INFLUENCE OF TEMPERATURE ON VOLATILE COMPOSITION OF GUACO

(M. laevigata and M. glomerata)

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ABSTRACT

Mikania laevigata and Mikania glomerata are popularly known as guaco and are mainly used for respiratory complications. As both are included in the 1st Phytotherapeutic of Brazilian Pharmacopoeia (2011), apparently being used indiscriminately, they will be studied in parallel. The aim of this study was to evaluate how the volatile compounds vary in different conditions of temperature. Clones of both species were submitted to 15 °C, 25 °C and 35 °C for 3 weeks. Volatile compounds were extracted by solid phase microextraction (SPME) and analyzed by GC-MS. By conducting an exploratory analysis of principal components (PCA), it was observed that the two species have very different chemical composition. Thus, M. laevigata and M. glomerata apparently cannot be used indiscriminately. Regarding the temperature treatment, there was no separation between the tests, indicating that this factor may not influence the volatiles composition of these species.

Keywords: Guaco, GC-MS, Volatiles compounds, Temperature.

RESUMO

Mikania laevigata e Mikania glomerata são popularmente conhecidas como guaco e utilizadas principalmente para complicações do trato respiratório. Como ambas constam no 1º Fitoterapic do Brazilian Pharmacopoeia (2011), aparentemente podendo ser usadas indiscriminadamente, elas serão estudadas em paralelo. O objetivo desse trabalho foi avaliar como os compostos voláteis
variam em diferentes condições de cultivo de temperatura. Clones das duas espécies foram submetidas a 15°C, 25°C e 35°C durante 3 semanas. A identificação dos voláteis se deu por microextração em fase sólida (SPME) e as análises por GC-MS. Realizando uma análise exploratória de componentes principais (PCA), observou-se que as duas espécies apresentam composição química bem diferente. Desse modo, *M. laevigata* e *M. glomerata* aparentemente não podem ser utilizadas indiscriminadamente. Quanto ao tratamento de temperatura, não houve uma separação entre os testes, indicando que esse fator pode não influenciar na composição dos voláteis dessas espécies.

**Palavras-chaves:** Guaco. GC-MS. Compostos voláteis. Temperatura.

*Mikanialaevigata* Schultz Bip. ex Baker and *Mikaniaglomerata*Spreng are Brazilian medicinal plants, also known as guaco (CASTRO, 2003). Both are included in the 1st Brazilian Phytotherapic Pharmacopoeia (ANVISA, 2004), and are indicated for anti-inflammatory diseases (OLIVEIRA; OGA; AKISUE, 1985), anti-allergic (FIERRO, *et al.*, 1999) and for bronchodilator action (LEITE, 1993). These activities are due to coumarin (1,2-benzopiron) (CELEGHINI; VILEGAS; LANÇAS, 2001), considered the chemical marker for this species. However, many studies show that this compound varies greatly between *M. laevigata*and *M. glomerata* (ANJOS, 2009; BOLINA; GARCIA; DUARTE, 2009; BERTOLUCCI, 2009). Our study group found high concentration of coumarin only in *M. laevigata* and much lower in *M. glomerata* (MELO; SAWAYA, 2015 – in press).

Besides the variation of the marker (coumarin) between the two species, studies indicate that variation occurs when plants are submitted to different treatments (ALMEIDA, 2015; BERTOLUCCI, 2013). Almeida, *et al.* (2017) compared these same two species and the results indicated that he highest content of coumarin was found in extracts of *M. laevigata*under 50% shade, at temperatures of 10 oC and/or 22 oC, and under lower water availability condition.
Therefore, this work aims to compare the volatile compounds of *M. laevigata* and *M. glomerata* under different temperature conditions. Furthermore, there are only few studies about volatile composition and even less about its variation under different environment conditions.

Clones of *M. laevigata* and *M. glomerata* were used to minimize genetic variation within the treatments. Vases with the clones were placed in growth chambers with controlled temperatures at 15°C, 25°C (control) and 35°C. The light intensity of the chamber was maintained at 180 mols photons m-2 s-1 using red and blue LED lights. Ten replicate for each species were made for each temperature. The pots were kept for three weeks under treatment.

