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## IN VITRO EVALUATION OF ANTIMUTAGENIC ACTIVITY AND PHYTOCHEMICAL STUDY OF Baccharis trimera (Less.) DC. ESSENTIAL OIL

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Abstract: Carqueja, Baccharis trimera (Less.) DC. (Asteraceae), is a native species widely used in folk medicine as anti-inflammatory and to treat digestive problems. This study evaluated the chemical composition and the antimutagenic activity of the essential oil of Baccharis trimera (EOBt) through the in vitro micronucleus assay (MN). Moreover, as the antioxidant activity is one important mechanism of antimutagenic action [1], the antioxidant effect of EOBt was assessed by 2,2-diphenyl-1-picrylhydrazil (DPPH) and Oxygen Radical Absorbance Capacity (ORAC) assays [2]. Aerial parts of B. trimera (cv. CPQBA 01) were collected in CPQBA experimental field in July/2012 and the EOBt was obtained by hydrodistillation (3 h). The chemical composition was determined by gas chromatography coupled to mass spectrometry (GC/MS) indicating bicyclogermacrene (15.19%), E-caryophyllene (13.70%), germacrene D (8.69 %), β-pinene (7.54%), globulol (7.52%), β-myrcene (6.56%) and δ-cadinene (6.01%) as the major compounds of EOBt. In antimutagenic activity assay, CHO-K1 cells (2x10<sup>5</sup> cells/well) were treated with EOBt (25 µg/ml) and methylmetanesulfonate (MMS, 25 µg/mL) in three different schedules of treatment [simultaneous (EOBt + MMS, 4 h), pre (EOBt, 2 h + MMS, 2 h) and post (MMS 2 h + EOBt, 2 h) treatments]. Then, the cells were treated with cytochalasin B (3  $\mu$ g/ml, 20 h), sodium citrate 1% and methanol/acetic acid 3:1 for slides preparation. Replication Index (RI), Cytokinesis-Block Proliferation Index (CBPI) and MN frequency (%) in binucleate cells were calculated and the statistical analysis was performed by ANOVA, post test Duncan (p < 0.05) [3]. All samples showed RI > 90% and CBPI > 1.7 (absence of cytotoxicity). Compared to untreated cells  $(1.07 \pm 0.30\%)$ , MMS increased MN frequency to 5.77  $\pm$  0.14% in simultaneous treatment and to 5.11  $\pm$  0.13% in pre and post-treatments. In the simultaneous treatment, EOBt decreased by 65.8% the frequency of MNs in relation to the positive control (MMS). In the post-treatment, there was a reduction of 54.5% in MN incidence while in pretreatment with EOBt, the reduction was only 35.1%. The antioxidant evaluation showed  $18.76 \pm 0.53$  and  $2927.94 \pm 14.80 \ \mu$ M of Trolox equivalent antioxidant capacity/g of EOBt in DPPH and ORAC assays, respectively. These results suggest that EOBt does not show a preventive effect, but may be more efficient to reverse damage caused by genotoxic or mutagenic agents. Moreover, this antimutagenic activity could be partially attributed to an antioxidant effect. (FAPESP grant #2013/13196-0, CAPES).

## References

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