Hassan Mansouri Tourghabeh

List of Publications by Year
in descending order

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42 papers	618 citations	840585 11 h-index	610775 24 g-index
42 all docs	42 docs citations	42 times ranked	615 citing authors

#	Article	IF	Citations
1	High levels of Von Willebrand factor markers in COVID-19: a systematic review and meta-analysis. Clinical and Experimental Medicine, 2022, 22, 347-357.	1.9	23
2	Elevated levels of <scp>C3</scp>, <scp>C4</scp>, and <scp>CH50</scp> of the complement system in <scp>ICU</scp> and <scp>nonâ€ICU</scp> patients with <scp>COVID</scp>â€19. Health Science Reports, 2022, 5, e519.	0.6	7
3	Female case with misdiagnosis of hemophilia A who underwent total knee arthroplasty: A case report. Clinical Case Reports (discontinued), 2022, 10, e05558.	0.2	0
4	Consanguineous marriage and rare bleeding disorders. Expert Review of Hematology, 2021, 14, 467-472.	1.0	6
5	Parvovirus 4 in Individuals with Severe Hemophilia A and Matched Control Group. International Journal of Hematology-Oncology and Stem Cell Research, 2021, 15, 192-198.	0.3	0
6	Determining causes of death among individuals with haemophilia in Afghanistan. Haemophilia, 2021, , .	1.0	0
7	Hemostatic System (Fibrinogen Level, D-Dimer, and FDP) in Severe and Non-Severe Patients With COVID-19: A Systematic Review and Meta-Analysis. Clinical and Applied Thrombosis/Hemostasis, 2021, 27, 107602962110109.	0.7	14
8	Current methods of measuring platelet activity: pros and cons. Blood Coagulation and Fibrinolysis, 2020, 31, 426-433.	0.5	6
9	Clinical Care of Bone Health in Patients on the Immune Tolerance Inductionâ€™s Protocols With an Immunosuppressive Agent for Inhibitor Eradication in Hemophilia. Clinical and Applied Thrombosis/Hemostasis, 2020, 26, 107602962091395.	0.7	3
10	D-dimer level in COVID-19 infection: a systematic review. Expert Review of Hematology, 2020, 13, 1265-1275.	1.0	242
11	Mining of mortality-related findings in rare bleeding disorders: a retrospective study from two centers. Blood Research, 2020, 55, 213-216.	0.5	4
12	Is the Detection of Factor IX Inhibitors in Hemophilia B Orphan than Factor VIII Inhibitors in Hemophilia A? A Concise, Systematic Review. Cardiovascular & Hematological Disorders Drug Targets, 2020, 20, 185-190.	0.2	1
13	Hemostasis critical values among Iranian clinical laboratories â€œNational Survey of 157 Clinical Laboratoriesâ€. International Journal of Laboratory Hematology, 2019, 41, 778-781.	0.7	1
14	The Most Common Allergenic Tree Pollen Grains in the Middle East: A Narrative Review. Iranian Journal of Medical Sciences, 2019, 44, 87-98.	0.3	6
15	Is It Rational to Study Coagulations Test Routinely before Operations and Invasive Procedure: Single Center Retrospective Study. International Journal of Hematology-Oncology and Stem Cell Research, 2019, 13, 172-173.	0.3	0
16	Causes of Death Among 379 Patients With Hemophilia: A Developing Countryâ€™s Report. Clinical and Applied Thrombosis/Hemostasis, 2018, 24, 612-617.	0.7	11
17	Common solvents for making extraction of allergenic proteins from plantsâ€™ pollens for prick tests and related factors: a technical review. Electronic Physician, 2017, 9, 4440-4446.	0.2	1
18	Bone Density Status in Bleeding Disorders: Where Are We and What Needs to Be Done?. Journal of Bone Metabolism, 2017, 24, 201.	0.5	9

