Sezaneh Haghpanah

List of Publications by Year
in descending order

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		394421	501196
172	1,513	19	28
papers	citations	h-index	g-index

185	185	185	2237
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Comparison of post-urethroplasty complication rates in pediatric cases with hypospadias using Vicryl or polydioxanone sutures. Asian Journal of Urology, 2022, 9, 165-169.	1.2	6
2	Comparative Effectiveness of Peg-Asparaginase and L-Asparaginase on Coagulation Markers Among Pediatric Patients with Acute Lymphoblastic Leukemia. Clinical Laboratory, 2022, 68, .	0.5	0
3	EFFICACY AND SAFETY OF SINOPHARM VACCINE FOR SARS-COV-2 AND BREAKTHROUGH INFECTIONS IN IRANIAN PATIENTS WITH HEMOGLOBINOPATHIES: A PRELIMINARY REPORT. Mediterranean Journal of Hematology and Infectious Diseases, 2022, 14, e2022026.	1.3	8
4	Oncogenic and tumor suppressor genes expression in myeloproliferative neoplasms: The hidden side of a complex pathology. Journal of Clinical Laboratory Analysis, 2022, 36, e24289.	2.1	1
5	Atopy manifestations in pediatric patients with acute lymphoblastic leukemia: correlation assessment with interleukin-4 (IL-4) and IgE level. BMC Pediatrics, 2022, 22, 149.	1.7	1
6	Effects of three months of treatment with vitamin E and N-acetyl cysteine on the oxidative balance in patients with transfusion-dependent Î²-thalassemia. Annals of Hematology, 2021, 100, 635-644.	1.8	4
7	Bayesian spatial modeling of transfusion-dependent Î²-thalassemia incidence rate in Fars Province, Southern Iran. Spatial and Spatio-temporal Epidemiology, 2021, 36, 100389.	1.7	2
8	Study of the Serum Immunoglobulin and Cell-Mediated Immunity in Patients with Congenital Severe Hemophilia. Clinical Laboratory, 2021, 67, .	0.5	0
9	A cost-effectiveness analysis of the prophylaxis versus on-demand regimens in severe hemophilia A patients under 12 years old in southern Iran. Hematology, 2021, 26, 240-248.	1.5	1
10	Blood Transfusion Practice in Operating Rooms in Nemazee Hospital in Southern Iran. Archives of Iranian Medicine, 2021, 24, 107-112.	0.6	4
11	Trace Elements in Children with Acute Lymphoblastic Leukemia. Asian Pacific Journal of Cancer Prevention, 2021, 22, 43-47.	1.2	9
12	A cost-analysis study of using adult red cell packs and Pedi-Packs in newborn intensive care units in Southern Iran. Cost Effectiveness and Resource Allocation, 2021, 19, 15.	1.5	1
13	Incidence Rate of COVID-19 Infection in Hemoglobinopathies: A Systematic Review and Meta-analysis. Hemoglobin, 2021, 45, 371-379.	0.8	12
14	Impact of antifungal stewardship interventions on the susceptibility of colonized Candida species in pediatric patients with malignancy. Scientific Reports, 2021, 11, 14099.	3.3	6
15	Comparison of the clinical features and outcome of children with hemophagocytic lymphohistiocytosis (HLH) secondary to visceral leishmaniasis and primary HLH: a single-center study. BMC Infectious Diseases, 2021, 21, 732.	2.9	9
16	Epidemiologic study of patients with thrombotic events referred to a tertiary hospital in Southern Iran. Heliyon, 2021, 7, e07734.	3.2	0
17	Long-term safety and efficacy of hydroxyurea in patients with non-transfusion-dependent Î²-thalassemia: a comprehensive single-center experience. Annals of Hematology, 2021, 100, 2901-2907.	1.8	5
18	Strategies for improvement of blood consumption management in the operating rooms: expertsÂ´ suggestions. Journal of Community Hospital Internal Medicine Perspectives, 2021, 11, 635-638.	0.8	2

