INTRODUCTION

Turmeric (Curcuma longa L., Zingiberaceae), whose scientific synonyms are C. domestica Valeton and Amomum curcuma Jacq., is a rhizomatous herbaceous perennial plant. This species is native to Southeast Asia and extensively cultivated in the tropical and subtropical regions of the world. In India, and in several other countries, it is one of the most important spices used as a natural yellow food pigment and in herbal medicine.

In the traditional systems of Asian medicine, turmeric has been indicated for the treatment of digestive disorders, ocular infection and inflammation, diarrhea, epilepsy, wound healing, fever, allergies, chronic cough, bronchial asthma, jaundice, arthritis and other diseases.

The biological activities of turmeric have been investigated in recent decades. Turmeric powder and crude extracts have been evaluated for some pharmacological activities such as hepatoprotective, antifungal, neuroprotective and memory improvement.

Regarding its chemical composition, the major secondary metabolite classes include curcuminoids and sesquiterpenes. Curcumin, the primary curcuminoid, represents 3%–5% of turmeric. Essentially, curcumin is a commercially available mixture of curcuminoids that contains 72%–78% of curcumin, 12%–18% of demethoxycurcumin, 3%–8% of bisdemethoxycurcumin and organic solvent residue. Curcumin can modulate multiple pathways, which can explain the diversity in its traditional indications and pharmacological activities. Furthermore, the safety of using curcumin has been demonstrated in clinical trials even at doses >8 g/day. Unfortunately, the efficacy could be questioned due to the small number of patients involved.

Low bioavailability of curcumin is another problem, and many papers are dealing with subject. Researchers have explored a few successful alternatives to increase the serum levels of curcumin, such as nanotechnology, structural analogues, association with sesquiterpenes derived from essential oil, extraction with water and enzymes from fresh rhizomes, encapsulate curcumin into chitosan, association with piperine, the major component of black pepper that increases bioavailability by 2000%. Besides, curcuminoids are also sensitive to light and air.

Dried rhizomes and leaves contain approximately 5%–6% and 1.0%–1.5% of volatile oil, respectively. The essential oils of rhizomes are constituted primarily by sesquiterpenes, typically α-turmerone, curcule (= β-turmerone), α-turmerone, γ-turmerone and β-sesquiphellandrene, and the monoterpenes β-pinene and para-cymene. In addition, studies have reported that the essential oil has anti-inflammatory, anti-nociceptive and anti-atherosclerotic in vivo effects and antioxidant, anti-proliferative and anti-angiogenic in vitro activities. Therefore, although curcuminoids are the most investigated compounds, currently, essential oil components have demonstrated significant results in pharmacological studies.

Several factors can interfere with the chemical variation of essential oils in plants, such as temperature, humidity, luminosity, altitude, pluviometry, ultraviolet radiation, soil and nutrient conditions, seasonality, circadian cycle, method of collection, drying and part of the plant. Due to these factors, the relative composition of turmeric varies considerably with the geographical origin and different agroclimatic zones. The composition of turmeric essential oil also can vary with maturity. As the plant ages, the concentrations of sesquiterpenes increase, whereas those of monoterpenes decline in the rhizomes. The maximum curcumin content was found at the age of approximately 9 months after planting, after which there was a decline in the total curcumin content. Furthermore, the essential
oil content positively correlates with the levels of phosphorous and potassium in the soil.\textsuperscript{35} Regarding the effect of solar radiation on the essential oil yield, \textit{C. longa} cultivated under full sunlight treatment showed a greater yield (3.0\%) than that under treatment with shading (50\% of sunlight) (1.90\%).\textsuperscript{37}

This work describes variation of the yield and chemical composition of the Brazilian \textit{C. longa} rhizomes. The results can be important for the quality control of herbal medicine produced in the savanna (Cerrado) region of Bahia.

**EXPERIMENTAL**

**Plant material**

About one kilo of each sample of turmeric rhizomes was collected in February 2011 from three commercial producers (Producer 1 São Manoel Farm, producer 2 Manchão Branco Farm, and producer 3 Raimundo Farm) in Jaborandi City (13°37’10’’S, 44°25’58’’W) located in the Brazilian savanna (Cerrado) biome. Twelve samples were collected in the late afternoon in a period of occasional rainfall. Rhizomes were sliced, dried at 40 °C for 10 h and stored in paper bags. All plant materials were identified by Prof. Patricia Baier Krepsky (Multidisciplinary Institute in Health, Federal University of Bahia, Brazil) based on the macroscopic and microscopic description presented in the Brazilian Pharmacopoeia.\textsuperscript{38} Samples have been deposited at the vegetable drug collection of the Federal University of Paraná with the numbers 122A10, 122A11, 122A12 and 122A13 (producer 1); 122A14, 122A15, 122A16 and 12A17 (producer 3) and 22A18, 122A19, 122A20 and 122A21 (producer 2). For this study, the authors obtained authorisation of access to the Brazilian System for the Management of Genetic Heritage and Associated Traditional Knowledge--SISGEN (# AB58325).

