



# Physicochemical characterization of *Lavandula* spp. honey with FT-Raman spectroscopy



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## ARTICLE INFO

### Keywords:

FT-Raman

Honey

*Lavandula* spp.

Chemical composition

Quality control

PLS regression models

## ABSTRACT

This study aimed to evaluate the potential of FT-Raman spectroscopy in the prediction of the chemical composition of *Lavandula* spp. monofloral honey. Partial Least Squares (PLS) regression models were performed for the quantitative estimation and the results were correlated with those obtained using reference methods.

Good calibration models were obtained for electrical conductivity, ash, total acidity, pH, reducing sugars, hydroxymethylfurfural (HMF), proline, diastase index, apparent sucrose, total flavonoids content and total phenol content. On the other hand, the model was less accurate for pH determination. The calibration models had high  $r^2$  (ranging between 92.8% and 99.9%), high residual prediction deviation - RPD (ranging between 4.2 and 26.8) and low root mean square errors.

These results confirm the hypothesis that FT-Raman is a useful technique for the quality control and chemical properties' evaluation of *Lavandula* spp honey. Its application may allow improving the efficiency, speed and cost of the current laboratory analysis.

## 1. Introduction

Honey is a natural food product produced by honey-bees that possesses a high amount of available sugars [1] and is a rich source of amino acids, vitamins, minerals and other biologically active compounds [2]. Honey carbohydrates are composed up of about 70% monosaccharides (mainly glucose and fructose), 10–15% disaccharides and a minor concentration of trisaccharides [1]. The chemical composition of this beehive product depends on the botanical and geographical origin [3], which may be evaluated through several methodologies.

Usually, a sample is classified as *Lavandula* spp. monofloral honey (common name: Lavender honey) when the percentage of pollen grains from *Lavandula* spp. is higher than 15% [4,5]. Even so, this monofloral honey may present a large variation in pollen spectrum resulting from the large variability in the ecosystems surrounding the apiaries. *Lavandula* spp. honey is characterized by a pleasant floral aroma, sweet taste and a light amber colour. Its chemical and sensory characteristics make it a much-appreciated honey with a high commercial value in Portugal and in the international market, which make it essential to

ensure an efficient and specific quality control for this product.

Vibrational molecular spectroscopy techniques are very useful for food and beverages' analysis due to their flexibility, efficiency and low cost [6]. Particularly, the use of FT-Raman is advantageous due to the small volume of sample required, the high data reproducibility and speed of analysis. Furthermore, in comparison to other spectroscopy techniques like FTIR or NIR, Raman has the advantage that spectral information avoids the interference related to the water molecule.

Spectroscopic techniques like FTIR, NIR or Raman spectroscopy have been used in the identification as well as quantification of the chemical composition of different products in food, pharmaceutical and other industries. Particularly, FT-Raman methodology is based on the scattering of light from near infrared radiation due to the vibrational energy of the molecules in the sample. FT-Raman has been used in food analysis, namely: quantitative analysis of vitamin A [7]; sugars in honey [8]; determination of erucic acid content in canola oil [9]; detection of vitamins B<sub>2</sub> and B<sub>12</sub> in cereals [10]; classification of different vegetable oils and identifying adulteration on virgin olive [11]; assessment of the quality of Southern Italian honey Types [12]; qualitative analysis of food fraud [10]; controlling protected designation of

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origin of wine [13].

Regarding honey analysis, Kizil et al. [14] evaluated the chemical changes induced by gamma irradiation on the fructose content of honey. Batsoulis et al. [15] applied and modified a standard HPLC-based method and used FT-Raman spectroscopy to evaluate fructose and glucose percentage. Also, Corvucci et al. [16] demonstrated that Raman spectral data, in combination with PCA models, could be a good tool to identify the botanical origins. In addition, more recently, Tahir et al. [17] applied FT-Raman for the prediction of phenolic compounds (catechin, syringic, vanillic, and chlorogenic acids) measured by HPLC–DAD, antioxidant activity and ferrous chelating capacity measured by Spectrophotometry in honey.