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19	The Outcome of Children With Malignant Bone Tumors: A Single-Center Experience. Global Pediatric Health, 2021, 8, 2333794X2110422.	0.7	0
20	Association of HFE Gene Mutations With Serum Ferritin Level and Heart and Liver Iron Overload in Patients With Transfusion-dependent Beta-Thalassemia. Journal of Pediatric Hematology/Oncology, 2021, 43, e26-e28.	0.6	3
21	Coronavirus disease 2019 (COVID-19) severity in patients with thalassemias: A Nationwide Iranian Experience. Mediterranean Journal of Hematology and Infectious Diseases, 2021, 13, e2021008.	1.3	12
22	A Proposed Managerial Model for Improvement of Blood Consumption in the Operating Rooms in Southern Iran. Health Scope, 2021, 10, .	0.6	0
23	The Prevalence of Hypothyroidism among Patients With Î²-Thalassemia: A Systematic Review and Meta-Analysis of Cross-Sectional Studies. Hemoglobin, 2021, 45, 275-286.	0.8	4
24	Investigating Trends of Incidence Rates of Esophageal Cancer Divided by Squamous Cell Carcinoma and Adenocarcinoma in Southern Iran: a 10-Year Experience. Journal of Gastrointestinal Cancer, 2021, , 1.	1.3	2
25	Evaluation of endocrine complications in beta-thalassemia intermedia (Î²-TI): a cross-sectional multicenter study. Endocrine, 2020, 69, 220-227.	2.3	8
26	Evaluation of Efficacy, Safety, and Satisfaction Taking Deferasirox Twice Daily Versus Once Daily in Patients With Transfusion-Dependent Thalassemia. Journal of Pediatric Hematology/Oncology, 2020, 42, 23-26.	0.6	7
27	Ocular findings in patients with transfusion-dependent Î²-thalassemia in southern Iran. BMC Ophthalmology, 2020, 20, 376.	1.4	4
28	Association of Exon 14 of the SOX6 Gene Sequence Variations with Response to Hydroxyurea Therapy in Patients Carrying Non Transfusion-Dependent Thalassemia. Hemoglobin, 2020, 44, 406-410.	0.8	1
29	Bone mineral density in transfusion-dependent thalassemia patients and its associated factors in Southern Iran. Archives of Osteoporosis, 2020, 15, 148.	2.4	2
30	Prevalence and mortality in Î²â€thalassaemias due to outbreak of novel coronavirus disease (COVIDâ€19): the nationwide Iranian experience. British Journal of Haematology, 2020, 190, e137-e140.	2.5	35
31	Parameters of tissue iron overload and cardiac function in patients with thalassemia major and intermedia. Acta Haematologica Polonica, 2020, 51, 95-101.	0.3	2
32	Prevalence of Low Bone Mass in Patients with Hemophilia and Its Related Ractors in Southern Iran. Journal of Comprehensive Pediatrics, 2020, 11, .	0.3	1
33	Prevalence and clinical features of COVID-19 in Iranian patients with congenital coagulation disorders. Blood Transfusion, 2020, 18, 413-414.	0.4	2
34	Prevalence and severity of Coronavirus disease 2019 (COVID-19) in Transfusion Dependent and Non-Transfusion Dependent Î²-thalassemia patients and effects of associated comorbidities: an Iranian nationwide study. Acta Biomedica, 2020, 91, e2020007.	0.3	5
35	Extramedullary manifestations in acute lymphoblastic leukemia in children: a systematic review and guideline-based approach of treatment. American Journal of Blood Research, 2020, 10, 360-374.	0.6	4
36	Frequency of silent brain lesions and aspirin protection evaluation over 3Âyears follow-up in beta thalassemia patients. Annals of Hematology, 2019, 98, 2267-2271.	1.8	7

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37	Correlation of bleeding score with frequency and severity of bleeding symptoms in FXIII deficiency assessing by the ISTH Bleeding Assessment Tool. Transfusion and Apheresis Science, 2019, 58, 495-497.	1.0	4
