

عنوان مقاله:

Histone deacetylase inhibitory and cytotoxic activities of the constituents from the roots of three species of *Ferula*

محل انتشار:

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خلاصه مقاله:

Objective(s): Histone deacetylase inhibitory and cytotoxic activities of 18 naturally occurring terpenoids (ferutinin, stylosin, tschimgine and guaiol), coumarins (umbelliprenin, farnesiferone B, conferone, feselol, ligupersin A, conferdione, conferoside) and sulfur-containing derivatives (latisulfies A-E, persicasulphides A and C) from the roots of three species of *Ferula* (*Ferula latisecta*, *Ferula ovina* and *Ferula flabelliloba*) were evaluated. **Materials and Methods:** The cytotoxic activity of compounds was evaluated against human cancer cell lines (HeLa, HCT116, A2780 and A549) by AlamarBlue® assay using vorinostat as the positive control. On the other hand, we aimed to evaluate their inhibitory activities against pan-HDAC. **Results:** The methanolic extract of the roots of *F. flabelliloba* was subjected to silica gel column chromatography. Further purification by preparative thin-layer chromatography (PTLC) and semipreparative RP-HPLC yielded twelve known compounds (1-12). This is the first report on the isolation of guaiol (1), persicasulphide C (3) and conferoside (10) from the roots of *F. flabelliloba*. Six compounds including persicasulfide A, conferone, feselol, latisulfide C, conferoside and ferutinin showed cytotoxic activity with IC₅₀ values in the range of 11.61-49.40 μM against cancer cells and pan-HDAC inhibitory activity with IC₅₀ values in the range of 1.06-35.27 μM. **Conclusion:** Results indicated that persicasulfide A (2), conferone (6) and feselol (7) showed moderate cytotoxicity with IC₅₀ values in the range of 11.76-39.24 μM against cancer cells and potent pan-HDAC inhibitory activity with IC₅₀ values in the range of 1.06-10.73 μM. Conferone was more active than others with a higher potency (for HDAC inhibition (1.06- 1.17 μM).

کلمات کلیدی:

Apiaceae, *Ferula latisecta*, *Ferula ovina*, *Ferula flabelliloba*, Histone deacetylase inhibitors, Cytotoxic activities

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