



Curriculum Vitae and Publications

Personal details:

Minoo Ahmadinejad, Born on 21 September 1967, Married, one child

Area of Expertise:

Education and University degrees:

1986-1995, MD, Tehran University of Medical Sciences (TUMS), Tehran, Iran

1995-1997, Board certificated of Clinical & Anatomical Pathology. TUMS, Tehran, Iran

2012 , Two months Fellowship course in Centro Emophilia e Trombosi Angelo Bianci Bonomi supported by ISTH as ISTH Reach-the-World Fellowship

Google Scholar **Citations 324**; Google Scholar **h-index 10**

<https://scholar.google.com/citations?hl=en&user=ieqMG8UAAAAJ>

Linkedin : <https://www.linkedin.com/in/minoo-ahmadinejad/>

Current Position:

- Associate Professor of pathology, Blood Transfusion Research Centre, High Institute for Research and Education in Transfusion Medicine Tehran- Iran
- President of Iranian Society of Thrombosis and Hemostasis (IRSTH)
- Head of Pathology Faculty Group of Blood Transfusion Research Centre
- Head of reference coagulation lab of IBTO

Professional experiences and administrative positions:

- Faculty member of IBTO research center 2011- present.
- Member and secretary of executive committee of Iranian Society of Thrombosis and Hemostasis (IRSTH) 2016-2025
- Director, Reference Coagulation Lab, February 2000-Present.
(Organizing IBTO reference coagulation lab's membership in WHO and WFH IEQAS program for blood coagulation)
- Director of Iranian Blood Transfusion Organization (IBTO) central laboratory with 120 employees and 18 different laboratories, 2011-2014
- IBTO lab, ISO 9001-2008, managers' representative member.
- Member of executive committee, Iranian blood transfusion scientific association 2000-2006
- Member of hematology committee of Iranian Ministry of Health. for three years
- Establishment of special laboratory for diagnosis of "Heparin Induced Thrombocytopenia" for the first time in Iran since 2021.

Research projects and dissertation supervised

- Director of "Assessment of Venous Thromboembolism (VTE) Prevention and Thromboprophylaxis Policies in Hospitals of Tehran City" Started since August 2025 continued
- Director of Frequency of Platelet Secretion Disorders in Patients Referred to the Special Coagulation Laboratory of the Iranian Blood Transfusion Organization (IBTO). 2022, completed
- Director of project "Set up of homemade FVIII: VWF binding assay" will be started on 2026 up to 18 months.
- Director of project "Molecular evaluation of patients referred to IBTO reference coagulation lab of IBTO suspected for type 2N VWD" 2022 , completed
- Director of project "Determination of Sensitivity and Specificity of Flow Cytometry HIT Confirm versus HIPA method as an alternative method in confirming of heparin-induced thrombocytopenia (HIT), started at 2022, Completed 2023
- Director of project "Evaluation of diagnostic value of VWF:CB assay along with VWD screening panel on improvement of diagnosis and/ or differentiation of Iranian patient with VWD type 1 and type 2 in comparison of common diagnostic panel" 2017-2018
- Director of project "The prevalence of discrepancy of F VII coagulation factor in patients with hereditary FVII deficiency" 2017, completed

- Director of project “Evaluation of Blood Storage Lesions in Leuko-depleted Red Blood Cell Units” 2016, completed
- Director of project:” Determination of discrepancy level in factor VIII assay with “one-stage” an “chromogenic” method in mild and moderate hemophilia A patients”. 2015 Completed
- Director of project: Prevalence of Heparin Induced Thrombocytopenia among Iranian patients with cardiac surgery. 2015 Completed
- Cooperation with “Prevalence of Inhibitor Level in Factor XIII Deficient Patients Receiving Prophylaxis” project-2014 Completed
- Cooperation in international project titled “3 WINTERS”
- Cooperation in international project on F VIII inhibitor rate in hemophilia patients treated with plasma-derived products vs. rF VIII (SIPPET project) 2012-2015.
- Co-supervision of a Master of Science thesis on "Evaluation of exon 28 molecular defects in Iranian patient’s with VWD type 2" 2011
- Director of Project, " Evaluation of diagnostic validity of a new Functional test, based on prothrombin-time for the detection of Factor V Leiden mutation (Activated Protein C Resistance) in thrombophilia patients.2009
- Director of Project "Evaluation of factors affecting post transfusion platelet increments and platelet refractoriness in thrombocytopenic patients of Shariati Hospital, 2006
- Co-supervision of a Master of Science thesis on "Structural Analysis of Glycated Hemoglobin and its Effect on Platelet Aggregation" 2006
- Co-supervision of a Master of Science thesis on "Prevalence of elevated factor VIII:C in Iranian patients with venous thromboembolism"2006
- Co-supervision of a Master of Science thesis on "Prevalence of thrombophilia in patients with thalassemia" 2008