As such, this study aimed to assess the potential of FT-Raman spectroscopy to be an accurate tool for the fast analysis of monofloral *lavandula* honey's quality.

## 2. Material and methods

### 2.1. Samples

One hundred ( $n = 100$ ) *Apis mellifera*'s honey samples were harvested by beekeepers from apiaries located in different regions of Portugal: Alentejo ( $n = 8$ ), Alentejo ( $n = 10$ ), Bragança ( $n = 12$ ), Castelo Branco ( $n = 7$ ), Chaves ( $n = 10$ ), Lisboa ( $n = 6$ ), Lousã ( $n = 6$ ), Marialva ( $n = 4$ ), Mirandela ( $n = 15$ ), Torres Vedras ( $n = 3$ ), Vila Flor ( $n = 12$ ) and Vimioso ( $n = 7$ ). Samples were delivered at the laboratory and kept in the dark at 5 °C until further analysis, which occurred in no more than one month after the extraction from the hives; none of the samples had signs of fermentation or spoilage.

In order to ensure that all samples could be classified as *Lavandula* spp. monofloral honey, palynological analysis was performed. Those samples that did not meet the requirements to be considered as monofloral *Lavandula* spp. honey were rejected.

The qualitative pollen analysis was performed using the acetolysis method, as recommended by Louveaux et al. [18] and Von der Ohe et al. [5]. The examination of the pollen slides was carried out with a Leitz Diaplan microscope (Leitz Messtechnik GmbH, Wetzlar, Germany) at 400 × and 1000 ×. A minimum of 1000 pollen grains were counted per sample. The recognition of the pollen grains was performed using the reference collection of the School of Agriculture of the Polytechnic Institute of Bragança as well as different pollen morphology guides and palynology atlas. The following terms were used for pollen frequency classes: predominant pollen (P, more than 45% of pollen grains counted), secondary pollen (S, 16–45%), important minor pollen (IM, 3–15%) and minor pollen (M, 1–3%).

### 2.2. Physicochemical analysis

The physicochemical parameters of Lavender honey samples assessed in a first phase were: ash content (%); electrical conductivity (mS/cm); 5-hydroxymethylfurfural content (HMF) (mg/kg); free acidity (meq/kg), diastase activity (Schade units/g); reducing sugars (%); apparent sucrose (%); pH and proline (mg/kg). The determinations were carried out in agreement with the Official Methods of Analysis of Association of Official Analytical Chemists (AOAC, 1990) [19], Harmonised methods of the International Honey Commission [20] and the Codex Alimentarius [21].

The protein content (mg/kg) was determined according to the method described by Nogueira et al. [22].

Regarding the total phenolic content of the honey samples, it was estimated following the Folin–Ciocalteu method while the total flavonoid were evaluated using the methodology proposed by Elamine et al. [23]. Three replicate analyses ( $n = 3$ ) were made using each sample. Results are expressed as mean value  $\pm$  standard deviation.

### 2.3. Raman data acquisition and processing

The Raman spectra of the honey samples were acquired using a FT-Raman spectrometer (BRUKER, MultiRAM) equipped with a 180° high-throughput collecting lens, a ultra-high sensitivity Liquid Nitrogen-cooled Ge Diode detector, an integrated 1064 nm ( $9392.5 \text{ cm}^{-1}$ ), diode pumped, Nd:YAG laser with a maximum output power of 500 mW, for a working spectral range of 3500–70  $\text{cm}^{-1}$  Stokes Shift. The instrumental parameters used for spectra acquisition were: spectral resolution: 4  $\text{cm}^{-1}$ ; scanner velocity: 5 kHz; number of sample scans: 100.

The system was operated using the OPUS software provided by the manufacturer. In order to minimize disturbances in the measurement conditions, an automatic motorized XY sample stage was used, accommodating well-plates with 96 sample positions, thus eliminating the need to constantly open the spectrometer for changing samples. Two measurements were performed for each sample. Mean spectra were used in all subsequent calculations.