Volatile compound were extract with solid phase micro-extraction (SPME) and the analysis were carried by gas chromatography with mass spectrometry (GC-MS). All extractions were carried out with a polydimethylsiloxane/divinylbenzene (PDMS/DVB) fiber, 65 µm film thickness (Supelco, Bellefonte, PA). Agilent GC 7890/ MS 5975 with Gerstelinjetor (MPS) (Agilent Technologies, EUA) was used for the analysis.

A quality control (QC) was made to ensure the equipment operation. For that, all samples from all treatments and all replicates were mixed to compose the QC. 20-mL vial with 1g of ground material were kept at 70°C for 5 minutes. Extraction time with the fiber was 10 min. The fiber was kept in the injector for 3 min for the optimal thermal desorption. The injection in 1:5 split mode used helium as carrier gas at a flow rate of 1.0 mL min-1. The injector temperature was 240 °C. A fused-silica capillary column (5% phenyl–95% polydimethylsiloxane, 30 m × 0.25 mm,0.25 µm) was employed in the separation of the compounds. The oven temperature was programmed from 100 °C to 170 °C at a rate of 3.5 °C min-1 and 170 °C to 250°C at a rate of 20 °C min-1. The mass spectrometer was used with electron ionization (70 eV) and mass scan range from 29 to 289 Da. The temperatures of the ion source and the GC-MS interface were 200 and 250 °C, respectively. Compounds were identified by comparing their mass spectra with the GC-MS spectral library (NIST, 2013).
Due to the large volume of data, it was not possible to differentiate the samples only by chromatogram analysis. Therefore, an exploratory data analysis was carried out by principal component analysis (PCA). The PCA was conducted using the autoscaled relative area of the compounds identified in each sample. The PCA routines were performed using the MetaboAnalyst 3.0 (XIA; WISHART, 2016).

In sample chromatograms it was possible to identify several volatile organic compounds extracted via SPME. The major volatile compound is Germacrene D, representing on average 44.9% of total area. Other mono and sesquiterpenes such as α-pinene, δ-elemene, α-humulene, δ-cadineno where identified too. Figure 1 shows a typical chromatogram of the guaco samples. All chromatograms from *M. laevigata* presented high peaks of coumarin, considered the chemical marker from this specie (ANVISA, 2004). On the other hand, none of the samples of *M. glomerata* showed the peaks of coumarin. Similar results were found by many others authors (MELO; SAWAYA, 2015; ALMEIDA, 2015; BERTOLUCCI, 2013).

**Figure 1.** Typical chromatogram obtained by SPME-GC/MS from *M. laevigata* samples

The PCA analysis (Figure 2) showed a good separation between the two species and the QC, indicating that *M. laevigata* and *M. glomerata* have different chemical composition. Figure 2a. Represents the scores (samples), in green are the *M. laevigata*, in red *M. glomerata* and the QC are colored in blue. In the loading plot (Figure 2b.), it is not clear which variables are responsible for
the samples separation. PC 1 explains 17.7% of variation and PC 2 only 9.5%. The large amount of data could be the reason of this low explanation and for the bad separation of the samples.

Regarding the temperature treatment, there was not a clear separation between the tests, indicating that this factor may not influence the volatiles composition of these species. It was observed that after the treatment, all leaves from all tests were damaged, with different colored areas (red to purple). Perhaps the artificial light used in the growing chamber affected the plant leaves, once the clones were young.

The present study demonstrated that *M. laevigata* and *M. glomerata* have differences between the volatile composition. PCA scores plot clearly divided the samples in three main groups, proving that the species are different and cannot be used without distinction. The temperature treatment didn’t show good separation between the tests. Maybe this factor does not influence the volatile composition. Further studies must be done to detect which variables are responsible for the species discrimination.

**Figure 2.** Principal Components Analysis of the main peak areas found in the GC–MS analysis of the temperature treatment samples (A) Scores plot (samples), *M. laevigata* colored in green, *M. glomerata* in red and QC in blue (B) Loadings plot; PC1 x PC2.
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