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Thrombocytopenia and ITP: Causes, Symptoms, and Treatment

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Deadline for manuscript submissions:
closed (31 October 2016)

Message from the Guest Editors

Dear Colleagues,

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

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Editors-in-Chief

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There has been an explosion of gene and target based research and therapeutics in the multitude of fields that compose clinical medicine. The *Journal of Clinical Medicine*'s (*JCM*) staff and editorial board are dedicated to providing cutting edge, timely, and peer-reviewed articles covering the diverse subspecialties of clinical medicine. The journal publishes concise, innovative, and exciting research articles as well as clinically significant articles and reviews that are pertinent to the myriad of disciplines within medicine. The articles published are relevant to both primary care physicians and specialists. The journal's full-texts are archived in PubMed Central and indexed in PubMed. Please consider submitting your manuscripts for publication to our journal and check us out on-line!

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