## Special Issue

## Computational and Omics Research on Rare Diseases

## Message from the Guest Editor

The comprehensive measurement of the molecular state of cells allowed by omics techniques, whether singly or together in a tissue, is rapidly redefining our understanding of disease and human development. We wish to focus such an approach on rare diseases. By definition, a rare disease affects fewer than one individual in 2000 people in the general population. However, as there are nearly 8000 rare diseases, it turns out that these "rare" diseases affect more than 300 million people worldwide. Genomics and all the emerging omics techniques hold the key to new diagnoses and therapies for rare diseases, but constant advances in analytical and computational methods are needed to support these techniques at increasing scale. Innovative approaches are also desirable to translate omics research into pharmacological therapies. We are seeking for papers that investigate rare diseases using an omics or integrated omics approach as well as computational omics involving artificial intelligence, big data, molecular simulations. Papers addressing specific pharmacological therapies involving drug-repositioning for rare diseases in light of omics studies are warmly welcomed.

## **Guest Editor**

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### Deadline for manuscript submissions

closed (30 April 2022)



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## Message from the Editor-in-Chief

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## **Editor-in-Chief**

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