

# Physicochemical properties and mineral and protein content of honey samples from Ceará State, Northeastern Brazil

*Propriedades físico-químicas, minerais e teor de proteínas em amostras de méis do Estado do Ceará, nordeste do Brasil*

Maria da Conceição Tavares Cavalcanti LIBERATO<sup>1\*</sup>, Selene Maia de MORAIS<sup>2</sup>, Carlos Emanuel de Carvalho MAGALHÃES<sup>1</sup>, Islay Lima MAGALHÃES<sup>1</sup>, Daniel Bomfim CAVALCANTI<sup>1</sup>, Marina Maciel de Oliveira SILVA<sup>1</sup>

## Abstract

This study evaluated the physicochemical properties and protein and mineral content of honey samples from Ceará State, Northeastern Brazil, one of the major honey exporters in the country. Nutritional importance of the minerals detected was also analyzed. Physicochemical properties were examined according to the AOAC and CAC official methods; the protein content was determined using the Bradford method, and the minerals were analyzed by atomic absorption spectrometry. All analyses were performed in triplicate. The levels of macrominerals sodium (Na), potassium (K), calcium (Ca), and magnesium (Mg) varied from 1.80-47.20, 21.30-1513.30, 14.58-304.82, and 2.48-28.33 mg/kg, respectively, and the trace elements iron (Fe), copper (Cu), manganese (Mn), zinc (Zn), selenium (Se), and chromium (Cr) varied from 0.12-8.76, 0.07-1.29, 0.06-1.96, 0.07-1.85 mg/kg,  $0.36 \times 10^{-3}$ - $62.00 \times 10^{-3}$  and  $22.50 \times 10^{-3}$ - $170.33 \times 10^{-3}$  µg/kg, respectively. *Myracrodruon urundeuva* honey sample had high contents of macrominerals (Na, K, Ca, and Mg). Protein content of the *Anacardium occidentale* honey sample was the highest (1121.00 µg/g) among the samples analyzed. Among the minerals detected in the honey samples, K showed the highest concentration, followed by Ca, Na, and Mg. The presence of trace elements can show environmental contamination. The honey samples studied were free of trace elements contamination, except for Mn; the *Piptadenia moniliformis* was the only honey sample that was in compliance with the law requirements. The variations of the chemical constituents in the honey samples are probably related to differences in the floral origin and mineral and protein contents and confirm the nutritional importance of Ceará State honey.

**Keywords:** *Apis mellifera* honey; floral origin; nutritional importance.

## Resumo

Este trabalho avaliou propriedades físico-químicas, teores de proteína e minerais em méis do Ceará, um dos principais exportadores do País. Também foi analisada a importância nutricional com relação aos minerais detectados. As propriedades físico-químicas foram determinadas pelos métodos da AOAC e CAC, o conteúdo de proteína pelo método de Bradford e os minerais por espectrometria de absorção atômica. Todas as análises foram feitas em triplicata. Os níveis dos macrominerais sódio (Na), potássio (K), cálcio (Ca), e magnésio (Mg) variaram de 1,80-47,20, 21,30-1513,30, 14,58-304,82 e 2,48-28,33 mg/kg, respectivamente, e os elementos traços ferro (Fe), cobre (Cu), manganês (Mn), zinco (Zn), selênio (Se), e cromo (Cr) variaram de 0,12-8,76, 0,07-1,29, 0,06-1,96, 0,07-1,85 mg/kg,  $0,36 \times 10^{-3}$ - $62,00 \times 10^{-3}$  e  $22,50 \times 10^{-3}$ - $170,33 \times 10^{-3}$  µg/kg, respectivamente. A amostra de *Myracrodruon urundeuva* apresentou altos teores dos principais minerais (Na, K, Ca, e Mg). O teor de proteínas para o mel de *Anacardium occidentale* foi o maior (1121,00 µg/g) entre as amostras de méis analisadas. Entre os minerais, detectados, K apresentou a maior concentração seguido por Ca, Na, e Mg. A pesquisa de elementos traços pode indicar contaminação ambiental. As amostras de méis estudadas apresentaram-se livres de contaminação por elementos traço, excetuando-se quanto ao Mn, em que apenas a amostra proveniente de *Piptadenia moniliformis* apresentou-se dentro da legislação. As variações dos constituintes químicos devem-se provavelmente às variadas origens florais e os teores de proteínas e minerais ratificam a importância nutricional dos méis do Ceará.

**Palavras-chave:** mel de *Apis mellifera*; origem floral; importância nutricional.

## 1 Introduction

Honey is rich in properties that result from its chemical composition. It is not considered a complete food according to human nutritional standards, but it does offer potential supplement for infants, senior citizens, and convalescents as an easily digestible foodstuff that may be ingested directly or used as a sweetener in a variety of products.