The spectra were collected at a constant room temperature of 20 °C.

PLS regression was done based on the spectral decomposition using OPUS 7.5.18 BRUKER software according the methodology used in Anjos et al. [24].

The spectral data were regressed against the measured parameters, using the pre-processing methods for PLS-R analysis: multiplicative scatter correction (MSC); minimum-maximum normalization (MinMax); vector normalization (VecNor); straight line subtraction (SLS); constant offset elimination (ConOff); first derivative (1stDer); second derivative (2ndDer); first derivation with multiplicative scattering correction (1stDer + MSC); first derivation with vector normalization (1stDer + VecNor); first derivation with straight line subtraction (1stDer + SLS).

The total number of samples was randomly split into two groups (Set 1 corresponding to 70% of samples; and Set 2 containing the remaining 30% of the samples). This separation into two groups was performed automatically by the software OPUS (v 7.5 Build 7, 5, 18 (20140810), Bruker Optik GmbH, Ettlingen, Alemanha), in order to ensure the representability of the samples.

The first group of 70 samples were used for internal validation (cross-validation) and a second one, the remaining 30 samples, for test (validation set). Wavelength selection was done iteratively by comparing and combining wavenumber ranges, and automatically by defining significant wavenumber ranges with the help of the Martens uncertainty test. In a first step the infrared dataset was regressed against the calibration components, and by means of full cross-validation with one sample omitted a significant number of PLS components was obtained.

The results of the cross-validation were tested for a maximum rank of 10, higher values of coefficient of determination ( $r^2$ ) and ratios of performance to deviation (RPD) and lower root mean square error of cross validation (RMSE) as the test set validation [25].

## 3. Results and discussion

Monofloral status generally refers to the presence of a single pollen type in quantities greater than 45% of the total pollen content in the pollen spectrum analysis. However, for honey samples containing under-represented pollen grains, like *Thymus vulgaris*, *Rosmarinus officinalis*, *Citrus* spp., *Lavandula* spp. and *Arbutus unedo*, the botanical classification must be achieved with a lower pollen frequency percentage - usually ranging between 10% and 20% [2]. The results of honey's pollen profile analysis allow determining its floral origin and confirmed that all samples could be classified as *Lavandula* spp. monofloral honey.

As mentioned before the *Lavandula* spp. monofloral honey needs to have a percentage of pollen grains from *Lavandula* spp. higher than 15% [4,5]. In this study the percentage of *Lavandula* spp. pollen grains of each honey sample ranged from 16% to 83%, evidencing that all samples analysed could be commercialized as *Lavandula* spp.

**Table 1**  
Percentage of other pollen grains found in the monofloral *Lavandula* spp. honey samples.

	Number of honey samples	Min-max	mean $\pm$ $\sigma$	CV
<i>Lavandula</i> spp.	105	15.8–83.1	33.4 $\pm$ 12.0	36.0
<i>Castanea sativa</i>	15	7.5–33.1	18.9 $\pm$ 7.9	41.8
<i>Carduus</i> spp.	3	2.4–11.2	6.8 $\pm$ 4.5	67.1
<i>Cistus</i> spp.	72	3.8–38.5	18.1 $\pm$ 9.2	50.6
<i>Cytisus</i> spp.	35	3.7–21.3	11.4 $\pm$ 4.8	42.2
<i>Echium</i> spp.	65	4.3–41.5	23.5 $\pm$ 11.1	47.1
<i>Erica</i> spp.	23	4.8–52.8	15.6 $\pm$ 11.8	75.9
<i>Eucaliptus</i> spp.	3	3.0–5.0	4.1 $\pm$ 0.9	22.3
<i>Leontodon</i> spp.	21	2.0–21.3	9.4 $\pm$ 5.5	58.3
<i>Prunus</i> spp.	32	2.8–38.0	13.9 $\pm$ 8.6	62.1
<i>Rubus</i> spp.	32	3.0–30.3	12.8 $\pm$ 6.4	50.2
<i>Thymus vulgaris</i>	4	3.0–14.6	7.8 $\pm$ 5.0	63.8
<i>Trifolium</i> spp.	25	2.6–23.1	11.0 $\pm$ 5.5	50.1
<i>Apium</i> spp.	17	2.8–18.9	7.2 $\pm$ 4.2	58.2
<i>Genista</i> spp.	7	6.7–19.5	13.3 $\pm$ 5.1	38.3
<i>Taraxacum officinale</i>	13	3.1–18.1	8.7 $\pm$ 4.2	48.2
<i>Medicago</i> spp.	5	4.7–16.7	10.0 $\pm$ 4.4	44.6
<i>Vicia</i> spp.	4	4.2–11.3	8.1 $\pm$ 2.9	36.1
<i>Quercus</i> spp.	7	3.3–32.1	12.8 $\pm$ 10.5	82.2
<i>Acacia</i> spp.	8	7.7–31.1	15.9 $\pm$ 8.3	52.2
<i>Pinus</i> spp.	1	24.3–25.0	24.6 $\pm$ 0.3	1.1
<i>Chamaespartium sagittale</i>	1	15.8–16.5	16.2 $\pm$ 0.3	1.8
<i>Anthemis</i> spp.	11	2.3–10.4	5.9 $\pm$ 2.6	43.7