38	Changing face of Candida colonization pattern in pediatric patients with hematological malignancy during repeated hospitalizations, results of a prospective observational study (2016â€“2017) in shiraz, Iran. BMC Infectious Diseases, 2019, 19, 759.	2.9	17
39	Effect of different iron chelation regimens on bone mass in transfusion-dependent thalassemia patients. Expert Review of Hematology, 2019, 12, 997-1003.	2.2	12
40	Prevalence of endocrine disorders and their associated factors in transfusion-dependent thalassemia patients: a historical cohort study in Southern Iran. Journal of Endocrinological Investigation, 2019, 42, 1467-1476.	3.3	20
41	A retrospective study on clinical manifestations of neonates with FXIII-A deficiency. Blood Cells, Molecules, and Diseases, 2019, 77, 78-81.	1.4	4
42	Comparative effectiveness of alendronate and zoledronic acid on bone mass improvement in transfusion-dependent thalassemia patients. Journal of Bone and Mineral Metabolism, 2019, 37, 996-1003.	2.7	4
43	Evaluation of intima-media thickness of common carotid artery in thalassemia patients compared to healthy individuals. Pars of Jahrom University of Medical Sciences, 2019, 17, 1-7.	0.1	0
44	Modified Primary Prophylaxis in Previously Untreated Patients With Severe Hemophilia A in Iran. Journal of Pediatric Hematology/Oncology, 2018, 40, 188-191.	0.6	5
45	Investigating the bone mineral density in children with solid tumors in southern Iran: a caseâ€“control study. Archives of Osteoporosis, 2018, 13, 8.	2.4	4
46	The Zinc and Copper Levels in Thalassemia Major Patients, Receiving Iron Chelation Therapy. Journal of Pediatric Hematology/Oncology, 2018, 40, 178-181.	0.6	8
47	Complementary and alternative medicine use in thalassemia patients in Shiraz, southern Iran: A cross-sectional study. Journal of Traditional and Complementary Medicine, 2018, 8, 141-146.	2.7	15
48	Blood transfusion versus hydroxyurea in beta-thalassemia in Iran: a cost-effectiveness study. Hematology, 2018, 23, 417-422.	1.5	3
49	Expression of antiapoptotic proteins livin and survivin in pediatric AML patients, as prognostic markers. Pediatric Hematology and Oncology, 2018, 35, 250-256.	0.8	7
50	Relationship of the Interaction Between Two Quantitative Trait Loci with Î³-Globin Expression in Î²-Thalassemia Intermedia Patients. Hemoglobin, 2018, 42, 108-112.	0.8	1
51	Efficacy and safety of resveratrol, an oral hemoglobin F-augmenting agent, in patients with beta-thalassemia intermedia. Annals of Hematology, 2018, 97, 1919-1924.	1.8	8
52	Effect of ursodeoxycholic acid and vitamin E in the prevention of liver injury from methotrexate in pediatric leukemia. Turkish Journal of Gastroenterology, 2018, 29, 203-209.	1.1	21
53	Evaluation of Endocrine Complications in Beta-Thalassemia Intermedia Patients: A Cross Sectional Multi-Center Study. Blood, 2018, 132, 2343-2343.	1.4	1
54	The frequency of hypothyroidism and its relationship with HCV positivity in patients with thalassemia major in southern Iran. Acta Biomedica, 2018, 89, 55-60.	0.3	2

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55	Quality of Life in Children and Adolescents With Rare Bleeding Disorders in Southern Iran. Clinical and Applied Thrombosis/Hemostasis, 2017, 23, 652-656.	1.7	7
56	Relationship Between Some Single-nucleotide Polymorphism and Response to Hydroxyurea Therapy in Iranian Patients With Î²-Thalassemia Intermedia. Journal of Pediatric Hematology/Oncology, 2017, 39, e171-e176.	0.6	9