Honey composition is variable due to the influence of plants, climate, and environmental conditions and the ability of the beekeeper. The variation of the physicochemical properties of honey depends on the nectar and pollen of the original plant, color, moisture, and protein and minerals contents (WHITE JUNIOR, 1978). Therefore, honey is related to its

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<sup>1</sup> Laboratório de Bioquímica e Biotecnologia – LABBIOTEC, Coordenação do Curso de Licenciatura em Química, Centro de Ciências e Tecnologia – CCT, Universidade Estadual do Ceará – UECE, Av. Paranjana, 1700, CEP 60740-000, Fortaleza, CE, Brasil, e-mail: liberato@secrel.com.br

<sup>2</sup> Laboratório de Produtos Naturais – LPN, Coordenação do Curso de Licenciatura em Química, Universidade Estadual do Ceará – UECE, Av. Paranjana, 1700, CEP 60740-000, Fortaleza, CE, Brasil

\*Corresponding author

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botanical origin, processing and storage, and climatic factors that occur during the flow of nectar, and to the temperature at which the honey matures in the hive (SEEMANN; NEIRA, 1988). Brazil is an important producer and exporter of *Apis mellifera* honey due to botanical diversity and differentiated vegetation that provides blossoming plants all year long.

The method described by Bradford was chosen for the quantitative determination of proteins. According to Azeredo et al. (2003), this method eliminates most of the problems involved in the other procedures.

The main minerals present in honey originate from soil and are transported to trees by the roots. The minerals make their way into the nectar and are subsequently incorporated into the honey produced by bees (STANKOVSKA; STAFILOV; SANJ, 2008). Accordingly, the composition and the content of metals in honey, particularly major and minor metals, is affected by the composition determined by geochemical and geological features (FERNÁNDEZ-TORRES et al., 2005). Since soil and plants are natural sources that have a great influence on the mineral composition of honey, information on the metal profile is suitable for categorizing honeys according to their floral and geographical provenance (POHL, 2009). There is little information about the inorganic content of Brazilian honey because usually the organic components or some physicochemical properties are investigated to characterize the honey. A lot of studies on the level of mineral in bee products are focused on investigating environmental pollution effects on these products quality since they have an important role as indicators of pollution in the areas where they are produced (CONTI; BOTRE, 2001). Honey samples originating from polluted sites, from flowers that provide nectar and pollen pellets, are subject to contamination by heavy metals from emissions of gases and particles (MAGALHÃES, 2010). These minerals present in bee products in levels above those permitted by legislation represent a threat to humans due to their negative and cumulative effects of contaminants on human health. Arsenic exposure, for example, increases the risk of cancer, diabetes, and hypertension among others diseases (DESCHAMPS; MATSCHLLAT, 2007). Heavy metals that can contaminate bee products are: lead (Pb), copper (Cu), chromium (Cr), manganese (Mn), nickel (Ni), zinc (Zn), silver (Ag), iron (Fe), cadmium (Cd), aluminum (Al), cobalt (Co), and strontium (Sr). Their levels in honey samples can indicate the degree of environmental pollution and the geographic origin of the honey samples (FRANCHINI et al., 2007).

This study investigated several honey samples of different floral origins from Ceará State, including honey samples from important Brazilian medicinal plants such as *Lippia sidoides* Cham. and *Myracrodruon urundeuva* Fr. All. (Table 1). *L. sidoides* (Verbenaceae) is a shrub native to semi-arid northeastern Brazil. *M. urundeuva* (Anacardiaceae) is native to northeastern Brazil, extending to São Paulo and Mato Grosso do Sul. It occurs widely in semi-arid and also in dry and subhumid forests. *L. sidoides* and *M. urundeuva* are widely used medicinal plants in northeastern Brazil, and honey samples from their flowers showed high total phenolic content, and antioxidant antiacetylcholinesterase activities (LIBERATO et al., 2011). The

consumption of foods with therapeutic benefits is an effective way to achieve a healthy lifestyle. Since a balanced diet is related to minerals and protein content in food, it is further studies are important to elucidate the potential of Ceará honey.

## 2 Materials and methods

### 2.1 Honey samples

Twenty-two samples of *Apis mellifera* honey from different floral origins were obtained from apiarists and beekeepers' associations in different places in Ceará State. Each honey sample was identified in terms of its floral origin and site of collection based on information obtained from the suppliers. The honey samples were derived from different botanical sources including: *Hyptis suaveolens* Poit. (S02), *Anacardium occidentale* L. (S04), *Spermacoce verticillata* L. (S05), *Mimosa verrucosa* Benth. (S08), *Piptadenia moniliformis* Benth. (S09), *Myracrodruon urundeuva* Fr. All. (S15 e S19), *Licania rigida* Benth. (S17), *Lippia sidoides* Cham. (S18), *Serjania* sp. (S20), and *Ziziphus joazeiro* Mart. (S22). The honey samples S01, S03, S06, S07, S10, S11, S12, S13, S14, S16, and S21, were heterofloral (Table 1). The samples were collected between July 2007 and March 2009, during the dry and rainy seasons. All samples were transferred to the laboratory, stored in amber flasks, and kept at 4-5 °C until analysis.

### 2.2 Moisture, ash, free acidity, pH, and insoluble solids

All analyses were carried out in triplicate. Moisture was determined according to the AOAC official method (ASSOCIATION..., 1998) using a standard Abbe-type refractometer at 20 °C. Water content (%) was obtained from the Chataway table.

Ash content was determined according to the CAC method (CODEX..., 1990). Honey samples of 5 g were incinerated in a muffle furnace (Quimis, São Paulo, Brasil) at 550 °C. A pH meter was used to determine the pH, and the acidity was determined according to the AOAC method.

