Special Issue

Advances of Phenylketonuria in Children

Message from the Guest Editors

From 1934 first detected phenyl pyruvic acid to 1963 a method for measuring phenylalanine from dried blood spots of newborns, it took almost 30 years. However, important drawbacks remain in the management of phenylketonuria (PKU) now. Adherence to dietary therapy, especially after the first decade of life, is often unsatisfactory, with consequences for neurocognitive function. Behavioral and emotional problems are still described in many continuously treated children and adolescents. The neuropathology of PKU remains a major knowledge gap. Sapropterin, an oral form of tetrahydrobiopterin, is an alternative pharmacological treatment for a subset of patients with PKU, mainly those with mild or moderate metabolic phenotypes. It is not known whether pegylated phenylalanine ammonia lyase, which requires daily subcutaneous injections, is safe and effective in children.

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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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