57	A cross-sectional study of complementary and alternative medicine use in patients with coagulation disorders in Southern Iran. Journal of Integrative Medicine, 2017, 15, 359-364.	3.1	3
58	A large case series on surgical outcomes in congenital factor XIII deficiency patients in Iran. Journal of Thrombosis and Haemostasis, 2017, 15, 2300-2305.	3.8	4
59	Spectrum of pediatric tumors diagnosed by fine-needle aspiration cytology. Medicine (United States), 2017, 96, e5480.	1.0	6
60	Evaluation of Proteinuria in Î²-Thalassemia Major Patients With and Without Diabetes Mellitus Taking Deferasirox. Journal of Pediatric Hematology/Oncology, 2017, 39, e11-e14.	0.6	3
61	Evaluation of bone mineral density in children with sickle-cell anemia and its associated factors in the south of Iran: a case-control study. Archives of Osteoporosis, 2017, 12, 70.	2.4	8
62	Correlation of serum ferritin levels with hepatic MRI T2 and liver iron concentration in nontransfusion beta-thalassemia intermediate patients: A contemporary issue. Pediatric Hematology and Oncology, 2017, 34, 292-297.	0.8	13
63	Introduction of novel Î±₁-hemoglobin gene mutation with transfusion-dependent phenotype. Hematology, 2017, 22, 168-171.	1.5	2
64	A comparison of heart function and arrhythmia in clinically asymptomatic patients with beta thalassemia intermedia and beta thalassemia major. Hematology, 2017, 22, 25-29.	1.5	10
65	A 25-year surveillance of disseminated Bacillus Calmetteâ€“GuÃ©rin disease treatment in children in Southern Iran. Medicine (United States), 2017, 96, e9035.	1.0	11
66	Transcranial Doppler Screening in 50 Patients With Sickle Cell Hemoglobinopathies in Iran. Journal of Pediatric Hematology/Oncology, 2017, 39, 506-512.	0.6	4
67	FMS-like Tyrosine Kinase 3 (FLT3) and Nucleophosmin 1 (NPM1) in Iranian Adult Acute Myeloid Leukemia (AML) Patients with Normal Karyotype; Mutation Status and Clinical and Laboratory Characteristics. Turkish Journal of Haematology, 2017, 34, 300-306.	0.5	14
68	Acquired Vitamin K Deficiency as Unusual Cause of Bleeding Tendency in Adults: A Case Report of a Nonhospitalized Student Presenting with Severe Menorrhagia. Case Reports in Obstetrics and Gynecology, 2017, 2017, 1-3.	0.3	4
69	THE EFFECTS OF OLIVE LEAF EXTRACT OINTMENT ON PAIN INTENSITY AND EARLY MATERNAL COMPLICATIONS IN PRIMIPAROUS WOMEN. International Journal of Pharmacy and Pharmaceutical Sciences, 2017, 9, 31.	0.3	10
70	Comparison of Quality of Life in Patients with Î²-Thalassemia Intermedia and Î²-Thalassemia Major in Southern Iran. Hemoglobin, 2017, 41, 169-174.	0.8	12
71	Evaluation of Metabolic Syndrome and Related Factors in Children Affected by Acute Lymphoblastic Leukemia. Indian Journal of Medical and Paediatric Oncology, 2017, 38, 97-102.	0.2	4
72	Use of Complementary and Alternative Medicine Among Iranian Cancer Patients in South of Iran. International Journal of Cancer Management, 2017, 10, .	0.4	8

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73	Phenotype Report on Patients with Congenital Factor V Deficiency in Southern Iran in the recent ten years experience. Turkish Journal of Haematology, 2017, 34, 250-253.	0.5	1
74	Epidemiology of Hereditary Coagulation Bleeding Disorders: A 15-Year Experience From Southern Iran. Hospital Practices and Research, 2017, 2, 113-117.	0.2	1
75	Correlation between Rs2108622 Locus of CYP4F2 Gene Single Nucleotide Polymorphism and Warfarin Dosage in Iranian Cardiovascular Patients. Iranian Journal of Pharmaceutical Research, 2017, 16, 1238-1246.	0.5	1