Seminars and workshops:

- IRSTH annual congresses 2020-2026
- Educational workshops on laboratory blood coagulation since 2010-present
- Coordinator and member of board director of several national seminars since 2004 – present.
- Scientific general secretary of two days seminar in "Bleeding Disorders" (2004).
- Scientific general secretary of two days’ workshop of "Coagulation Laboratory"(2005).
- Presentation of lectures about coagulation aspects in different Seminars (2005 till present).
- Administrative general secretary of seminar on Blood transfusion and Nurses (2008).
- Scientific general secretary of two days’ workshop of " Hemostasis and Thrombosis coagulation lab” (2010)

- Coordinator and member of board director of “Patient Blood Management and Coagulopathy “session in 18th Annual Pathology seminar, 2017, Tehran-Iran
- Preliminary Coagulation workshop (principles of coagulation screening tests), 2021/09/16-19, Special Coagulation Laboratory of IBTO, Tehran, Iran

Technical skills:

- Establishing referral HIT diagnostic lab, 2018 in collaboration with Greiswald university, Germany
- RTW Fellowship course (granted by ISTH) in molecular pathology of bleeding disorders 2014
- Teaching of theoretical aspects of "Basic Principles of Primary and Secondary Hemostasis ", "Coagulation Disorders" and" Platelet Disorders" for MSc and PhD students
- Teaching of practical aspects of coagulation test for hematologist/oncologist fellowships, pathology residents and MSc students.
- vWF Multimer Analysis (learned in a three months course in Angelo Bianchi Bonomi Hemophilia and Thrombosis Center of the Maggiore Hospital, Milan University, 2008)
- Platelet fibrinogen, nucleotide and serotonin level tests (learned in a three months course in Angelo Bianchi Bonomi Hemophilia and Thrombosis Center of the Maggiore Hospital, Milan University, 2008)
- ADAMTS 13 activity and inhibitor assay (learned in a three months course in Angelo Bianchi Bonomi Hemophilia and Thrombosis Centre of the Maggiore Hospital, Milan University, 2008).
- Learning of basic principles of DNA sequencing for diagnosis of VWD, Milan University, 2012)

Publications:

Papers:

- Platelet secretion defects and increased CD63 expression in Hermansky-Pudlak syndrome: a case report–Erratum, *Blood Coagulation & Fibrinolysis* 37(1):p 47, January 2026.
- Leukocyte adhesion deficiency type III in an infant presenting with intestinal perforation and low percentage of natural killer cells: first case report from Iran." *BMC pediatrics* 25.1 (2025): 315.
- A randomized, two-armed, double-blind, single-dose, cross-over, bioequivalence clinical trial to compare pharmacokinetic parameters and safety of recombinant human factor VIII with Fc fusion produced by AryoGen Pharmed Company versus Elocta ® (reference product) in previously treated patients with severe haemophilia A." *Annals of Hematology* 104.2 (2025): 1195-1202.
- Flow cytometry-based functional assay is a valuable diagnostic approach for confirmation of Heparin-Induced Thrombocytopenia - *Iranian Journal of Blood and Cancer*, 2024
- Responses of blood coagulation, fibrinolysis systems and von Willebrand to different interval and

continuous swimming protocols in elderly male Wistar rats , *Journal of Practical Studies of Biosciences in Sport Research Paper Spring 2024*, 12 (29), 80-92

