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Hereditary Hemorrhagic Telangiectasia: Recent Advances and Future Challenges

Guest Editors:

Dr. Hans-Jurgen Mager
Department of Pulmonology, St. Antonius Ziekenhuis, Utrecht, The Netherlands

Prof. Dr. Carmelo Bernabeu
Centro de Investigaciones Biológicas "Margarita Salas", Consejo Superior de Investigaciones Científicas (CSIC), 28040 Madrid, Spain

Dr. Marco Post
Department of Cardiology, St. Antonius Ziekenhuis, Nieuwegein/Utrecht and University Medical Center, Utrecht, The Netherlands

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Message from the Guest Editors

Hereditary Hemorrhagic Telangiectasia (HHT) is an autosomal heritable disease, leading to vascular malformations, ranging from mucocutaneous telangiectases to large arteriovenous malformations, which can occur in different organs. HHT is associated with a decreased quality of life and severe complications. If untreated, the disease leads to a decreased life expectancy. Recent years have brought advances in diagnosis and treatment, but not a cure for HHT. The exact molecular etiology is still unknown, but important steps in unravelling the mechanisms of disease haven been made.

This Special Issue aims to highlight not only the current knowledge regarding diagnosis and treatment of HHT, but also the newest insights in the molecular basis of HHT, because understanding the mechanisms of disease is essential for the development of new medicines or therapeutic strategies.

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Special Issue

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Editors-in-Chief

Prof. Dr. Emmanuel Andrès
Internal Medicine Department, University Hospital Strasbourg, 67081 Strasbourg, France

Prof. Dr. Kent Doi
Department of Emergency and Critical Care Medicine, The University of Tokyo, Tokyo, Japan

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Journal of Clinical Medicine Editorial Office
MDPI, Grosspeteranlage 5
4052 Basel, Switzerland

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