76	Evaluation of Plasma Platelet Microparticles in Thrombotic Thrombocytopenic Purpura. Annals of Clinical and Laboratory Science, 2017, 47, 62-67.	0.2	4
77	Evaluation of Thrombin Generation Assay in Patients With Hemophilia. Clinical and Applied Thrombosis/Hemostasis, 2016, 22, 322-326.	1.7	11
78	Evaluation of Knowledge of Patients with Hemophilia Regarding Their Diseases and Treatment in Iran. Turkish Journal of Haematology, 2016, 33, 355-356.	0.5	2
79	A comparison between MRI, sonography and Functional Independence Score in Haemophilia methods in diagnosis, evaluation and classification of arthropathy in severe haemophilia A and B. Blood Coagulation and Fibrinolysis, 2016, 27, 131-135.	1.0	5
80	Efficacy and safety of factor eight inhibitor bypassing activity prophylaxis evaluation in young patients with hemophilia and high titer inhibitor. Blood Coagulation and Fibrinolysis, 2016, 27, 232-233.	1.0	0
81	Wound Healing Studies Using Punica granatum Peel. Advances in Skin and Wound Care, 2016, 29, 217-225.	1.0	15
82	Current strategies against invasive fungal infections in patients with aplastic anemia, strong power and weak weapon, a case report and review of literature. Medical Mycology Case Reports, 2016, 11, 16-20.	1.3	3
83	Evaluation of the Relationship Between Hb F Levels and Nucleated Red Blood Cells with Morbidity in Non Transfusion-Dependent Thalassemia Patients. Hemoglobin, 2016, 40, 250-256.	0.8	3
84	Frequency of silent cerebral ischemia in patients with transfusion-dependent Î²-thalassemia major compared to healthy individuals. Annals of Hematology, 2016, 95, 1387-1387.	1.8	6
85	Bone mineral density in children with acute leukemia and its associated factors in Iran: a case-control study. Archives of Osteoporosis, 2016, 11, 36.	2.4	7
86	Evaluation of bone mineral density in patients with hemoglobin H disease. Annals of Hematology, 2016, 95, 1329-1332.	1.8	2
87	The frequency of silent cerebral ischemia in patients with transfusion-dependent Î²-thalassemia major. Annals of Hematology, 2016, 95, 135-139.	1.8	18
88	Prevalence and Prognostic Impact of Wilms' Tumor 1 (WT1) Gene, Including SNP rs16754 in Cytogenetically Normal Acute Myeloblastic Leukemia (CN-AML): An Iranian Experience. Clinical Lymphoma, Myeloma and Leukemia, 2016, 16, e21-e26.	0.4	8
89	Correlation of Serum Ferritin Levels with Liver and Heart Mri T2 and Liver Iron Concentration in Beta Thalassemia Intermediate Patients: A Contemporary Issue. Blood, 2016, 128, 4829-4829.	1.4	1
90	Impact of Education on Awareness Towards Reproductive Health in Women With Beta-Thalassemia Major. Women's Health Bulletin, 2016, In Press, .	0.7	0

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91	Impact of Education on Awareness Towards Reproductive Health in Women With Beta-Thalassemia Major. Women's Health Bulletin, 2016, 4, .	0.7	0
92	Impact of clinical supervision on field training of nursing students at Urmia University of Medical Sciences. Journal of Advances in Medical Education and Professionalism, 2016, 4, 88-92.	0.2	4
93	Hemoaction Game: An educational Step to Improve Hemophilia Children and Nurses Self-Efficacy. Journal of Advances in Medical Education and Professionalism, 2016, 4, 206.	0.2	1
94	A randomised controlled trial of oral zinc sulphate for primary dysmenorrhoea in adolescent females. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2015, 55, 369-373.	1.0	16
95	Relationship Between Serum Hepcidin and Ferritin Levels in Patients With Thalassemia Major and Intermedia in Southern Iran. Iranian Red Crescent Medical Journal, 2015, 17, e28343.	0.5	16