- Utility of the international society on thrombosis and hemostasis-bleeding assessment tool in the diagnosis of patients who suspected of platelet function disorders , *Blood Coagulation & Fibrinolysis*, 2024
- The prevalence of Lupus Anticoagulant in Iranian children and adolescents; Current status and literature review of diagnosis *Iranian Journal of Blood and Cancer* 2023, 15(4): 262-271
- Analysis and Identification of Putative Novel Peptides Purified from Iranian Endemic Echis Carinatus Sochureki Snake Venom by MALDI-TOF Mass Spectrometry *Archives of Razi Institute*, 1481-1505
- Biochemistry of the Thrombin-Like enzyme and Its Purification from Iranian Echis carinatus snake venom: its interaction with platelet receptors, *Archives of Razi Institute* (2023)
- Analysis and identification of Some Putative Novel Peptides purified from Iranian endemic Echis carinatus sochureki snake venom by MALDI-TOF Mass Spectrometry, *Archives of Razi Institute DOI:10.22092/ARI.2023.361954.2701*
- Responses of blood coagulation, fibrinolysis systems and von Willebrand to different interval and continuous swimming protocols in elderly male wistar rats , *Journal of Practical Studies of Biosciences in Sport*, 2023;DOI : 10.22077/jpsbs.2023.6080.1768
- Identification of Diagnostic SNPs in Exon 28 of the Von Willebrand Factor Gene for Diagnosis of VWD Type 2B in Patients Referred to IBTO Coagulation Lab: In-silico Approach, *International Journal of Medical Laboratory* 2023 ;10 (2):127 -14 3 .
- A review on “Acquired Von Willebrand Syndrome” *Scientific Journal of Iran Blood Transfus Organ* 20 (2),(2023), 155-177
- von Willebrand factor neutralizing and non-neutralizing alloantibodies in 213 subjects with type 3 von Willebrand disease enrolled in 3WINTERS-IPS, *Journal of Thrombosis and Haemostasis* 21.4 (2023): 787-799.
- Isolation of the Anticoagulant and Procoagulant Fractions of the Venom of Iranian Endemic Echis carinatus *Iranian Journal of Pharmaceutical Research* 2022; 21 (1)
- Von Willebrand factor propeptide and pathophysiological mechanisms in European and Iranian patients with type 3 von Willebrand disease enrolled in the 3WINTERS-IPS study *Journal of thrombosis and haemostasis* 2022; 20 (5), 1106-1114
- Homozygous SOD1 mutation L144S produces a severe form of ALS in an Iranian family. *Neurology:Genetics* 8(1), 2022
- The activity of labile coagulation factors and fibrinogen in thawed plasma during a 5 day storage period in the hospital blood bank refrigerator. *Transfusion Medicine*. 2021
- Heparin-Induced Thrombocytopenia in Iranian Cardiac Surgery Patients Using the 4Ts Clinical Scoring System and Laboratory Methods. *International Journal of Hematology-Oncology and Stem Cell Research- 2012;15(4):230-238.*
- Genetic analysis of non-severe hemophilia A phenotype with A discrepancy between one-stage and chromogenic factor VIII activity assays. *Transfus Apher Sci*. 2021 Oct;60(5):103194.

