Human Genetics Asia 2023

Special interest group:

Genomic medicine for sick newborn infants

Precise genetic diagnosis is continuously revolutionizing healthcare. It is undoubtful that genomic medicine provided great benefits to the most vulnerable population, namely, sick newborn infants. Indeed, rapid genetic diagnosis is increasingly recognized as a powerful and indispensable clinical tool in acute care settings in many countries. At the same time, we are facing new challenges that need to be solved in the future. This SIG aims to provide a focused and constructive discussion on the benefits and possibilities of future international collaboration. Those who are interested in the genomic medicine for sick newborn infants are welcomed.