96	Incidence of testicular microlithiasis in patients with Î²-thalassemia major. Annals of Hematology, 2015, 94, 1785-1789.	1.8	3
97	The effects of economic sanctions on disease specific clinical outcomes of patients with thalassemia and hemophilia in Iran. Health Policy, 2015, 119, 239-243.	3.0	29
98	Distribution of alpha-thalassemia mutations in Iranian population. Hematology, 2015, 20, 359-362.	1.5	23
99	Efficacy of Deferasirox (ExjadeÂ®) in Modulation of Iron Overload in Patients withÎ²-Thalassemia Intermedia. Hemoglobin, 2015, 39, 327-329.	0.8	9
100	Combination therapyÂâ€“Âdeferasirox and deferoxamineÂâ€“Âin thalassemia major patients in emerging countries with limited resources. Transfusion Medicine, 2015, 25, 8-12.	1.1	17
101	Serum Ferritin Levels Correlation With Heart and Liver MRI and LIC in Patients With Transfusion-Dependent Thalassemia. Iranian Red Crescent Medical Journal, 2015, 17, e24959.	0.5	37
102	Comparative Study of Radiographic and Laboratory Findings Between Beta Thalassemia Major and Beta Thalassemia Intermedia Patients With and Without Treatment by Hydroxyurea. Iranian Red Crescent Medical Journal, 2015, 17, e23607.	0.5	4
103	Quality of Life in Children with Rare Bleeding Disorders. Blood, 2015, 126, 4698-4698.	1.4	0
104	THE FREQUENCY OF ADRENAL INSUFFICIENCY IN ADOLESCENTS AND YOUNG ADULTS WITH THALASSEMIA MAJOR VERSUS THALASSEMIA INTERMEDIA IN IRAN. Mediterranean Journal of Hematology and Infectious Diseases, 2014, 7, e2015005.	1.3	10
105	Compliance and satisfaction with deferasirox (Exjade^{Â®}) compared with deferoxamine in patients with transfusion-dependent beta-thalassemia. Hematology, 2014, 19, 187-191.	1.5	24
106	Genotype and phenotype report on patients with combined deficiency of factor V and factor VIII in Iran. Blood Coagulation and Fibrinolysis, 2014, 25, 360-363.	1.0	14
107	Predictors of excessive renal displacement during access in percutaneous nephrolithotomy: a randomized clinical trial. Urolithiasis, 2014, 42, 61-65.	2.0	9
108	Intracranial hemorrhage pattern in the patients with factor XIII deficiency. Annals of Hematology, 2014, 93, 693-697.	1.8	21

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109	An experience of using Traumastem P in control of spontaneous nose bleeding in patients with inherited bleeding disorders in southern <scp>I</scp>ran. Haemophilia, 2014, 20, e79-80.	2.1	3
110	Comparison of Thrombin Generation Assay With Conventional Coagulation Tests in Evaluation of Bleeding Risk in Patients With Rare Bleeding Disorders. Clinical and Applied Thrombosis/Hemostasis, 2014, 20, 637-644.	1.7	20
111	Epidemiology of Hemoglobinopathies and Thalassemias in Individuals Referred to the Haematology Research Centre, Shiraz University of Medical Sciences, Shiraz, Iran From 2006 to 2011. Hemoglobin, 2014, 38, 287-288.	0.8	2
112	Frequency of Cholelithiasis in Patients With Beta-Thalassemia Intermedia With and Without Hydroxyurea. Iranian Red Crescent Medical Journal, 2014, 16, e18712.	0.5	6
113	Hydroxyurea Treatment in Transfusion-Dependent Î²-Thalassemia Patients. Iranian Red Crescent Medical Journal, 2014, 16, e18028.	0.5	24
114	A Comparison Between MRI, Sonography and FISH Methods in Diagnosis, Evaluation and Classification of Arthropathy in Severe Haemophilia a and B. Blood, 2014, 124, 5035-5035.	1.4	0
115	Evaluation of Thrombin Generation Assay in Patients with Hemophilia. Blood, 2014, 124, 5061-5061.	1.4	0