**Variability in the essential oil yield**

To determine the essential oil yield, five grams of each sample with about 100 mL of water was subjected to the hydro-distillation method for 4 h, using a Clevenger-type apparatus with 0.5 mL of xylene to solve the essential oil extracted, according to the Brazilian Pharmacopoeia procedure.\textsuperscript{39} After extraction, the essentials oils were measured directly in the extraction apparatus, and the content (\%) was calculated as volume (mL) of essential oil per 100 g of dry plant material. These extractions were done in triplicate for each sample 1 month after the collection. All samples were triturated for no more than 1 day before analysis because of the sensitivity of the turmeric components. Average, standard deviation and relative standard deviation were calculated for the group of samples obtained from each producer as well as for the total number of collected samples. Comparisons between results obtained for each producer’s samples were performed by the analysis of variance (ANOVA), applying Tukey’s test to compare mean values, using the GraphPad Prism software, version 5. These results were compared with the essential oil yield recommended by the Brazilian Pharmacopoeia.\textsuperscript{38} Essential oils were stored in the freezer before conducting further chromatographic analysis.

**Analysis of the chemical profile of essential oil and curcumin by thin-layer chromatography**

Thin-layer chromatography (TLC) was used to verify the essential oil profile using silica gel GF254 of approximately 250-nm thickness.\textsuperscript{30} The mobile phase was optimised previously to hexane/ethyl acetate (9:1), and the essential oils obtained from all the 12 samples were diluted in xylene (1:1) before spotting onto the TLC plates. The plates were visualised by spraying anisaldehyde solution.

The chemical profiles were also compared qualitatively using a commercial mix of curcuminoids (curcumin, demethoxycurcumin and bisdemethoxycurcumin, Sigma Aldrich\textsuperscript{36}) as reference substances. Rhizomes were powdered and sequentially the samples were prepared with 0.5 g in 5 mL of methanol, under stirring for 30 min, centrifuged at 2500 rpm for 10 minutes and, filtered. Reference solution was prepared by using 5 mg of curcuminoids (purchased from Sigma Aldrich\textsuperscript{36}) in 5 mL of methanol. Samples and reference solution were spotted on silica gel plates GF254. Mobile phase consisted of chloroform, ethanol, and acetic acid (95:5:0.5). After development, plate was removed, dried and, spots were visualized in UV light (366 nm).\textsuperscript{38}

**Qualitative and quantitative analysis by gas chromatography coupled to a mass spectrometer**

The essential oils were analysed using a Shimadzu\textsuperscript{37} QP2010 gas chromatography apparatus directly interfacing with mass (MS), equipped with a 5% diphenyl 95% dimethyl polysiloxane capillary column (30 m × 0.25 mm, 0.25 μm of film thickness). Helium was used as the carrier gas. The mass detector of the GC-MS equipment was operated in an electron-impact mode, with a scan range of 45–500 amu, an ionisation energy of 70 eV and a scan rate of 0.30 s per scan. The temperatures of the ionisation source and the injector were maintained at 200 °C and 240 °C, respectively. Other GC conditions such as the flow rate, the concentration of the sample and the separation temperature programme were optimised to provide a better separation of components in a shorter run time. Chemical constituents were identified by referring to compounds reported in the literature and by comparing their mass spectra with those of known compounds available in the database of NIST 2008 (National Institute Standard and Technology) and Flavour and Fragrances of Natural and Synthetic Compounds (FFNSC) 1.3 libraries. The identification was further supported by the calculation of their retention indices under identical experimental conditions using n-alkanes (C10-C40) series (Sigma Aldrich, USA). Retention Index (RI) was calculated for each main compound and compared to those reported in the literature data.

The relative percentage of each compound present in rhizome oil was calculated using the corresponding peak area integration, performed automatically using their own software (GCMSolution\textsuperscript{38}). A cheque of the integration of each peak was conducted and corrected manually if necessary. The composition was reported as a relative percentage of the total peak area.

Average, standard deviation and relative standard deviation (%) of the relative proportion areas (%) were calculated for each major peak identified and their respective retention times.

**Plant drug stability**

Analyses were repeated for five samples 6 months later to verify the chemical stability. A paired Student’s t-test was used to assess whether the samples had stability in terms of the essential oil yield and the relative area of the major oil components by gas chromatography. Besides, the effect of the Clevenger extraction on the volatile compounds were evaluated analysing the composition of these volatiles after extraction of 1 g of three samples with 20 mL hexane. The extract was injected in the GC-MS employing the same conditions.