monofloral honey. Even so, despite the monofloral classification, other pollen grains were also present in different proportion. The Pollen spectrum and their frequency on the analysed honey samples are presented in Table 1.

According to the legislation the moisture content in honey must not exceed 20%. For all samples, the moisture content was in agreement the legal limits (Table 2). This parameter was measured only to attest the quality of honey, however it was not used in calibration model because water is not a good Raman scatterer. Water does not cause interference in the Raman spectra.

Hydroxymethylfurfural (HMF) and Diastase Activity act as quality indicators, suggesting honey's freshness and/or overheating. Almost all the samples under assessment were within the legal limits established for these variables. However, exceptionally, one of the samples presented 6.7 Gothe degrees. In spite of this, this sample was considered for further assessment since its HMF concentration was in accordance with the standards (Table 2).

Ash and electrical conductivity values depend on the mineral content of the honey, the first one measures the inorganic residue after carbonization and the second evaluates ionisable organic and inorganic substances values. The values obtained (Table 2) for these two parameters are in accordance to the values for nectar honeys according to

**Table 2**  
Results obtained for the different parameters evaluated.

Parameter	Mean $\pm$ $\sigma$	Min – max	Coefficient of variation	Legal limits
Moisture (%)	16.4 $\pm$ 0.8	15.2–19.12	4.9	< 20
HMF (mg/kg)	4.6 $\pm$ 4.4	0.5–17.3	100.0	< 40
Electrical conductivity (mS/cm)	0.27 $\pm$ 0.04	0.20–0.37	15.7	< 0.8
Ash (%)	0.15 $\pm$ 0.07	0.06–0.4	59.8	< 15
Total acidity (meq/kg)	24.2 $\pm$ 7.1	10.5–36.8	29.1	< 50
pH	3.61 $\pm$ 0.59	2.12–6.54	16.2	
Proline (mg/kg)	256.8 $\pm$ 23.6	227.9–304.8	9.2	
Diastase index (Schade units/g)	11.2 $\pm$ 2.8	6.7–17.0	31.1	> 8
Reducing sugars (%)	69.7 $\pm$ 3.9	61.7–77.5	5.6	> 60
Apparent sucrose (%)	3.4 $\pm$ 1.1	1.0–6.8	30.5	< 10
Total phenol content (mg/100 g)	152.0 $\pm$ 41.2	87.9–229.0	27.1	
Total flavonoids content (mg/100 g)	11.6 $\pm$ 2.2	5.8–15.8	18.7	

the Codex Alimentarius [21].

The *Lavandula* spp. honey presented pH values ranging from 2.12 to 6.54 (Table 2), which is in accordance with the values reported for Portuguese honeys and for Lavender honeys [2,4].