116	Hypothyroidism in Î²-Thalassemia Intermedia Patients with and without Hydroxyurea. Iranian Journal of Medical Sciences, 2014, 39, 60-3.	0.4	5
117	Serum cancer antigen 15.3 concentrations in patients with betathalassemia minor compared to those with cancer and healthy individuals. Medical Journal of the Islamic Republic of Iran, 2014, 28, 91.	0.9	1
118	Percutaneous nephrolithotomy: is distilled water as safe as saline for irrigation?. Urology Journal, 2014, 11, 1551-6.	0.4	5
119	Cerebral Artery Velocity Determined by Transcranial Doppler Ultrasonography in Patients With Î²-Thalassemia Intermedia Compared to Î²-Thalassemia Major. Clinical and Applied Thrombosis/Hemostasis, 2013, 19, 367-373.	1.7	1
120	Experience on Using Prothrombin Complex Concentrate in Urgent Warfarin Reversal. Clinical and Applied Thrombosis/Hemostasis, 2013, 19, 277-281.	1.7	4
121	Serological investigation for hepatitis E virus infection in the patients with chronic maintenance hemodialysis from southwest of Iran. Asian Journal of Transfusion Science, 2013, 7, 21.	0.3	14
122	Family Planning Practices in Families with Children Affected by Î²-Thalassemia Major in Southern Iran. Hemoglobin, 2013, 37, 74-79.	0.8	3
123	Attitudes and practices with regard to circumcision in haemophilia patients in Southern Iran. Haemophilia, 2013, 19, e177-8.	2.1	2
124	Evaluation of the FXIII deficiency prophylaxis intervals in large number of FXIII deficiency patients from Iran. Haemophilia, 2013, 19, e175-6.	2.1	5
125	Frequency of combined factor V and factor VIII deficiency in southern Iran. Blood Coagulation and Fibrinolysis, 2013, 24, 458-459.	1.0	1
126	Quality of life among Iranian patients with beta-thalassemia major using the SF-36 questionnaire. Sao Paulo Medical Journal, 2013, 131, 166-172.	0.9	33

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127	Inherited Thrombophilia and Recurrent Pregnancy Loss. Iranian Red Crescent Medical Journal, 2013, 15, e13708.	0.5	20
128	Management of Bleeding in Post-liver Disease, Surgery and Biopsy in Patients With High Uncorrected International Normalized Ratio With Prothrombin Complex Concentrate: An Iranian Experience. Iranian Red Crescent Medical Journal, 2013, 15, e12260.	0.5	2
129	Is zinc an essential factor in maternal health status and fetal and neonatal growth?. Trace Elements and Electrolytes, 2012, 29, 239-245.	0.1	1
130	Sociocultural challenges of beta-thalassaemia major birth in carriers of beta-thalassaemia in Iran. Journal of Medical Screening, 2012, 19, 109-111.	2.3	10
131	Î²-Thalassemia: New Therapeutic Modalities, Genetics, Complications, and Quality of Life. Anemia, 2012, 2012, 1-1.	1.7	38
132	Comparative study of hypogonadism in beta-thalassemia intermedia patients with and without hydroxyurea. Hematology, 2012, 17, 122-124.	1.5	6
133	Iranian experience of deferasirox (Exjade^{Â®}) in transfusion-dependent patients with iron overload: what is the most effective dose based on serum ferritin levels?. Hematology, 2012, 17, 367-371.	1.5	5
134	Cerebral thrombosis in patients with Î²-thalassemia. Blood Coagulation and Fibrinolysis, 2012, 23, 212-217.	1.0	31
135	Genotypeâ€“phenotype correlation related to lipid profile in beta-thalassemia major and intermedia in southern Iran. Journal of Clinical Lipidology, 2012, 6, 108-113.	1.5	7
136	Thyroid function and stress hormones in children with stress hyperglycemia. Endocrine, 2012, 42, 653-657.	2.3	4
137	Frequency and distribution of asymptomatic brain lesions in patients with Î²-thalassemia intermedia. Annals of Hematology, 2012, 91, 1833-1838.	1.8	18