The gravimetric method was used for the determination of the insoluble solids according to the CAC method. Honey samples (20 g) were diluted with the minimum amount of water at 80 °C and transferred to porous previously weighed crucibles. After this procedure, the samples were flushed with distilled water at 80 °C to remove all sugars. The porous crucibles were placed in an oven at 135 °C for 1 hour and then cooled and weighed.

### 2.3 Analysis of honey color intensity

Honey color intensity was determined using a Spectrophotometer (model Thermo Biomate 5, New York, USA). The honey samples were diluted to 50% (w/v) with warm water (45-50 °C), sonicated for 5 minutes, and filtered to eliminate large particles. The net absorbance was defined as the difference between the spectrophotometric absorbance at 450 and 720 nm (BERETTA et al., 2005).

**Table 1.** Honey samples, place of collection, period of collection, and local vegetation.

Samples	Place of collection*	Period	Common vegetation
Heterofloral (S01)	Parambu	July 2008	<i>Bumelia sartorum</i> Mart., <i>Croton sonderianus</i> Muell. Arg., <i>Hyptis suaveolens</i> Poit., <i>Mimosa caesalpiniaefolia</i> Benth., <i>Mimosa scabrella</i> Benth., <i>Mimosa tenuiflora</i> (Willd.) Poir.
Heterofloral (S02)	Meruoca	May 2008	<i>Mimosa verrucosa</i> Benth., <i>Borreria verticillata</i> Mayer, <i>Malpighia glabra</i> L.
Heterofloral (S03)	Itapiuna	October 2008	<i>Croton sonderianus</i> Muell. Arg., <i>Hyptis suaveolens</i> Poit., <i>Bumelia sartorum</i> var. <i>latifolia</i> , <i>Mimosa caesalpiniaefolia</i> Benth., <i>Mimosa tenuiflora</i> (Willd.) Poir.
<i>M. verrucosa</i> (S04)	Ibiapina	August 2008	<i>Mimosa verrucosa</i> Benth.,
Heterofloral (S05)	Morada Nova	September 2008	<i>Merremia aegyptia</i> L. Urban, <i>Mimosa hostilis</i> (Willd.) Poir., <i>Hyptis suaveolens</i> Poit., <i>Croton sonderianus</i> Muell. Arg.
<i>P. moniliformis</i> (S06)	Pindoretama	April 2008	<i>Piptadenia moniliformis</i> Benth.
<i>Serjania</i> sp. (S07)	Crato	June 2008	<i>Serjania</i> sp.
<i>M. urundeuva</i> (S08)	Tauá	November 2007	<i>Myracrodruon urundeuva</i> Fr. All.
<i>A. occidentale</i> (S09)	Pacajus	January 2008	<i>Anacardium occidentale</i> L.
<i>H. suaveolens</i> (S10)	Morada Nova	June 2008	<i>Hyptis suaveolens</i> Poit.
<i>S. verticillata</i> (S11)	Horizonte	April 2009	<i>Spermacoce verticillata</i> L.
Heterofloral (S12)	Canindé	February 2008	<i>Alternanthera brasiliana</i> (L.) Kuntze, <i>Bumelia sartorum</i> var. <i>latifolia</i> , <i>Croton campestris</i> St. Hill., <i>Croton sonderianus</i> Muell. Arg., <i>Mimosa tenuiflora</i> (Willd.) Poir., <i>Mimosa caesalpiniaefolia</i> Benth.
Heterofloral (S13)	Barbalha	July 2008	<i>Mimosa caesalpiniaefolia</i> Benth., <i>Mimosa verrucosa</i> Benth., <i>Serjania</i> sp., <i>Borreria verticillata</i> Mayer, <i>Combretum leprosum</i> Mart., <i>Mimosa tenuiflora</i> (Willd.) Poir., <i>Mimosa scabrella</i> Benth.
Heterofloral (S14)	Ibaretama	April 2009	<i>Licania rigida</i> Benth., <i>Mimosa caesalpiniaefolia</i> Benth., <i>Mimosa tenuiflora</i> (Willd.) Poir., <i>Bumelia sartorum</i> var. <i>latifolia</i> Miq.
Heterofloral (S15)	Irauçuba	April 2009	<i>Bumelia sartorum</i> var. <i>latifolia</i> Miq., <i>Croton sonderianus</i> Muell., <i>Hyptis suaveolens</i> Poit., <i>Mimosa scabrella</i> Benth., <i>Mimosa caesalpiniaefolia</i> Benth., <i>Mimosa tenuiflora</i> (Willd.) Poir.
<i>H. suaveolens</i> (S16)	Milagres	June 2008	<i>Hyptis suaveolens</i> Poit.
<i>L. rigida</i> (S17)	Choró	July 2007	<i>Licania rigida</i> Benth.
<i>Z. joazeiro</i> (S18)	Várzea Alegre	July 2007	<i>Ziziphus joazeiro</i> Mart.
Heterofloral (S19)	Cascavel	January 2009	<i>Anadenanthera macrocarpa</i> Benth., <i>Anacardium occidentale</i> L., <i>Cocos nucifera</i> L., <i>Byrsonima intermedia</i> A. Juss.
Heterofloral (S20)	Aracati	October 2008	<i>Cocos nucifera</i> L., <i>Moringa oleifera</i> Lam., <i>Byrsonima crassifolia</i> L. Kunth.
<i>L. sidoides</i> (S21)	Monsenhor Tabosa	November 2008	<i>Lippia sidoides</i> Cham.
<i>M. urundeuva</i> (S22)	Monsenhor Tabosa	June 2008	<i>Myracrodruon urundeuva</i> Fr. All.
Heterofloral (S23)	Itapipoca	August 2008	<i>Anacardium occidentale</i> L., <i>Cocos nucifera</i> L., <i>Byrsonima crassifolia</i> L. Kunth., <i>Borreria verticillata</i> Mayer

\*All municipalities were located in Ceará State.

