Special Issue

Mechanisms and Novel Therapeutic Approaches for Muscle Disease

Message from the Guest Editors

Muscular dystrophy is a group of inherited rare diseases that causes progressive weakness and the degeneration of skeletal and cardiac muscles, with or without damaging the nerve supplied to the muscles. The diseases often affect a particular group of muscles before spreading more widely to other muscle groups, increasing the level of disability and possible deformity or even becoming life-threatening when cardiac or/and respiratory muscles are severely impaired. This Special Issue on "Mechanisms and Novel Therapeutic Approaches for Muscle Disease" therefore welcomes basic to translational research studies that aim to provide up-to-date insights into pathophysiological mechanisms, innovative methodologies, cellular and animal models, and novel therapeutic strategies that have a great potential for the development of a cure or modifying treatments for muscular dystrophies.

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Biomedicines (ISSN 2227-9059) is an open access iournal devoted to all aspects of research on human health and disease, the discovery and characterization of new therapeutic targets, therapeutic strategies, and research of naturally driven biomedicines, pharmaceuticals, and biopharmaceutical products. Topics include pathogenesis mechanisms of diseases, translational medical research, biomaterial in biomedical research, natural bioactive molecules, biologics, vaccines, gene therapies, cell-based therapies, targeted specific antibodies, recombinant therapeutic proteins, nanobiotechnology driven products, targeted therapy, bioimaging, biosensors, biomarkers, and biosimilars. The journal is open for publication of studies conducted at the basic science and preclinical research levels. We invite you to consider submitting your work to Biomedicines, be it original research, review articles, or developing Special Issues of current key topics.

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