

Special Issue

Targeted Therapies for Acute Leukemias

Message from the Guest Editor

Targeted therapies aim to interfere with the molecular pathways that drive the growth and survival of leukemia cells, while simultaneously mitigating any adverse effects on healthy cells. Several such agents have revolutionized treatment approaches and improved outcomes for many patients with acute leukemias. Acute myeloid leukemia (AML) is the most common acute leukemia in adults. Recent advances have significantly impacted treatment options. Acute lymphoblastic leukemia (ALL) is the most common acute leukemia in children. The targeted agents used in B-ALL are tyrosine kinase inhibitors when Philadelphia-positive, rituximab, blinatumomab, inotuzumab ozogamicin and, most recently, chimeric antigen receptor (CAR)-T cell therapies. Nevertheless, questions remain about optimal sequencing, effective combinations, maintenance therapy post-remission, especially in AML, and treatment options post-CAR-T cell therapy failure in ALL. This Special Issue focuses on known or developing targeted treatments in acute leukemias to provide clinicians a comprehensive perspective.

Guest Editor

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Diseases is an international, peer-reviewed, open access, multidisciplinary journal that focuses on the latest outstanding research concerning diseases and conditions. Research articles, reviews, and other contents are released on the Internet immediately after acceptance. This journal aims to cover international conferences and symposia as new targets.

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