## 2.4 Protein content

The Bradford method (1976) was used for protein determination. To a 0.1 mL solution of protein extract (honey sample 50% w/v) were added 5 mL of Coomassie Brilliant Blue (200 mg of Coomassie Brilliant Blue G-250 dissolved in 100 mL 95% ethanol, and then 200 mL 85% H<sub>3</sub>PO<sub>4</sub> were added. The resulting solution was diluted to a final volume of 2 L). The Coomassie Brilliant Blue forms a protein-dye complex. After 2 minutes of incubation, absorbance was measured at

595 nm against an albumin standard solution of bovine serum (10-100 µg/0.1 mL) in 0.15 M NaCl.

## 2.5 Mineral analysis

The ashes were dissolved in an acid medium (0.1M HCl), and the volume was obtained in a 10 mL volumetric balloon. From the ash solutions, Cr and Se were quantified by Graphite Furnace Atomic Absorption Spectrometry (SpectrAA220 ZL model, equipped with a longitudinal Zeeman-effect background

correction system, Varian, Mulgrave, Australia), and the other mineral constituents (Na, K, Ca, Mg, Fe, Cu, Zn, and Mn) were determined by Flame Atomic Absorption Spectrometry (SpectrAA55 model, Varian, Mulgrave, Australia). In both cases, the analyses were carried out following the manufacturer's recommendations. The standard solutions of minerals were prepared from several dilutions of stock standard solutions (1000 mg/L) supplied by J. T. Baker (England).

### 3 Results and discussion

#### 3.1 Floral origins of honey samples from Ceará State

The description of the honey samples, their places of collection, period of collection, and vegetation characteristics where the samples were collected are shown in Table 1.

#### 3.2 Physicochemical properties

The physicochemical parameters of the honey samples (moisture, pH, ash, free acidity, color, and insoluble solids) are shown in Table 2. Most of the moisture values were below 20%, which is the maximum value allowed by CAC and Brazilian

regulations for honey (BRASIL, 2000). *L. sidoides* (S18) (20.80%), a heterofloral sample (S10) (20.40%), and *L. rigida* (S17) (19.78%) had the highest moisture values, while *Serjania sp* (S20) had the lowest value (13.63%). Moisture content in honey is associated to its botanical source (ABU-TARBOUSH.; AL-KAHTANI; EL-SARRAGE, 1993) and to several other factors such as the degree of honey ripeness, processing techniques, stockpiling conditions, relative air moisture, blossom, and amount of available nectar. There is a great variation in climate and soil all over Ceará State. Therefore, there are some places with high and others with low relative air moisture. Moisture is the key criterion that determines the ability of honey to remain fresh and free of fermentation (BOGDANOV et al., 1999). Bendini and Souza (2008) found values of moisture varying from 16.50 to 19.20% in honey samples originating from cashew flowers from the city of Cascavel, Ceará State. These results are close to those obtained in the present study. Arruda et al. (2005) found moisture contents ranging from 14.97 to 17.23% in honey samples from the region of Chapada do Araripe/Santana do Cariri, Ceará State. Some of these results are lower than the results obtained in the present study, which includes honey