138	Attitudes of haemophilic patients towards their health and socioâ€economic problems in Iran. Haemophilia, 2012, 18, 122-128.	2.1	9
139	Efficacy of prophylaxis and genotypeâ€phenotype correlation in patients with severe Factor X deficiency in Iran. Haemophilia, 2012, 18, 211-215.	2.1	34
140	Polymorphisms associated with sickle cell disease in Southern Iran. Russian Journal of Genetics, 2012, 48, 755-757.	0.6	0
141	Evaluation of Health Related Quality of Life in 6â€“18 Years Old Patients with Acute Leukemia during Chemotherapy. Indian Journal of Pediatrics, 2012, 79, 177-182.	0.8	13
142	Genotypeâ€“phenotype relationship of patients with Î²-thalassemia taking hydroxyurea: a 13-year experience in Iran. International Journal of Hematology, 2012, 95, 51-56.	1.6	50
143	Frequency and Distribution of Asymptomatic Brain Lesions in Patients with Î²-Thalassemia Intermedia. Blood, 2012, 120, 5141-5141.	1.4	0
144	Hemorrhagic symptoms and bleeding risk in obligatory carriers of type 3 von Willebrand disease in southern Iran. Blood Coagulation and Fibrinolysis, 2011, 22, 325-330.	1.0	1

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145	The correlation between gene mutations and inhibitor development in patients with haemophilia A in southern Iran. Haemophilia, 2011, 17, 820-821.	2.1	2
146	Erectile dysfunction among hemodialysis patients. International Urology and Nephrology, 2011, 43, 117-123.	1.4	16
147	Use of Glubran 2 and Glubran tissue skin adhesive in patients with hereditary bleeding disorders undergoing circumcision and dental extraction. Annals of Hematology, 2011, 90, 463-468.	1.8	12
148	Protein C concentration in newborn infants with sepsis-like illness. Journal of Neonatal-Perinatal Medicine, 2011, 4, 55-58.	0.8	0
149	Oral health status in Iranian hemodialysis patients. Indian Journal of Nephrology, 2011, 21, 235.	0.5	9
150	Cerebral Artery Velocity in Patients with Î²-Thalassemia Intermedia Compared to Î²-Thalassemia Major Using Transcranial Doppler Ultrasonography,. Blood, 2011, 118, 3335-3335.	1.4	0
151	Epidemiology of erectile dysfunction in hemodialysis patients using IIEF questionnaire. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2011, 22, 232-6.	0.3	11
152	Effect of combination therapy of hydroxyurea with <scp>l</scp>â€carnitine and magnesium chloride on hematologic parameters and cardiac function of patients with Î²â€thalassemia intermedia. European Journal of Haematology, 2010, 84, 52-58.	2.2	22
153	ADVERSE EFFECTS OF HYDROXYUREA IN Î²-THALASSEMIA INTERMEDIA PATIENTS: 10 Yearsâ€™ Experience. Pediatric Hematology and Oncology, 2010, 27, 205-211.	0.8	33
154	Long Term Use of Hydroxyurea In Patients with Î²-Thalassemia In Southern Iran. Blood, 2010, 116, 2061-2061.	1.4	7
155	Serum lipid profiles in patients with beta-thalassemia major and intermedia in southern Iran. Journal of Research in Medical Sciences, 2010, 15, 150-4.	0.9	17
156	Effect of intralesional verapamil for treatment of Peyronieâ€™s disease: a randomized single-blind, placebo-controlled study. International Urology and Nephrology, 2009, 41, 467-471.	1.4	110
157	Application of amniotic membrane as xenograft for urethroplasty in rabbit. International Urology and Nephrology, 2009, 41, 895-901.	1.4	26
158	Is red cell from an otherwise healthy G6PD-deficient donor efficient for transfusion to fauvism patients?. Indian Journal of Hematology and Blood Transfusion, 2009, 25, 23-26.	0.6	1
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