



## Genetic, Functional and Therapeutic Aspects of Procoagulant and Anticoagulant Factors

Guest Editors:

**Dr. Rima Dardik**

National Hemophilia Center,  
Sheba Medical Center, Ramat  
Gan 52621, Israel

**Prof. Tami Livnat**

1. Director, Amalia Biron  
Research Institute of Thrombosis  
and Hemostasis, Faculty of  
Medical and Health Sciences, Tel-  
Aviv University Sheba Medical  
Center, Tel Hashomer 52621,  
Israel  
2. Rabin Medical Centre,  
Ophthalmology Department and  
Laboratory of Eye Research  
Felsenstein Medical Research  
Centre, Petah-Tikva 49100, Israel

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### Message from the Guest Editors

Dear Colleagues,

Normal hemostasis is highly dependent on the balance between procoagulant systems (e.g. platelets, procoagulant factors) and anticoagulant systems (e.g. protein C, protein S, antithrombin). Lack or dysfunction of a major procoagulant factor results in a bleeding disorder (e.g. factor VIII deficiency leading to hemophilia A), whereas a defect in an essential anticoagulant system (e.g. protein C deficiency) leads to a thrombotic disorder. Beyond their function in hemostasis, cell signaling pathways which are induced by procoagulant (e.g. thrombin, FVII) and anticoagulant factors (e.g. APC), mediated by specific receptors, have become the focus of increasing attention, with some of them being explored as promising therapeutic targets.

This special issue will focus on genetic, functional and therapeutic aspects of procoagulant and anticoagulant systems. We invite research studies and reviews on genetics of bleeding and thrombotic disorders, functional and therapeutic studies of procoagulant and anticoagulant factors, as well as studies exploring their involvement in the crosstalk between hemostasis and inflammation.

Dr. Rima Dardik

Dr. Tami Livnat

*Guest Editors*





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### **Prof. Dr. Maurizio Battino**

Department of  
Odontostomatologic and  
Specialized Clinical Sciences,  
Sez-Biochimica, Faculty of  
Medicine, Università Politecnica  
delle Marche, Via Ranieri 65,  
60100 Ancona, Italy

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*International Journal of Molecular  
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