

Special Issue

Thrombocytopenia and ITP: Causes, Symptoms, and Treatment

Message from the Guest Editors

Thrombocytopenia encompasses a diverse range of conditions, ranging from acquired to constitutive. With the availability of high throughput molecular techniques, new insights into the etiology of immune and inherited thrombocytopenia syndromes have markedly improved our ability to diagnose and counsel patients accurately and promise to aid in the development of novel therapeutics. This Special Issue of the Journal of Clinical Medicine will cover the following important aspects of this field:

- Molecular understanding of thrombopoiesis
- Molecular basis of inherited platelet defects
- Leukemia predisposition in inherited thrombocytopenia syndromes
- Mechanisms of immune dysregulation in immune thrombocytopenia
- New therapeutic developments

Guest Editors

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Journal of Clinical Medicine (ISSN 2077-0383) is an international, peer-reviewed, open access journal dedicated to advancing clinical and preclinical research. Our Editorial Board and staff are committed to delivering cutting-edge, timely, peer-reviewed articles spanning the diverse subspecialties of clinical medicine. The journal publishes concise, innovative, and impactful research articles, as well as reviews that are clinically significant and relevant to the full spectrum of medical disciplines. All full texts are archived in PubMed Central and indexed in SCIE (Web of Science), Scopus, PubMed, and other major databases. We invite you to submit your manuscript for publication, as well as join our community as a reviewer or reader online.

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