**Table 2.** Physicochemical parameters of honey bee of different plant origins from Ceará State - Brazil.

Sample	Moisture	Ash	Free acidity	pH	Color	Ins. solids	Protein
	(%, w/w)	(%, w/w)	(meq/kg)		(mUA)	(%)	( $\mu\text{g g}^{-1}$ )
S01	14.39 ± 0.89	0.28 ± 0.02	34.00 ± 2.45	3.66 ± 0.01	573 ± 19	0.07 ± 0.01	708 ± 61.20
S02	18.39 ± 1.41	0.33 ± 0.06	44.40 ± 1.60	3.73 ± 0.17	1185 ± 265	0.06 ± 0.03	476 ± 26.13
S03	16.55 ± 0.39	0.17 ± 0.04	35.00 ± 1.45	3.43 ± 0.05	1257 ± 121	0.03 ± 0.01	255 ± 32.40
S04	15.62 ± 0.31	0.01 ± 0.02	18.20 ± 0.53	3.15 ± 0.02	2600 ± 135	0.08 ± 0.02	1121 ± 14.50
S05	15.89 ± 0.45	0.67 ± 0.02	40.70 ± 2.23	3.83 ± 0.33	2033 ± 175	0.10 ± 0.01	288 ± 40.62
S06	14.97 ± 1.01	0.15 ± 0.15	28.80 ± 2.01	3.45 ± 0.16	801 ± 42	0.03 ± 0.02	121 ± 15.09
S07	19.00 ± 0.87	0.01 ± 0.21	17.00 ± 1.19	3.50 ± 0.28	1432 ± 11	0.14 ± 0.01	330 ± 21.30
S08	18.50 ± 0.81	0.12 ± 0.01	25.00 ± 0.14	3.18 ± 0.07	153 ± 19	0.09 ± 0.01	248 ± 48.31
S09	16.95 ± 0.22	0.07 ± 0.03	19.10 ± 1.02	3.68 ± 0.26	306 ± 58	0.04 ± 0.03	301 ± 31.05
S10	20.40 ± 0.56	0.71 ± 0.01	54.50 ± 1.65	3.38 ± 0.42	327 ± 29	0.17 ± 0.01	198 ± 15.01
S11	18.00 ± 1.12	0.01 ± 0.04	16.67 ± 2.15	3.40 ± 0.31	355 ± 53	0.14 ± 0.00	289 ± 31.11
S12	17.40 ± 0.84	0.07 ± 1.11	37.51 ± 0.98	3.61 ± 0.25	228 ± 41	0.09 ± 0.01	382 ± 51.07
S13	18.20 ± 0.78	0.06 ± 0.02	38.96 ± 1.76	3.65 ± 0.20	212 ± 19	0.13 ± 0.01	356 ± 27.02
S14	16.83 ± 1.08	0.01 ± 0.01	25.14 ± 3.43	3.21 ± 0.05	178 ± 45	0.10 ± 0.02	295 ± 37.21
S15	15.65 ± 0.15	0.52 ± 0.21	49.25 ± 0.75	4.21 ± 0.41	2945 ± 141	0.03 ± 0.05	451 ± 21.54
S16	14.79 ± 0.25	0.05 ± 0.14	33.54 ± 2.72	3.01 ± 0.69	610 ± 55	0.07 ± 0.02	312 ± 44.08
S17	19.78 ± 0.92	0.24 ± 0.03	52.00 ± 2.51	4.17 ± 0.15	638 ± 102	0.18 ± 0.01	397 ± 35.14
S18	20.80 ± 0.65	0.13 ± 0.05	51.03 ± 1.00	3.70 ± 0.10	902 ± 33	0.13 ± 0.00	471 ± 44.56
S19	18.23 ± 2.58	0.30 ± 0.21	50.25 ± 0.75	3.80 ± 0.13	2062 ± 441	0.05 ± 0.04	845 ± 38.21
S20	13.63 ± 0.34	0.01 ± 0.01	31.50 ± 0.50	3.67 ± 0.25	412 ± 44	0.06 ± 0.01	178 ± 24.08
S21	17.85 ± 0.43	0.32 ± 0.17	20.42 ± 0.77	3.43 ± 0.40	876 ± 53	0.17 ± 0.00	355 ± 30.17
S22	17.23 ± 0.12	0.12 ± 0.08	40.10 ± 0.02	3.78 ± 0.17	2830 ± 258	0.10 ± 0.02	724 ± 48.16

Honey floral origin: *Hyptis suaveolens* (S02), *Anacardium occidentale* (S04), *Spermacoce verticillata* (S05), *Mimosa verrucosa* (S08), *Piptadenia moniliformis* (S09), *Myracrodruon urundeuva* (S15 and S19), *Licania rigida* (S17), *Lippia sidoides* (S18), *Serjania sp* (S20), *Ziziphus joazeiro* (S22).

samples from different regions of Ceará State and not only from the Cariri region, which has a typical vegetation.

The honey from a heterofloral sample (S10) had greatest free acidity value (54.50 mEq/kg). *L. rigida* (S17) (52.00 mEq/kg) and *L. sidoides* (S18) (51.03 mEq/kg) also had acidity levels above the maximum value. AOAC and Brazilian regulations for honey (BRASIL, 2000) allow 50.00 mEq/kg as the maximum free acidity value. A high level of acidity in honey may indicate that fermentation has occurred. The honey samples with free acidity values above 50.00 mEq/kg, except for the honey sample from *L. rigida*, had moisture contents a little higher than 50.00 mEq/kg. Sodré, Marchini and Carvalho (2002) found free acidity values between 13.00 and 43.00 mEq/kg in honey samples from Northeastern Brazil. Abadio Finco, Moura and Silva (2010), studying honey samples from Southern Tocantins State, found free acidity values between 35.00 and 59.00 mEq/kg. Acidity is influenced by chemical properties of the organic and inorganic acids, e.g. the tridimensional molecular structure and the ionization rate in the honey samples and also by the amino acids content provided by the nectar and salivary enzymes of *A. mellifera* (CRANE, 1983). According to Pamplona (1989), the content of gluconic acid, produced from glucose by the action of the glucose-oxidase enzyme, tends to increase during the storage of honey because this enzyme remains in activity after processing. Thus, honey acidity increases during the storage and as a result the pH decreases.

The pH analysis of the honey samples from Ceará were performed although the Brazilian legislation (BRASIL, 2000) does not require evaluation of quality of honey samples. The honey samples showed pH values between 3.01 and 4.21. Values of pH represent a measure of the acidity of dissolved hydrogen ions in water. Sodré, Marchini and Carvalho (2002) found similar pH values ranging from 3.37 to 4.46 in honey samples from the North Coast of Bahia State. Borsato et al. (2010), studying honey samples from Paraná State, found values of pH varying from 3.60 to 5.35. Increasing trends in pH have been detected in samples with higher values of ashes, which are in accordance with the results found in the present study (Table 2) (BORSATO et al., 2010).

