Special Issue

Advances in Clinical Management of Pulmonary Arterial Hypertension

Message from the Guest Editor

Pulmonary arterial hypertension (PAH) is characterized by an abnormal and chronic increase in the mean pulmonary artery pressure. PAH has been associated with severe cardiac remodelling and life-threatening comorbidities. Although scientific knowledge has evolved throughout past decades, new strategies will be required to improve the management and quality of the life of PAH patients. This Special Issue, titled Advances in Clinical Management of Pulmonary Arterial Hypertension, aims to highlight recent clinical and experimental discoveries regarding the mechanisms of PAH and the potential translational applications that could lead to developments of new treatment approaches, thus improving the diagnosis and clinical management of PAH. In this Special Issue, we welcome original research articles and reviews. Research areas may include (but are not limited to) the following: pulmonary arterial hypertension, inflammation, resolution, cardiopulmonary disorders, and new therapeutic strategies in PAH management. I look forward to receiving your contributions. Sincerely,

Guest Editor

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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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