The total acidity of honey suggests the absence of unwanted fermentations. The analysed honey samples presented on average 24.2  $\pm$  7.1 meq/kg (Table 2) and are in accordance with the legislation.

Honey is mainly composed of the monosaccharides glucose and fructose and others di- and tri-saccharides [1]. The Reducing Sugars content of the honeys analysed ranged from 61.7% to 77.5%, higher than the 60% required by the legislation.

According to the European Directives, for most honey types the apparent sucrose content must be under 5%. However, for Lavender honey, due to their floral origin, this value must be under 10%. All the samples met this directive.

The Protein content is one of the minor components in honey and came from the honeybee and from the pollen content in honey [22]. The protein content in the analysed honey samples ranged between 0.21 mg/kg and 0.53 mg/kg and no accurate models could be found for this parameter.

Total phenols and flavonoids are key parameters because they are related to the honey bioactive properties. The total phenol content of the analysed honey varied from 87.9 to 229.0 mg/100 g and the total flavonoid content varied from 5.8 to 15.8 mg/kg (Table 2). These values are in accordance with those reported by Gomes et al [4]. and Estevinho et al. [2].

### 3.1. Raman spectra characterization and analysis

FT-Raman spectra of Lavender honey are shown in Fig. 1. Honey samples show a majority of the spectral peaks in the 200–1500  $\text{cm}^{-1}$  region. In the region between 300 and 1500  $\text{cm}^{-1}$ , peaks were observed at 341, 422, 521, 626, 705, 776, 825, 867, 915, 979, 1072, 1124, 1266, 1366, and 1460  $\text{cm}^{-1}$ , matching those observed in literature [8,17,26,27].

The FT-Raman spectra obtained for honey samples were similar to those reported in the literature [17,28]. The peaks identified in the spectra can be assigned as follows:

- From 200  $\text{cm}^{-1}$  to 500  $\text{cm}^{-1}$  was observed skeletal vibrational modes, namely C-C-O and C- C-C, C-O and C-C [29];
- 521 – deformation of C-C-O and C- C-C [17];
- 626 was assigned to ring deformations [17];
- around 705  $\text{cm}^{-1}$  corresponds to the stretching of C-O and C-C-O, O-C-O bending [30];
- the band at 776 was assigned to the C-C stretching and C-H vibrations present in glucose [30,31];
- at 867 and 825  $\text{cm}^{-1}$  were assigned to the vibration of C-H and  $\text{CH}_2$  [31] deformation and C-O-H bending [30];

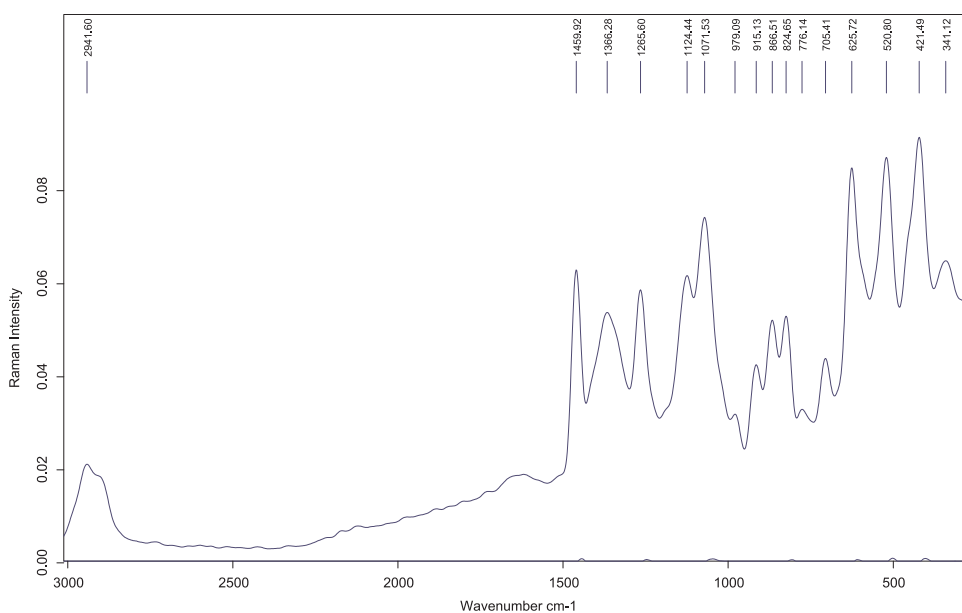


Fig. 1. Average FT-RAMAN spectrum of the *Lavandula* spp. honey.

