Bioactive Constituents of Conyza Albida

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Abstract: Alkynes and spathulenol were isolated from *Conyza albida* (Asteraceae); some of the compounds were lethal against *Artemia* sp. and cytotoxic against KB cells.

Introduction

Conyza albida Willd. ex Sprengel (Compositae) is a species growing in Argentina. Formerly, it was included as a synonym of *C. bonariensis* var. *microcephala* Cabr., but, there is enough evidence to consider it a valid entity at species level [1].

Conyza albida is reported to have expectorant, antitussive, and antiinflamatory activities [2,3]. Since *C. albida* usually grows together with *C. bonariensis* populations, it is believed that both species are useful in the treatment of urinary affections, liver diseases, stomach ulcers, and to wash sores [4] as well as an antihelmintic, digestive and diuretic [5,6,7].

There are no phytochemical studies, nor information on the active constituents of *C. albida*. We now present the results on the bioactivity-guided fractionation of an active extract of the leaves of *C. albida* and the evaluation of the activity of the pure compounds against *Artemia* sp., KB cells and as topoisomerase I inhibitors.

Experimental Procedures

Dry leaves of *Conyza albida* were extracted with CH_2Cl_2 . The total extract MeOH-H₂O 20% was partitioned between hexane, Et₂O, EtOAc and H₂O. All the extracts, including the water extract, were concentrated to dryness and tested in the brine shrimp toxicity test (BSTT). The hexane and Et₂O extracts gave positive results with $LC_{50} = 99 \ \mu g/ml$ and $LC_{50} = 96 \ \mu g/ml$, respectively. They were fractionated, guided by the BSTT, by vacuum liquid, centrifugal planar and preparative thin layer chromatographies. The isolates were identified by a combination of the following spectroscopic methods:

GC-MS, IR, UV, ¹H NMR and ¹³C NMR.

Results and Discussion

The ethyl ether extract afforded two bioactive fractions with similar chemical composition. After further purification the following compounds were identified: alkenynes **1**, **2** [8,9], **3** [8,9], and spathulenol **4** [10]. The hexane fraction contained alkenynes **1**-**3** and 1-dodecen-7,11-dimethyl-3-methylene [11] which was inactive. This is the first report on compound **1**, although the *trans* isomer was obtained by synthesis [12].

Compound	BSTT (µg/ml)	KB (µg/ml)	DNA Topoisomer- ase I (%)
	1.3	7.3	-
о осн ₃ Н Н 2	1.2	9.2	-
	5.2	19.1	21
	4.2	9	39

Positive controls: BSTT, berberine $LC_{50} = 8.4 \,\mu g/ml$; against KB cells, colchicine $IC_{50} = 0.02 \,\mu g/ml$.

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