



Molecular Players in Diagnostics and Therapeutics of Malignant Pleural Mesothelioma

Guest Editors:

Prof. Dr. Eric Santoni-Rugiu

1. Department of Pathology,
Diagnostic Center,
Rigshospitalet, Copenhagen
University Hospital, Blegdamsvej
9, 2100 Copenhagen E., Denmark
2. Biotech Research & Innovation
Center, University of
Copenhagen, Ole Maaløes Vej 5,
DK-2200 Copenhagen N.,
Denmark

Prof. Dr. Jens Benn Sørensen

Department of Oncology, Center
for Cancer and Organ Diseases,
Rigshospitalet, Copenhagen
University Hospital, Blegdamsvej
9, 2100 Copenhagen E., Denmark

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Message from the Guest Editors

Malignant pleural mesothelioma (MPM) is a very aggressive cancer originating from the mesothelial cells lining the pleural cavity, usually associated with exposure to asbestos or asbestos-like fibers and characterized by challenging diagnostics, poor response to current standard therapies, and dismal prognosis. MPM is difficult to distinguish from reactive pleural conditions using conventional imaging- and histopathology-based methodologies.

However, recent seminal comprehensive studies have enlightened the role played in MPM by important genetic, epigenetic, and immunological alterations as well as by the interplay between genetic cancer susceptibility and environmental factors. This allows delineating novel molecular and clinical subtypes, as well as promising diagnostic and therapeutic strategies for MPM.

This Special Issue aims at collecting and welcomes original research articles, state-of-the-art review articles in this field.





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Editor-in-Chief

Prof. Dr. Maurizio Battino

Department of
Odontostomatologic and
Specialized Clinical Sciences,
Sez-Biochimica, Faculty of
Medicine, Università Politecnica
delle Marche, Via Ranieri 65,
60100 Ancona, Italy

Message from the Editor-in-Chief

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