

## عنوان مقاله:

In silico analysis of identification miRNA-mRNA regulatory network in acute myeloid leukemia (AML) patients

## محل انتشار:

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## خلاصه مقاله:

Backgrounds: miRNAs are a class of non-coding RNA that is involved in various biological processes and diseases including cancers. Therefore, many studies illustrating the role of miRNAs in acute myeloid leukemia (AML) focused on identifying AML-specific miRNA expression patterns. AML is an invasive disease characterized by the increased proliferation and malignancy of immature myeloid cells. Therefore this study aims to investigate putative target genes and interaction networks where they are involved in AML. Also, Because of the numerous possible interactions between a single miRNA and target genes, bioinformatics analysis is very valuable to identifying putative pathways. Materials and Methods: The original data set GSE142699 was selected from the GEO dataset (NCBI), and then the differentially expressed miRNAs in cytogenetically normal acute myeloid leukemia patients were identified using the GEO2R. Their target genes were predicted from four (Targetscan, miRWalk, miRDB, miRmap) miRNA target prediction databases. Then, functional analysis was accomplished for the target genes using by the construction of a miRNAs-target gene network. Results: In current study, described five miRs (miR-382-5p, hsa-miR-151a-3p, hsa-miR-495-3p, hsa-miR-409-3p, and miR-135) with down-regulation and three miRs (hsa-miR-196b-5p, hsa-miR-34a-5p, and hsa-miR-181a-3p) with up-regulation in patients with AML. The miRNAs were exposed to the most used predictions software and >200 overlap target genes predicted. Then, enrichment analysis was performed revealing the KEGG pathway, comprising the cell cycle, Transcriptional dysregulation in cancer, and cellular senescence. Network construction was generated and links between the selected miRNAs and the predicted targets. Conclusion: In this study, we merged miRNA expression analysis with a bioinformatics-based workflow. Some genes (CDK6, HOXA9, RUNX1, and ITGB3), pathways, and interactions, putatively involved in AML development, were identified

## کلمات کلیدی:

miRNA, acute myeloid leukemia, network, bioinformatics

## لینک ثابت مقاله در پایگاه سیویلیکا:

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