- Von Willebrand disease combined with coagulation defects in Iran. *Blood Transfus* 2021 Sep;19(4):428-434.
- Genotypes of European and Iranian patients with type 3 von Willebrand disease enrolled in 3WINTERS-IPS. *Blood Adv.* 2021 Aug 10;5(15):2987-3001
- Severe Prekallikrein Deficiency Associated with Low Level of Factor XII: A Case Report. *Iran J Pathol.* 2021 Summer; 16(3): 332–336.
- Discrepancy between the results of one-stage and chromogenic factor VIII: C assays in patients with mild/moderate hemophilia A. *Blood Coagulation & Fibrinolysis* December 2020 - Volume 31 - Issue 8 - p 530-535
- Evaluation of Normal Range of Bleeding Scores in Healthy Iranian Adults using the International Society on Thrombosis and Hemostasis Bleeding Assessment Tool. *IJBC* 2020, 12(4): 126-130
- Are Iranian patients with von Willebrand disease type 2N properly differentiated from hemophilia A and do they receive appropriate treatment? *Blood Coagulation & Fibrinolysis* August 2020 - Volume 31 - Issue 6 - p 382-38
- 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
- Evaluation of Hemostatic Behavior of Micro and Nano Gelatin/Silica Hybrid in Severe Bleeding. *Current stem cell research & therapy*, 2019
- Procoagulant activity of red blood cell-derived microvesicles during red cell storage. *Transfusion Medicine and Hemotherapy*, 2019
- World Thrombosis Day, a Movement to the World without Thrombosis, *EC Cardiology* 6.12 (2019): 01-03.
- Heparin induced thrombocytopenia. *Sci J Iran Blood Transfus Organ* 2019;16(2): 133-148.
- Coagulation inhibitors in Thalassemia. *Sci J Iran Blood Transfus Organ* 2019;7(2): 78-84.
- Comparative evaluation of the safety and efficacy of recombinant FVIII in severe hemophilia A patients. *JOURNAL OF PHARMACOPUNCTURE*. 2018 Jun 1;21(2):76-81.
- Evaluation of diagnostic value of 4T score for heparin induced thrombocytopenia. *Sci J Iran Blood Transfus Organ* 2018; 14(4): 272-280
- “Is it Time to Renew the Guidelines for Diagnosis of HIT?”. *EC Cardiology* 5.7 (2018): 371-373.
- Evaluation of Blood Storage Lesions in Leuko-depleted Red Blood Cell Units. *Iranian Journal of Pediatric Hematology & Oncology*. 2017, Vol. 7 Issue 3, p171-179. 9p.
- Combined occurrence of Bernard-Soulier syndrome and prekallikrein deficiency. *Blood Res.* 2017 Sep;52(3):229-231
- Evaluation of blood use in cardiac surgery. *Sci J Iran Blood Transfus Organ* 2017; 14(3): 188-194
- THE RESULTS OF TWO METHODS OF FACTOR VIII: C IN PATIENTS WITH HEMOPHILIA A. *THE SCIENTIFIC JOURNAL OF IRANIAN BLOOD TRANSFUSION ORGANIZATION (KHOON) WINTER 2017, Volume 13 , Number 4; Page(s) 277 To 283.*

- Investigating the Frequency of Resistance to Activated Protein C and Protein S, C and Antithrombin Deficiencies in Patients with Deep Vein Thrombosis Referred to the Iranian Blood Transfusion Organization. *Journal of Advances in Medical and Biomedical Research* 2017;25(112):21-29.
- A Randomized Trial of Factor VIII and Neutralizing Antibodies in Hemophilia A. *N Engl J Med* 2016; 374:2054-2064
- Platelet Aggregation Increased by Advanced Glycated Hemoglobin. *Asadifar et al., J Blood Disord Transfus* 2015, 6:4
- Long-term prophylaxis in patients with severe congenital factor XIII deficiency is not complicated by inhibitor formation *Blood Coagul Fibrinolysis* 27:000–000 Copyright 2016
- Elevated Factor VIII activity and venous thromboembolism in patients referred to the Iranian Blood Transfusion Organization: A case control study. *Acta Haematologica Polonica* 47 (2 0 1 6) 1 5 – 1 8
- PREPARATION AND CHARACTERIZATION OF NOVEL GELATIN/SILICA NANO-HYBRID AS A STYPTIC FOR MASSIVE BLEEDING, *Digest Journal of Nanomaterials and Biostructures*, Vol 11, Issue 4 – 2016
- Combined high molecular weight Kininogen and factor XI deficiency *Haemophilia* (2016), 22, e36--e79
- A Comparison Between Recombinant Activated Factor VII (Aryoseven) and Novoseven in Patients with Congenital Factor Deficiency. *Clin Appl Thromb Hemost*, Vol 21, Issue 8, 2015.
- A comparison of efficacy between recombinant activated factor VII (Aryoseven) and Novoseven in patients with hereditary FVIII deficiency with inhibitor, *Clinical and Applied Thrombosis/Hemostasis* 2016, Vol. 22(2) 184-190
- Evaluation of Aggregometry Parameters and Efficacy of Plavix versus Clopidex in Patients Suffering from Ischemic Heart Disease: A Randomized Double-Blind Clinical Trial *Iran Red Crescent Med J.* 2014 February; 16(2): e15277.
- The caspase-3 inhibitor (peptide Z-DEVD-FMK) affects the survival and function of platelets in platelet concentrate during storage *Blood Res.* 2014Mar;49(1):49-53.
- Effects of caspase 3 inhibitor on the survival of platelet concentrate during storage. *Sci J Iran Blood Transfus Organ* 2014; 10(4): 319-325
- Squamous metaplasia in thyroid gland as histologic diagnostic pitfall a case report. *Tehran University Medical Journal*, October 2013; Vol. 71, No. 7: 480-484
- The prevalence of platelet function defects in patients referred to the special reference coagulation laboratory of Iranian Blood Transfusion Organization. *Scientific Journal of Iranian Blood Transfusion Organization.* Spring2014, Vol. 11 Issue 1, p23-31. 9p.
- Frequency of Bleeding Disorders in Women Presenting with Menorrhagia in North of Iran *World Applied Sciences Journal* 23 (1): 01-06, 2013
- Squamous metaplasia in thyroid gland as histologic diagnostic pitfall: a case report. *Tehran University Medical Journal*, October 2013; Vol. 71, No. 7: 480-484
- Acquired factor V inhibitor developing in a patient with esophageal squamous cell carcinoma, *Blood Coagulation and Fibrinolysis Journal*, 2013 Jan;24(1):97-9.

