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Pulmonary Arterial Hypertension: Old Drugs and New Treatment Strategies

Guest Editor:

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Deadline for manuscript submissions:
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Message from the Guest Editor

Pulmonary Hypertension (PH) is a hemodynamic condition characterized by an increase in pressure in the pulmonary artery secondary to various pathological conditions. A correct diagnosis and a correct clinical framing of PH are prerequisites for adequate treatment able to improve not only the symptoms and quality of life, but also the survival rates of patients. The implementation of aggressive therapy protocols focused on the combination of drugs with synergistic action from the beginning and the availability of new molecules acting on alternative cellular and molecular targets are now changing the outcomes of the disease, and are crucial challenges on which research and clinical practice must focus efforts and resources. In this Special Issue, we welcome authors to submit papers on the clinical advance of PH in terms of both diagnosis, risk assessment and treatment.

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