Special Issue

Advances in Pediatric Neuromuscular Disorders: Second Edition

Message from the Guest Editor

In the past decade, there has been a quantum leap in therapies for pediatric neuromuscular disorders. Diseases which were once untreatable, such as spinal muscular atrophy and Pompe disease, now have multiple therapies available including enzyme replacement therapy, post-transcription RNA modulars and gene therapy. Life expectancy and quality of life have improved for patients with Duchenne muscular dystrophy and other muscular dystrophies. Newer and safer agents are now available for previously treatable disorders, such as myasthenia gravis. Ongoing multifaceted research is taking place regarding early diagnosis, treatment and remedies to improve the quality of life and outcomes for patients with various neuromuscular disorders. Additionally, like everything in medicine, COVID-19 has introduced a new paradigm of neuromuscular diseases. This Special Issue aims to explore the frontier of the evolving field of neuromuscular medicine.

Guest Editor

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Deadline for manuscript submissions

closed (25 March 2025)

Children

an Open Access Journal by MDPI

Impact Factor 2.1
CiteScore 3.8
Indexed in PubMed

mdpi.com/si/215126

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Message from the Editor-in-Chief

You are invited to contribute a research article or comprehensive review for consideration and publication in *Children* (ISSN 2227-9067). *Children* is an open access journal—research articles, reviews, and other content are published online immediately after acceptance. The scientific community and the general public have unlimited free access to the content as soon as it is published. The journal focuses on sharing clinical, epidemiological, and translational science relevant to children's health. We would be pleased to welcome you as one of our authors.

Editor-in-Chief

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