The honey samples analyzed had insoluble solid contents ranging from 0.03 to 0.18%. Honey sample from flowers of *L. rigida* (S17) (0.18) showed the greatest insoluble solid content. Some other honey samples analyzed, such as S07, S10, S11, S13, S17, S18, and S21, also had values higher than the maximum limit of insoluble solids allowed by Brazilian legislation (0.10%) (BRASIL, 2000). The measurement of insoluble matter is an important means to detect honey impurities that are higher than the permitted maximum value. It was used when a significant portion of the honey collected worldwide was harvested by pressing the combs. However, nowadays almost all commercial honey is harvested by centrifugation. Wax, which is not determined by the *Codex* method, is a major source of water-insoluble contamination (BOGDANOV et al., 1999). Silva, Queiroz and Figueirêdo, studying honey samples from Piauí State, found 0.08% as the mean value of insoluble solids in water with a range of variation from 0.06 to 0.09%.

The results of color intensity showed that the absorbance of a 50% (w/v) honey solution varied from 153 (*Mimosa verrucosa*, S08) to 2945 mUA (*Myracrodruon urundeuva*, S15). The predominance of honeys colors varying from amber to dark amber was observed. Color is the physical property immediately perceived by the consumer. Indeed, the pricing of honey depends to a great extent on honey color (BOGDANOV; RUOFF; PERSANO ODDO, 2004). Beretta et al. (2005) investigated 14 commercial honey samples of different floral and geographic origins and found values of absorbance varying from 25 mAU, for the pale-white honey sample (acacia), to 3413 mAU, for the dark-brown strawberry tree honey sample. This marked difference might be a reliable evidence of the presence of pigments such as carotenoids and Maillard reaction products (GHELDOLF; WANG; ENGESETH; 2002). According to Beretta et al. (2005), this could be due to specific contaminating pigments arising from handling, processing, and storage, and/or from biochemical reactions during honey maturation.

The ash values of the honey samples in this study varied from 0.01 to 0.71%, with a average value of 0.19%. The highest content of ashes was found in the heterofloral honey sample (S10) (0.71%), followed for *Spermacoce verticillata* (S05) (0.67%), and *M. urundeuva* (S15) (0.52%) honey samples. These samples had dark colors which, according White Junior (1978), reflect their higher mineral content; however, other honey samples, such as *Anacardium occidentale* (S04) (0.01%) and *Ziziphus joazeiro* (S22) (0.12%), with low ash contents had the two highest color values (2600 and 2830 mUA, respectively). The maximum ash content allowed by the European Honey Commission (BOGDANOV; MARTIN; LÜLLMANN, 1997) for floral honey is 0.60%. Two samples, among the honey samples analyzed in the present study, exceeded the maximum value permitted by law. Analyzing honey samples from Piauí State, Northeastern Brazil, Silva, Queiroz and Figueirêdo (2004) found average ash values ranging from 0.06 to 0.14%.

### 3.3 Protein contents

The quantitative determination of protein contents are shown in Table 2. The protein content found in *A. occidentale* (S04) honey sample was the highest (1121.00 µg/g), followed by the honey samples of *M. urundeuva* (S19) (845.80 µg/g) and *Z. joazeiro* (S22) (724.50 µg/g). In general, the results are within the range established by the official methods used in Brazil (BRASIL, 2000). Azeredo et al. (2003) studied the presence of proteins in honey samples of different floral origins, commercialized in several states in Brazil, using the Bradford method, and they found that the colorimetric determination of the protein content of honey samples using this method was efficient. To Azeredo et al. (2003) considered high protein contents to be those higher than 1000.00 µg/g and found high values of protein content in honey samples of *Borreria verticillata* (2236.00 µg/g), from Piauí.

### 3.4 Mineral analysis

The macromineral content of the honey samples analyzed are shown in Table 3. The most abundant mineral in the floral honey samples was K, with levels ranging from 21.30 mg/kg,

for *P. moniliformis* (S09), to 1513.30 mg/kg, for the heterofloral sample (S10). The mineral with the second highest content in the honey samples studied was Ca, with values ranging from 14.58 mg/kg, for a sample of honey of *H. suaveolens* (S02), to 304.82 mg/kg, for the honey sample of *Z. joazeiro* Mart (S22); and the third most abundant mineral was Na, with levels varying from 1.80 mg/kg, for *Z. joazeiro* (S22) honey sample, to 47.20 mg/kg, for *M. urundeuva* (S15) honey sample, followed by *M. urundeuva* (S19) (38.40 mg/kg) from a different place of collection. The levels of Zn in the honey samples analyzed varied from 0.07 mg/kg, for *P. moniliformis* honey sample (S09), to 1.85 mg/kg, for the heterofloral honey sample (S21). They are shown in Table 4. Several authors found that the levels of minerals in honey vary according to the botanical origin and soil composition (POHL, 2009). Calcium is a macronutrient essential for plants as well as for animals. It occupies much of the exchange sites in neutral and calcareous soils. Zinc is an important micronutrient essential for plant growth and development. Iron is an essential micronutrient for both plants and animals. In neutral-to-alkaline soils, it is present as insoluble Fe<sup>3+</sup> compounds. Manganese is a heavy metal and an essential micronutrient (SARWAR et al., 2010). However, there is little information about the mineral content of honey from Ceará State.

