# Hassan Mansouri Tourghabeh

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4743189/publications.pdf

Version: 2024-02-01

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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 | D-dimer level in COVID-19 infection: a systematic review. Expert Review of Hematology, 2020, 13, 1265-1275. | 1.0 | 242 |
| 2 | Haemorrhagic symptoms in patients with combined factors V and VIII deficiency in north-eastern Iran. Haemophilia, 2004, 10, 271-275. | 1.0 | 54 |
| 3 | 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 |
| 4 | Reduced bone density in individuals with severe hemophilia B. International Journal of Rheumatic Diseases, 2009, 12, 125-129. | 0.9 | 25 |
| 5 | 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 |
| 6 | Clinical and laboratory approaches to hemophilia a. Iranian Journal of Medical Sciences, 2015, 40, 194-205. | 0.3 | 23 |
| 7 | Are individuals with severe haemophilia A prone to reduced bone density?. Rheumatology International, 2008, 28, 1079-1083. | 1.5 | 22 |
| 8 | 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 |
| 9 | CIRCUMCISION IN MALES WITH BLEEDING DISORDERS. Mediterranean Journal of Hematology and Infectious Diseases, 2013, 5, e2013004. | 0.5 | 15 |
| 10 | 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 |
| 11 | 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 |
| 12 | Pattern of factor VIII inhibitors in patients with hemophilia A in the north east of Iran. Hematology, 2006, 11, 215-217. | 0.7 | 12 |
| 13 | Haemorrhagic manifestations and prevalence of factor V deficiency in northâ€eastern Iran. Haemophilia, 2010, 16, 376-380. | 1.0 | 12 |
| 14 | Combined factor V and VIII deficiency: a new family and their haemorrhagic manifestations. Haemophilia, 2006, 12, 169-171. | 1.0 | 11 |
| 15 | A close insight to factor VIII inhibitor in the congenital hemophilia A. Expert Review of Hematology, 2016, 9, 903-913. | 1.0 | 11 |
| 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 | 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 |
| 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | 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 |
| 20 | 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 |
| 21 | Current methods of measuring platelet activity: pros and cons. Blood Coagulation and Fibrinolysis, 2020, 31, 426-433. | 0.5 | 6 |
| 22 | Consanguineous marriage and rare bleeding disorders. Expert Review of Hematology, 2021, 14, 467-472. | 1.0 | 6 |
| 23 | 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 |
| 24 | Idiopathic Factor VIII Inhibitor Autoantibody in a Man Presented After Accident. Clinical and Applied Thrombosis/Hemostasis, 2009, 15, 588-590. | 0.7 | 5 |
| 25 | Bleeding disorders and reduced bone density. Rheumatology International, 2011, 31, 283-287. | 1.5 | 5 |
| 26 | Acquired haemophilia A in a woman with autoimmune hepatitis and systemic lupus erythematosus. Blood Coagulation and Fibrinolysis, 2011, 22, 738-741. | 0.5 | 4 |
| 27 | Mining of mortality-related findings in rare bleeding disorders: a retrospective study from two centers. Blood Research, 2020, 55, 213-216. | 0.5 | 4 |
| 28 | 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 |
| 29 | 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 |
| 30 | Inherited factor V deficient neonate with galactosaemia. Clinical Biochemistry, 2012, 45, 356-358. | 0.8 | 1 |
| 31 | 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 |
| 32 | 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 |
| 33 | 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 |
| 34 | 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 |
| 35 | 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 |
| 36 | Detection of hepatitis G virus envelope protein E2 antibody in blood donors. International Journal of Infectious Diseases, 2008, 12, 445. | 1.5 | 0 |

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|---|---|---|---|
| 37 | Frequency of combined factor V and factor VIII deficiency in southern Iran. Blood Coagulation and Fibrinolysis, 2014, 25, 92-93. | 0.5 | 0 |
| 38 | 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 |
| 39 | Normal range of bleeding time in west of Iran. ARYA Atherosclerosis, 2016, 12, 156-157. | 0.4 | 0 |
| 40 | 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 |
| 41 | Determining causes of death among individuals with haemophilia in Afghanistan. Haemophilia, 2021, , . | 1.0 | 0 |
| 42 | 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 |