



Optical Genome Mapping in Hematological Malignancies

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Message from the Guest Editors

For more than 40 years, the clinical evaluation of structural variation in hematological malignancies has primarily been driven by karyotyping. Karyotyping is a robust tool for evaluating structural and numerical changes in leukemia and other malignancies due to its ability to visualize and catalogue recurrent changes to chromosomes. As a result, many types of leukemia and lymphoma have a cytogenetic (i.e., karyotyping-based) classification system for diagnosis and prognosis. However, karyotyping suffers from several challenges, including the need for dividing cells (metaphases) and the relatively “low resolution” of the technique. Thus, ancillary testing, such as FISH or RT-PCR, is required to identify specific rearrangements between genes. Optical Genome Mapping is a new technology that can detect structural variation within a sample at a much higher resolution than karyotyping. As such, it is a promising new technique for the clinical evaluation of chromosomal changes in hematologic malignancies.

The aim of this Special Issue is stimulate discussion around the use and adoption of Optical Genome Mapping for clinical diagnostics in hematological malignancies.





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Message from the Editor-in-Chief

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