

عنوان مقاله:

Strategies to overcome drug resistance to imatinib in chronic myeloid leukemia cells

محل انتشار:

کنفرانس بین المللی ژنتیک و ژنومیکس انسانی (سال: 1400)

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خلاصه مقاله:

Chronic myeloid leukemia (CML) is a type of leukemia that results from the clonal proliferation of bone marrow hematopoietic stem cells and is diagnosed with the BCR/ABL hybrid gene, which has abnormal tyrosine kinase activity. Using tyrosine kinase inhibitors (TKIs) such as imatinib (IM), significant advances have been made in the treatment of CML. However, drug resistance to IM has been identified as an important and basic barrier in the treatment of this disease. Recent evidence has shown that disruption of miRNAs regulation is associated with drug resistance of CML cells. miR-221, miR-199a/b-5p, miR-153-3p, and miR-57Y were found to be downregulated in IM-resistant cells. On the other hand, upregulation of miR-221, miR-199a/b-5p, miR-153-3p, and miR-57Y increased IM efficiency and markedly reduced the proliferation and survival of resistant cells. In contrast, the downregulation of these miRNAs reversed these effects in resistant cells. Moreover, inhibition of autophagy by inhibitors including 3-methyladenine and chloroquine significantly increased IM sensitivity in CML cells. Also, overexpression of miR-199a/b-5p and miR-153-3p led to induction of apoptosis and decreased autophagy in resistant cells. To sum up, the results of these studies demonstrated that inhibition of autophagy and overexpression of miR-153-3p, miR-199a/b-5p, miR-221, or miR-57Y combined with IM treatment could be a novel potential strategy to increase the sensitivity of CML cells to IM.

کلمات کلیدی:

chronic myeloid leukemia (CML), imatinib (IM), drug resistance, autophagy, miRNA

لینک ثابت مقاله در پایگاه سیویلیکا:

<https://civilica.com/doc/1516508>