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19	A close insight to factor VIII inhibitor in the congenital hemophilia A. Expert Review of Hematology, 2016, 9, 903-913.	1.0	11
20	Desmopressin acetate as a haemostatic elevator in individuals with combined deficiency of factors V and VIII: a clinical trial. Journal of Thrombosis and Haemostasis, 2016, 14, 336-339.	1.9	15
21	Normal range of bleeding time in west of Iran. ARYA Atherosclerosis, 2016, 12, 156-157.	0.4	0
22	Coincidence of Glanzmann's thrombasthenia with hereditary haemorrhagic telengiectasia in a man with gastrointestinal bleeding. Blood Coagulation and Fibrinolysis, 2015, 26, 98-100.	0.5	1
23	Clinical and laboratory approaches to hemophilia a. Iranian Journal of Medical Sciences, 2015, 40, 194-205.	0.3	23
24	Frequency of combined factor V and factor VIII deficiency in southern Iran. Blood Coagulation and Fibrinolysis, 2014, 25, 92-93.	0.5	0
25	The Association between Neutrophilic Hypersegmentation and Iron Deficiency with Regard to Folate Status in 16 - 30 Year-Old Women. Clinical Laboratory, 2014, 60, 517-22.	0.2	1
26	CIRCUMCISION IN MALES WITH BLEEDING DISORDERS. Mediterranean Journal of Hematology and Infectious Diseases, 2013, 5, e2013004.	0.5	15
27	An investigation of the spectrum of common and rare inherited coagulation disorders in north-eastern Iran. Blood Transfusion, 2013, 11, 233-40.	0.3	16
28	Acquired haemophilia A in a woman with autoimmune hepatitis and systemic lupus erythematosus; review of literature. Blood Coagulation and Fibrinolysis, 2012, 23, 71-74.	0.5	6
29	Inherited factor V deficient neonate with galactosaemia. Clinical Biochemistry, 2012, 45, 356-358.	0.8	1
30	Acquired haemophilia A in a woman with autoimmune hepatitis and systemic lupus erythematosus. Blood Coagulation and Fibrinolysis, 2011, 22, 738-741.	0.5	4
31	Bleeding disorders and reduced bone density. Rheumatology International, 2011, 31, 283-287.	1.5	5
32	Spectrum of inherited bleeding disorders in southern Iran, before and after the establishment of comprehensive coagulation laboratory. Blood Coagulation and Fibrinolysis, 2010, 21, 296.	0.5	2
33	Haemorrhagic manifestations and prevalence of factor V deficiency in northâ€eastern Iran. Haemophilia, 2010, 16, 376-380.	1.0	12
34	Acquired Hemophilia A: Clinical Features, Surgery and Treatment of 34 Cases, and Experience of Using Recombinant Factor VIIa. Clinical and Applied Thrombosis/Hemostasis, 2010, 16, 294-300.	0.7	34
35	Successful Use of Factor VIII Concentrate and Fresh Frozen Plasma for Four Dental Extractions in an Individual with Combined Factor V and VIII Deficiency. Transfusion Medicine and Hemotherapy, 2009, 36, 138-139.	0.7	9
36	Idiopathic Factor VIII Inhibitor Autoantibody in a Man Presented After Accident. Clinical and Applied Thrombosis/Hemostasis, 2009, 15, 588-590.	0.7	5

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37	Reduced bone density in individuals with severe hemophilia B. International Journal of Rheumatic Diseases, 2009, 12, 125-129.	0.9	25
38	Are individuals with severe haemophilia A prone to reduced bone density?. Rheumatology International, 2008, 28, 1079-1083.	1.5	22
39	Detection of hepatitis G virus envelope protein E2 antibody in blood donors. International Journal of Infectious Diseases, 2008, 12, 445.	1.5	0
40	Pattern of factor VIII inhibitors in patients with hemophilia A in the north east of Iran. Hematology, 2006, 11, 215-217.	0.7	12
41	Combined factor V and VIII deficiency: a new family and their haemorrhagic manifestations. Haemophilia, 2006, 12, 169-171.	1.0	11
42	Haemorrhagic symptoms in patients with combined factors V and VIII deficiency in north-eastern Iran. Haemophilia, 2004, 10, 271-275.	1.0	54