- Prevalence of Clopidogrel resistance in patient's candidate for percutaneous coronary intervention. *Pejouhandeh* 2011;16(1):47-51.
- Analysis of levels of natural anticoagulant proteins and activated protein C resistance in venous thromboembolism patients referred to Iranian Blood Transfusion Organization, Tehran *Sci J Iran Blood Transfus Org* 2011; 8(1): 1-9
- Effects of L-Carnitine supplement on plasma coagulation and anticoagulation factors in hemodialysis patients *Renal Failure*, 32:9, 1109-1114,
- Effects of L-carnitine supplement on plasma coagulation and anticoagulation factors in hemodialysis patients, *Iranian Journal of nutrition and Food Technology*, Vol 4, No.3 Autumn 2009,25-34
- Effects of soy consumption on oxidative stress, blood homocysteine, coagulation factors, and phosphorus in peritoneal dialysis patients. *J Ren Nutr.* 2009 Sep;19(5):389-95. Epub 2009 Jul 3
- PREVALENCE OF ELEVATED FACTOR VIII IN IRANIAN PATIENTS WITH IDIOPATHIC VENOUS THROMBOEMBOLISM. *Iranian Blood journal (quarterly)*, 2008, 3,149-156
- Anticoagulant system activity in splenectomized versus non-splenectomized Iranian patients with beta-thalassemia major. *Pediatric Hematol Oncol.* 2007 Dec;24(8):645-7
- Comparison of the Cytology Technique and the Frozen Section Results in Intraoperative Consultation Of the Breast Lesions *Acta Medica Iranica* 2002; 40(3):203-206

Abstracts:

- Presence of Subjects With von Willebrand Factor Polymorphism p. D1472H Among Patients With Low VWF: GPIbR Levels." *HAEMOPHILIA*. Vol. 31. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2025.
- Platelet Storage Pool Defects (SPDs) in Syndromic Patients." *HAEMOPHILIA*. Vol. 31. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2025.
- Application of von Willebrand Factor Propeptide in Identification of Type 1C von Willebrand Disease Patients." *HAEMOPHILIA*. Vol. 31. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2025.
- Familial Multiple Coagulation Factor Deficiencies (VWD/FVIID): A Study From Five Families." *HAEMOPHILIA*. Vol. 31. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2025.
- Clustered Referral of Family Members Suspected of Type 2B von Willebrand Disease *HAEMOPHILIA*, 2025 Vol. 31. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2025.
- Bleeding Disorder of Unknown Cause, a Report From a Uni-Centre National Referral Coagulation LAB - *HAEMOPHILIA*, Vol. 31. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2025.
- Leukocyte adhesion deficiency, type III in an infant presenting with intestinal obstruction: A case report from Iran." *HAEMOPHILIA*. Vol. 30. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2024.
- Heavy menstrual bleeding in an adolescent with platelet GP VI deficiency and thrombocytopenia: A challenge between hereditary or acquired." *HAEMOPHILIA*. Vol. 30. 111 RIVER ST, HOBOKEN 07030-5774, NJ USA: WILEY, 2024.