**Table 3.** Macrominerals in honey samples of different origin from Ceará State.

Samples	Na (mg/kg)	K (mg/kg)	Ca (mg/kg)	Mg (mg/kg)
S01	12.40	44.50	21.24	2.48
S02	6.60	302.90	14.58	7.71
S03	12.70	328.40	64.01	4.19
S04	14.00	53.20	44.78	22.73
S05	5.00	1057.40	59.93	13.32
S06	23.10	43.00	37.55	10.83
S07	2.10	103.40	30.59	11.74
S08	15.00	198.00	56.63	15.58
S09	11.20	21.30	37.98	5.45
S10	13.10	1513.30	60.29	25.16
S11	14.70	45.30	16.67	7.85
S12	4.30	126.70	57.51	2.79
S13	15.40	169.80	38.96	10.24
S14	4.20	62.50	58.14	14.76
S15	47.20	1234.60	69.29	28.33
S16	18.00	236.00	87.54	13.16
S17	35.20	559.10	103.45	12.95
S18	22.60	269.90	15.99	9.91
S19	38.40	137.80	141.30	24.76
S20	4.90	48.10	15.55	9.74
S21	10.70	315.50	79.99	22.02
S22	1.80	251.70	304.82	25.47
<b>MV</b>	<b>15.06 ± 11.72</b>	<b>310.30 ± 407.36</b>	<b>62.00 ± 61.82</b>	<b>13.53 ± 7.70</b>

Honey floral origin: *Hyptis suaveolens* (S02), *Anacardium occidentale* (S04), *Spermacoce verticillata* (S05), *Mimosa verrucosa* (S08), *Piptadenia moniliformis* (S09), *Myracrodruon urundeuva* (S15 and S19), *Licania rigida* (S17), *Lippia sidoides* (S18), *Serjania sp* (S20), *Ziziphus joazeiro* (S22).

Minerals are a big class of micronutrients, which are largely considered essential nutrients. They are traditionally divided into macrominerals (volume elements) and microminerals (trace elements) (MAHAN; ESCOTT-STUMP, 2005). Minerals represent near 4.00 to 5.00 % of body weight or 2.80 to 3.50 kg in adults, men and women respectively. Approximately 50% of this weight is calcium. The best way to obtain a variety of minerals is through various foods instead of the intake of dietary supplements (MAHAN; ESCOTT-STUMP, 2005).

Adequate Intake (AI) for Ca, for men and women is 1000.00 mg for adults of 19 to 50 years old and 1200.00 mg for adults above 51 years old; for Mn is 2.30 mg for men and 1.80 mg for women; for Cr 35.00 µg for men and 25.00 µg for women. Recommended Dietary Allowance (RDA) for minerals is: from 1600.00 to 2000.00 mg (K); 500.00 mg (Na); from 400.00 to 420.00 mg (Mg) for men and from 310.00 to 320.00 mg for women (14 to 70 years old); 10.00 mg (Fe) for men and 15.00 mg for women; 11 mg (Zn) for men and 8.00 mg for women; 900.00 µg (Cu) for men and women; and 55.00 µg (Se) for men and women (FRANCO, 2008; MAHAN; ESCOTT-STUMP, 2005). If an adult individual ingests a tablespoon of honey every day, according to the findings obtained in this study, this person could have an intake of 0.65 mg of Na, 13.33 mg of K, 2.67 mg of Ca, 0.58 mg of Mg, 0.68 mg of Fe, 0.02 mg of Cu, 0.03 mg of Mn, 0.02 mg of Zn, 3.32 µg of Cr, and 0.19 µg of Se. A tablespoon of honey from Ceará State is sufficient to meet the daily potassium and calcium requirements, as recommended by RDA.

The most abundant metal in honey is K (FERNÁNDEZ-TORRES et al., 2005). Other major metals present in honey are Na (second most common), Ca, and Mg. Cu, Fe, Zn, and Mn are present in intermediate quantities (POHL, 2009). Almeida-Anacleto and Marchini (2004) studied honey samples from the city of Pirassununga, State of São Paulo, and found K to be the most abundant element. These results are consistent with those found by Abu-Tarboush, Al-Kahtani and El-Sarrage (1993). In the present study, K was the most abundant metal found in the honey samples from Ceará.

Almeida-Anacleto and Marchini (2004) found Ca to be the second most abundant element in honey samples from São Paulo, southeastern Brazil. Conti (2000) investigated honey samples of different botanical origins in the Lazio region (Central Italy) and found the following average values (µg/g weight): K (472.00), Na (96.00), Ca (47.70), Mg (37.00), Cu (0.31), Fe (4.51), Mn (3.00), and Zn (3.14). The level of K was lower than those reported for the honey samples from Galicia by Rodriguez-Otero et al. (1994), who found 1572.00 µg/g.

