

## عنوان مقاله:

Moraea sisyrinchium inhibits proliferation, cell cycle, and migration of cancerous cells, and decreases angiogenesis in chick chorioallantoic membrane

## محل انتشار:

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## خلاصه مقاله:

**Objective(s):** Experimental studies reported that some plants in the genus of *Moraea* (Iridaceae family) show anticancer potential. This study aimed to evaluate the effects of *Moraea sisyrinchium* on U87 glioblastoma multiforme and HepG2 liver cancer cells. **Materials and Methods:** The cells were incubated for 24 hr with hydroalcoholic extract of the stem, flower, and bulb of *M. sisyrinchium*. Then, the cell proliferation (MTT) assay, cell cycle analysis (propidium iodide staining), cell migration test (scratch), Western blotting (Bax and Bcl-2 expression), and gelatin zymography (for matrix metalloproteinases, MMPs) were performed. Oxidative stress was evaluated by determining the levels of reactive oxygen species and lipid peroxidation. Angiogenesis was evaluated on chick embryo chorioallantoic membrane. **Results:** The extracts of the flower, stem, and bulb significantly decreased the proliferation of HepG2 and U87 cells. This effect was more for U87 than HepG2 and for the bulb and stem than the flower. In U87 cells, the bulb extract increased oxidative stress, cell cycle arrest, and the Bax/Bcl-2 ratio. Also, this extract suppressed the migration

ability of HepG2 and U87 cells, which was associated with the inhibition of MMP2 activity. In addition, it significantly reduced the number and diameter of vessels in the chorioallantoic membrane. Liquid chromatography-mass spectrometry revealed the presence of xanthenes (bellidifolin and mangiferin), flavonoids (quercetin and luteolin), isoflavones (iridin and tectorigenin), and phytosterols (e.g., stigmasterol) in the bulb. Conclusion: M. sisyrinchium bulb decreased the proliferation and survival of cancer cells by inducing oxidative stress. It also reduced the migration ability of the cells and inhibited angiogenesis.

### کلمات کلیدی:

Glioblastoma, Hepatocellular carcinoma, HepG2, Iridaceae, U87

### لینک ثابت مقاله در پایگاه سیویلیکا:

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