- Frequency of Inherited platelet secretion disorders in patients referred to the special coagulation laboratory of the Iranian blood transfusion organization (IBTO). ISTH 2023 Congress
- Acquired von Willebrand syndrome in essential thrombocythemia patients referred to special coagulation laboratory of Iranian Blood Transfusion Organization (IBTO). ISTH 2022 Congress
- Heparin Induced Thrombocytopenia Before and During COVID-19 Pandemia in Iran. ISTH 2022 Congress
- Clinical and Phenotypic Evaluation of Patients with Low von Willebrand Factor. *Res Pract Thromb Haemost.* 2020; 4 (Suppl 1)
- von Willebrand Disease Combined with Other Coagulation Defects: Experience at a Single Reference Coagulation Laboratory in Iran. *Res Pract Thromb Haemost.* 2020; 4 (Suppl 1)
- The Utility of ISTH-BAT in Iranian Patients with von Willebrand Disease. *Res Pract Thromb Haemost.* 2020; 4 (Suppl 1)
- The Prevalence of Lupus Anticoagulant in Children and Adolescents Referred to the Reference Coagulation Laboratory of Iranian Blood Transfusion Organization [abstract]. *Res Pract Thromb Haemost.* 2020; 4 (Suppl 1)
- Activated recombinant Factor VII (rFVIIa) Inhibitor in Iran: Immunogenicity is a Major Challenge through Treated Bleeds Covered with rFVIIa [abstract]. *Res Pract Thromb Haemost.* 2020; 4 (Suppl 1).
- Diagnostic problems of hemophilia in Iran (Case report). *The Second National Congress of Thrombosis and Homeostasis of Iran 2019, January 2-4.*
- Methods and interpretation of platelet function tests. *The Second National Congress of Thrombosis and Homeostasis of Iran 2019, January 2-4.*
- Evaluation of coagulability in patients with thalassemia intermedia. *The Second National Congress of Thrombosis and Homeostasis of Iran 2019, January 2-4.*
- Heparin induced thrombocytopenia: clinical evaluation and laboratory diagnosis. *The 11th International and 16th National Congress on Quality Improvement in Clinical Laboratories 2018, April 19-22.*
- Time course evaluation of erythrocyte derived microvesicles in stored packed cell concentrates . *XIVth Annual Asian Association of Transfusion Medicine (AATM) conference and 4th Iran International Congress of Transfusion Medicine .Tehran 2-4/10/2018.*
- A clinical challenge for haemostatic level and significance of bleeding episodes in women with heterozygote factor XIII deficiency. *XIVth Annual Asian Association of Transfusion Medicine (AATM) conference and 4th Iran International Congress of Transfusion Medicine .Tehran 2-4/10/2018.*
- The rate of anti PF4 antibody among the patients suspected for HIT referred to special coagulation lab of IBTO. *XIVth Annual Asian Association of Transfusion Medicine (AATM) conference and 4th Iran International Congress of Transfusion Medicine .Tehran 2-4/10/2018.*
- The rate of hereditary fibrinogen deficiency in Iran. *XIVth Annual Asian Association of Transfusion Medicine (AATM) conference and 4th Iran International Congress of Transfusion Medicine .Tehran 2-4/10/2018.*

