Special Issue

Pulmonary Hypertension: From Bench to Bedside: 2nd Edition

Message from the Guest Editor

Pulmonary arterial hypertension is life-threatening diseases caused by constriction and remodeling of pulmonary arteries. Prostacyclin, endothelin receptor antagonist, phosphodiesterase 5 inhibitor and soluble guanylate cyclase stimulator is an available for treatment of pulmonary arterial hypertension, which improve hemodynamics and exercise capacity. However, sufficient improvement of long-term prognosis is not achieved in all patients with pulmonary arterial hypertension. New findings from basic research contribute to therapeutic advances of pulmonary arterial hypertension. In this special issue, we focus on basic research in pulmonary arterial hypertension. Volume I: https://www.mdpi.com/journal/life/special_issues/pulmonary_hypertension_2022

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Deadline for manuscript submissions

closed (1 July 2024)

Life

an Open Access Journal by MDPI

Impact Factor 3.4
CiteScore 6.0
Indexed in PubMed

mdpi.com/si/145808

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