Almeida-Anacleto and Marchini (2004) found Mg as the third most abundant mineral in their honey samples. Fernández-Torres et al. (2005) determined the levels of eleven elements (Zn, P, B, Mn, Mg, Cu, Ca, Ba, Sr, and K) in 40 honey samples from four different plant sources from different locations in Spain. These authors found that K, Ca, and P showed the highest concentrations: 434.10-1935.00 mg/kg for K, 42.59-341.00 mg/kg for Ca, and 51.17-154.30 mg/kg for P. The levels of Cu (0.53-2.12 mg/kg), Ba (0.11-1.26 mg/kg), and Sr (0.26-1.46 mg/kg) were the lowest in all honey samples. Concentrations of Zn (1.33-7.29 mg/kg), Mn (1.33-9.47 mg/kg), Mg (13.26-74.38 mg/kg)

**Table 4.** Concentration of trace elements in honey samples of different floral origins from Ceará State.

Samples	Cu	Mn	Zn	Cr	Se	Fe
	(mg/kg)	(mg/kg)	(mg/kg)	(µg/kg)	(µg/kg)	(mg/kg)
S01	0.07	0.26	0.22	76.28	0.84	0.73
S02	0.44	0.61	1.14	22.50	1.68	1.10
S03	0.57	1.42	0.71	54.65	5.42	2.57
S04	0.52	0.55	0.92	73.20	1.84	2.44
S05	0.10	0.57	0.32	46.45	4.02	1.52
S06	0.38	0.50	0.61	170.33	1.44	1.01
S07	0.36	0.43	0.26	84.12	0.84	0.32
S08	0.72	0.97	0.57	110.55	2.20	0.89
S09	0.16	0.06	0.07	140.21	1.96	0.31
S10	0.53	1.96	1.17	84.43	15.82	3.07
S11	0.33	0.18	0.40	74.36	15.40	0.38
S12	0.16	0.71	0.21	52.15	17.40	2.28
S13	0.23	0.41	0.41	30.19	11.20	0.12
S14	0.31	0.47	0.61	32.20	0.36	0.17
S15	1.29	0.18	0.30	144.15	2.48	4.70
S16	0.55	0.68	0.14	100.60	2.02	0.61
S17	0.50	1.83	0.61	98.13	0.88	1.80
S18	0.34	0.73	0.27	96.00	0.56	0.45
S19	0.76	1.65	0.29	116.72	1.90	1.39
S20	0.43	0.11	0.11	34.50	1.28	0.32
S21	0.56	1.30	1.85	43.67	0.76	8.76
S22	0.57	1.62	1.24	44.06	62.00	0.62
<b>MV</b>	<b>0.43 ± 0.27</b>	<b>0.80 ± 0.58</b>	<b>0.56 ± 0.44</b>	<b>77.17 ± 40.25</b>	<b>4.31 ± 5.28</b>	<b>1.58 ± 1.93</b>

Honey floral origin: *Hyptis suaveolens* (S02), *Anacardium occidentale* (S04), *Spermacoce verticillata* (S05), *Mimosa verrucosa* (S08), *Piptadenia moniliformis* (S09), *Myracrodruon urundeuva* (S15 and S19), *Licania rigida* (S17), *Lippia sidoides* (S18), *Serjania sp* (S20), *Ziziphus joazeiro* (S22).

kg), and Na (11.69-218.50 mg/kg) showed a strong dependence on botanical origin. Franchini et al. (2007), analyzing 14 Brazilian commercial honey samples, determined trace metals by capillary zone electrophoresis. The average contents of 656.00, 69.10, 71.80, 36.00, 21.40, and 1.70 mg/kg were found, respectively for K<sup>+</sup>, Ca<sup>2+</sup>, Na<sup>+</sup>, Mg<sup>2+</sup>, Mn<sup>2+</sup>, and Li<sup>+</sup> in the honey samples analyzed.

The honey samples from Ceará had lower values than those reported by Fernández-Torres et al. (2005). Przybyłowski and Wilczyńska (2001), studying the mineral levels in honey samples from the Pomerania region, found an average value for Zn of 7.76 mg/kg, which exceeded the limits accepted by legislation in that region. Nanda et al. (2003) studied the presence of mineral content in six floral honey samples from different botanical sources in Northern India and found considerable variations in the concentrations of Na, K, Ca, Fe, Cu, and Zn.

The levels of K, Ca, Cu, Fe, Mg, Mn, and Zn have been evaluated in several Brazilian plant leaves (LOPES et al., 2002; SILVEIRA et al., 2009). Among these minerals, K had the highest content, followed by Mg and Ca. Lopes et al. (2002) analyzed the mineral composition of several medicinal plants grown in Ceará State. The main minerals found in these plants in order of abundance were Ca > K > Mg > Na; although the order differs from that found in the present study for the honey samples from Ceará (K > Ca > Na > Mg), the main minerals are the same.

## 4 Conclusions

The average values of the physicochemical parameters and minerals found in the honey samples studied, with a few exceptions, were similar to those reported in the literature for good quality honeys. Among the minerals detected in the honey samples in the present study, K showed the highest concentration, followed by Ca, Na, and Mg, which is in agreement with the main minerals found in plants in general. Comparing mineral contents, *M. urundeuva* honey sample had the highest content among the four macrominerals studied (Na, K, Ca, and Mg). The protein content of *A. occidentale* honey was the highest, followed by the honey samples from *M. urundeuva* and *Z. joazeiro*. One tablespoon spoon of honey from Ceará state a day is enough to obtain the amounts of macrominerals needed each day, mainly calcium and potassium. In a great number of studies, the assessment of the levels of different metals, especially heavy metals, has been a general objective aimed at characterizing different types of honey in view of safety implications. The analyses of the trace elements of honey samples from Ceará State showed that their production areas are free of the environmental pollution.

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