liver iron concentration responses to deferasirox therapy in patients with transfusionâ€dependent thalassaemia. European Journal of Haematology, 2017, 98, 280-288.	1.1	29
153	Enhancing Effect of Hydroxyurea on Hb F in Sickle Cell Disease: Ten-Year Egyptian Experience. Hemoglobin, 2017, 41, 267-273.	0.4	6
154	Iron Overload and Chelation Therapy in Non-Transfusion Dependent Thalassemia. International Journal of Molecular Sciences, 2017, 18, 2778.	1.8	20
155	Iron overload in thalassemia: different organs at different rates. Hematology American Society of Hematology Education Program, 2017, 2017, 265-271.	0.9	167
156	Quality of Life in Patients with Î ² -Thalassemia: Transfusion Dependent Versus Non-Transfusion Dependent. Blood, 2017, 130, 751-751.	0.6	1
157	Efficacy and safety of iron-chelation therapy with deferoxamine, deferiprone, and deferasirox for the treatment of iron-loaded patients with nontransfusion-dependent thalassemia syndromes. Drug Design, Development and Therapy, 2016, Volume 10, 4073-4078.	2.0	15
158	Deferasirox: an orphan drug for chronic iron overload in non-transfusion dependent thalassemia syndromes. Expert Opinion on Orphan Drugs, 2016, 4, 677-686.	0.5	2
159	Morbidities in nonâ€transfusionâ€dependent thalassemia. Annals of the New York Academy of Sciences, 2016, 1368, 82-94.	1.8	20
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