Special Issue

Precision Medicine in Neurodevelopmental Disorders: Challenges and Opportunities

Message from the Guest Editors

The scientific reports included in this Special Issue show advances in studies on pediatric diseases with intellectual disability. Since Dr. George A. Jervis diagnosed the first case of alkaptonuria in the last century, more than a hundred disorders, such as fragile X, intellectual disability X-linked syndromic 10, and HSD10 deficiency, have been discovered. Major advances in medical science have provided opportunities to prevent such refractory diseases. This Special Issue of Journal of Personalized Medicine, "Precision Medicine in Neurodevelopmental Disorders: Challenges and Opportunities", will promote the diagnosis and treatment of these diseases, helping patients with developmental disabilities to live richer lives.

Guest Editors

Prof. Dr. Song-Yu Yang

Dr. Xue-Ying He

Prof. Dr. Manfred Philipp

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About the Journal

Message from the Editor-in-Chief

Journal of Personalized Medicine (JPM; ISSN 2075-4426) is an international, open access journal aimed at bringing all aspects of personalized medicine to one platform. JPM publishes cutting edge, innovative preclinical and translational scientific research and technologies related to personalized medicine (e.g., precision medicine, pharmacogenomics/proteomics, systems biology, 'omics association analysis). JPM is covered in Scopus, the Science Citation Index Expanded (SCIE), PubMed, PMC, Embase, and other databases.

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