



Bridging the Gap between the Clinical and Regulatory Use of Plasma Rich in Growth Factors. Where Are We?

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Deadline for manuscript
submissions:

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Message from the Guest Editors

Platelet-rich plasma (PRP) therapies have been attracting great attention in both the research and clinical community. However, the field faces important challenges at the basic, clinical, and regulatory frameworks. These three fields are progressing at a different pace, thereby generating a gap that in certain occasions is difficult to bridge effectively and efficiently. An evidence-based approach and appropriate regulatory framework would ensure both patient safety and the consolidation of current treatments and development of future applications. At the same time, quality studies are needed to promote the safe advancement of these types of biologic therapies.

Manuscripts are welcome to be submitted to cover the topics of platelet-rich plasma therapies, including in vitro research, preclinical in vivo, clinical studies or regulatory issues regarding the use of PRPs. However, the topics of interest are not limited to these. This issue aims to put in the same space and time coordinated basic, translational, and clinical research together with a regulatory/ethical overview in an effort to approximate the needs and bridge the gap.





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Message from the Editor-in-Chief

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