**RESULTS AND DISCUSSION**

**Variability in the essential oil yield**

\textit{C. longa} populations collected from three different producers...
were investigated for their essential oil yield and chemical composition. Extractions of essential oils from each sample were carried out in triplicate, and their average yield of oils obtained from all samples was 3.97% ± 0.61% and the variation among the producers was 3.0%–5.16% (relative standard deviation = 15.3%) in the dry material. The average yield for each producer was as follows: producer 1, 3.83% ± 0.17% (4.3%); producer 2, 4.58% ± 0.58% (14.7%) and producer 3, 3.48% ± 0.59% (16.9%). Comparisons between producers performed by ANOVA showed significant differences. A significant difference (p = 0.05) was found only between producers 2 and 3 (Tukey’s test). Despite the significant variation in oil yields among the producers, all the analysed samples had an oil yield within the values specified by the Brazilian Pharmacopoeia. Therefore, the turmeric samples grown in the Brazilian savanna region of Bahia state are suitable for use in herbal medicine preparations, considering that the essential oil content should be at least 2.5% as recommended by the Brazilian Pharmacopoeia.

According to earlier studies, *C. longa* essential oil yield in dry rhizomes varied from 1.5% to 5.0%. Similarly, other researchers found 2.9% and 3.8% of essential oil yields. Similar variations ranging from 2.1% to 4.4% were found in Brazil. Therefore, *C. longa* cultivated in the Brazilian savanna could produce high-quality turmeric in terms of the essential oil yield.

In India, some researchers have reported essential oil yields between 0.61% and 1.45% in the fresh rhizomes of 27 accessions from the northern part of the country and between 0.37% and 0.8% on different agroclimatic zones.

### Analysis of the chemical profile of essential oil and curcumin by TLC

TLC analysis of the essential oil samples resulted in at least four main violet spots. The chromatographic profiles demonstrated significant similarity among all samples. Subsequent analysis of the methanol extract revealed the presence of three major curcuminoids, which is consistent with that described in the Brazilian Pharmacopoeia.

### Qualitative and quantitative analysis by gas chromatography coupled to a mass spectrometer

After testing and evaluation of several different conditions of chromatographic parameters (concentration of the sample, inlet mode, flow rate and separation temperature programme), the best parameters were as follows: a flow rate of 1.5 mL min⁻¹, a concentration of 2.5% of the sample in dichloromethane and a temperature programme as described in Table 1. Using this developed method, it was possible to separate all the substances satisfactorily within 20 min as shown in Figure 1. This method proved to be more advantageous than other methods described in the literature because of the shorter analysis time.

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Seven compounds were identified in the essential oil by comparing the RI (literature and RI taken from NIST 2008 library) and mass spectra of each peak with the NIST and 1.3 FFNSC libraries (similarity >90%). The major compounds identified were ar-turmerone (5), turmerone (α-turmerone) (6) and curlone (β-turmerone) (7). Some minor components were also identified, including α-phellandrene (1), α-curcumene (ar-curcumene, curcumene) (2), β-bisabolene and (3) β-sesquiphellandrene (4) (Table 2).

Analyses of the samples were performed by peak area normalisation. Table 2 shows the average, standard deviation and the relative standard deviation (RSD%) values of the relative proportion area % (RPA).

Table 3 shows the results of the comparison between the relative percentages of *C. longa* essential oil obtained in this study and those reported in the literature using MS. This comparison revealed that the major compounds reported in the literature were similar to those observed in the present study.

In order to verify if there was no conversion of turmerol in

![Figure 1. Essential oil GC-MS chromatogram of C. longa rhizomes](image)
turmerone, extraction with hexane solvent was carried out by maceration at room temperature. GC-MS analysis of this extract showed the peak of ar-turmerone and curlone but not of turmerone. These results indicate the possibility of the conversion of turmerol to turmerone may have occurred during the hydrodistillation processes. However, previous studies in literature do not mention this type of conversion.

**Plant drug stability**

As mentioned earlier, the average volatile oil yield was 3.97%. The repeated analysis performed after 6 months showed an average yield of 3.50% with variations between 2.98% and 3.94% (RSD%: 10.42). Although there was a significant reduction (p = 0.06, t-test) in the volatile oil yield, the remaining were within the standards recommended by the Brazilian Pharmacopoeia. The chromatographic profile analysed by TLC remained similar. Concerning about the analysis by GC-MS, from a qualitative point of view, the only difference was the absence of α-phellandrene. However, the percentage of this compound was too small since the first testing of each sample (average 0.35%). On the basis of all these results, it can be concluded that the plant drug exhibited satisfactory stability after the 6-month storage period, despite the small decrease in the essential oil content.

**CONCLUSIONS**

The yield (3.97% ± 0.61%) and the chemical composition of the essential oils obtained from the rhizomes of *C. longa* were similar to those of high-yield accessions described in the literature. Therefore, based on the parameters analysed in this study, the samples found in the Brazilian savanna region fulfilled the quality requirements. Moreover, the primary advantage of the current method of GC-MS analysis was its shorter analysis time, i.e. only 20 min. Regarding stability, the content was maintained adequate and the chromatographic profile remained similar even 6 months after harvest.

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**REFERENCES**

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Essential oil of Curcuma longa L. rhizomes chemical composition, yield variation and stability