- Storage of red cell concentrate affects RBC microvesiculation, and hemolysis rat. *The 9th International Congress on Laboratories and Clinics and 2nd National Congress on Basic Medical Sciences and Knowledge-based Production 2017 February 21-24.*
- The Diagnostic Value of Rotational Thromboelastometry in Patients with Non-severe Hemophilia A, *Res Pract Thromb Haemost.* 2017;1(Suppl. 1):813.
- Prevalence of Heparin Induced Thrombocytopenia among Iranian Patients with Cardiac Surgery, *Res Pract Thromb Haemost.* 2017;1(Suppl. 1):1366.
- The role of platelet function testing in PBM. *Tehran 2016.*
- The role of POC testing in antiplatelet drug management. *The 18th Annual Conference and the 1st International Conference on Pathology and Laboratory Medicine 2016 December*
- Laboratory evaluation of homeostasis. *Conference on Therapeutic Challenges Facing Doctors Working in Hemophilia Centers 2016 December 16*
- Prevalence of Discrepancy Between the Results of One-Stage and Chromogenic Factor VIII: C Assays in Iranian Patients with Mild/Moderate Hemophilia A, *Blood* 2016 128:4982;
- The role of laboratory in diagnosis and treatment of hemophilia. *3rd International Congress of Transfusion Medicine, Evidence-Based use of Blood Components and Plasma Derived Medicine. Tehran 15-17/12/2015.*
- THE INVESTIGATION OF NATURAL ANTICOAGULANT DEFICIENCIES IN PATIENTS WITH DEEP VEIN THROMBOSIS *THROMBOSIS RESEARCH Volume 11 PagesS45-S45*
- Prevalence Of Anemia Among Non-pregnant Women In Child Bearing Age In Iran: A Two-year Cross Sectional Study *Vox Sanguinis Vol 110 Pages 89-2014.*
- Evaluation of 195 referral cases with suspected hereditary platelet function disorders using platelet aggregometry and Atp release *2013 Journal of Thrombosis and Haemostasis Volume 11 Pages1149-1150. Winner of a “Reach The World” award*
- PREVALENCE OF FACTOR V LEIDEN IN 318 IRANIAN INDIVIDUAL REFERRED CASES AND THE DIAGNOSTIC POWER OF TWO DIFFERENT APC-R SCREENING TESTS, abstract N°SSC12-1230, **Winner of a “Reach the World” award**
- Comparison of diagnostic power of the two different APC-R screening tests in diagnosis of factor V Leiden mutation in patient’s referred coagulation lab of Iranian Blood Transfusion Organization. *2nd National Congress of Hematology. Yazd. 16-18/05/2012*
- ISTH 2011 P-TH-211 ACQUIRED FACTOR V INHIBITOR DEVELOPING IN A PATIENT WITH ESOPHAGEAL SQUAMOUS CELL CARCINOMA. *JTH, 2011;9 Supplement s2:P-124 , Winner of DWSA award*
- ISTH 2011 P-TH-322 THROMBOPHILIA IN IRANIAN WOMEN WITH PREGNANCY. *JTH, 2011;9 Supplement s2:P-160*
- ISTH 2011 P-TH-322 PREVALENCE OF THROMBOPHILIC DEFECTS IN WOMEN WITH ADVERSE PREGNANCY OUTCOMES. *JTH, 2011;9 Supplement s2:P-169*
- ISTH 2011 P-MO-002 PLATELET REACTIVITY INCREASED BY ADVANCE GLYCATED HEMOGLOBIN. *JTH, 2011;9 Supplement s2:P-59 Iron Stores In Male Blood Donors Vox Sanguinis. 101():122–123, Jul 2011*

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- INHERITED THROMBOPHILIC DEFECTS IN IRANIAN PATIENTS WITH PULMUNARY EMBOLISM

Vox Sanguinis 2008; 95 Supplement 1: P-472

- EVALUATION OF PLASMA HOMOCYSTEINE AND NATURAL COAGULATION INHIBITORS IN PATIENTS WITH THROMBOSIS REFERRED TO COAGULATION LAB OF BLOOD TRANSFUSION *Vox Sanguinis* 2008; 95 Supplement 1: P-359
- DESIGN OF A NEW LINER IN IRANIAN BLOOD TRANSFUSION ORGANISATION (IBTO) TO PREVENT FOLDING OF BLOOD BAGS WHILE CENTRIFUGING: *Vox Sanguinis*. 93 (Suppl. 1):99-100, July 2007.
- HEREDITARY DEFICIENCIES OF ANTITHROMBIN, PROTEIN S, AND THE PROTEIN C PATHWAY IN IRANIAN DVT PATIENTS. *J Thromb Haemost* 2007;5 Supplement 2: P-S-572
- HIGH PREVALENCE OF ELEVATED FACTOR VIII IN IRANIAN PATIENTS WITH IDIOPATHIC VENOUS THROMBOEMBOLISM *J Thromb Haemost* 2007; 5 Supplement 2: P-W-466
- HEREDITARY THROMBOPHILIC DEFECTS IN IRANIAN PATIENTS WITH CEREBROVASCULAR ACCIDENT. *J Thromb Haemost* 2007; 5 Supplement 2: P-T-576
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References:

Professor P.M. Mannucci (Milan University, piermannuccio.mannucci@policlinico.mi.it)

Professor F.Peyvandi (Milan University, flora.peyvandi@unimi.it)

Dr.S.Shariat Torbaghan (Tehran University of Medical Sciences, siamak_shariat@yahoo.com)

Dr.P.Eshghi (Shaheed Beheshti University of Medical Sciences, peyman64@gmail.com)