a signal around  $915\text{ cm}^{-1}$  was associated with the vibration of C-H and C-O-H [32];

- The peak at  $979\text{ cm}^{-1}$  was due to vibration in the two anomers of fructose and glucose [30];
- around  $1072\text{ cm}^{-1}$  were assigned to the carbohydrates bending vibration of C-H and C-O-H and also a minor contribution by the proteins and amino acids vibration of C-N bond in amino acids and proteins [31];
- usually the peak at  $1124\text{ cm}^{-1}$  was attributed to a combination of stretching vibration of C-O and C-O-C and vibration of C-N of protein and amino acids [31];
- around  $1266\text{ cm}^{-1}$  could be attributed to the vibration of C-O-H, C-C-H and O-C-H;
- The signal at  $1366\text{ cm}^{-1}$  corresponded to the bending of C-H and O-H bonds [28];
- At  $1460\text{ cm}^{-1}$  was found the signal associated to a combination of the vibration of COO<sup>-</sup> group of bending vibration of CH<sub>2</sub> group [31,33]. This region was attributed to the presence of flavanols and organic acids [33].

### 3.2. Calibration model development

The PLS regression was used to develop the calibration models after

spectra pre-treatments were applied in order to increase the performance of the predictive models. A different spectral range was selected for each parameter analysed (Table 3).

Table 3 summarizes the best models obtained for each analytical parameter in both cross-validation and test set validation.

Each model was selected according to the criteria explained in material and methods section, namely present a higher coefficient of determination for both sets, with higher residual prediction deviation and lower root mean square error of cross-validation and prediction and bias.

In Fig. 2 was represented the predict model by FT-Raman, for the reference method of some of the chemical characteristics that was reported in the Portuguese legislation for honey quality control.

All analytical parameters evaluated have high correlation coefficient ranging between 92.8% and 99.9% for cross validation and between 93.8 to 99.9% in test set validation, with a rank that varied between 9 and 10. Although for pH the model's parameters are also presented in the Table 3, the authors think the calculated model does not predict values accurately enough to consider it a good model. The values of  $r^2$  for pH were 83.8 (with RPD of 2.5) in Cross-Validation and 98.5 in test set validation (with RPD of 8.2). In this case the analytical pH values ranged only between 2.12 and 6.54, which could be a too narrow interval given the dispersion of the values. In fact, the values

Table 3

Results of the Cross-validation and the validation set for the calculated models regarding the different parameters.

	Spectral range (cm <sup>-1</sup> )	Pre-process	Cross-validation (n = 70)					Validation set (n = 30)			
			Rk	r <sup>2</sup>	RMSECV	RPD	Bias	r <sup>2</sup>	RMSEP	RPD	Bias
Electrical conductivity (mS/cm)	1356–1210 + 1065–772 + 627–480 + 350–250	VecNor	10	92.8	0.0106	3.7	-0.0009	93.8	0.0111	4.2	-0.0030
Ash (%)	1500–772 + 350–250	SLS	10	95.1	0.0107	4.6	0.0008	94.1	0.0008	4.2	-0.0008
Total acidity (meq/kg)	1500–1210 + 1065–919 + 774–336	MSC	10	99.9	0.223	25.9	-0.0211	99.8	0.283	28.4	0.1300
pH	1500–772 + 480–190	MSC	9	83.8	0.119	2.5	-0.0012	98.5	0.0977	8.2	0.0091
Reducing sugars (%)	1500–1354 + 1210–919 + 774–336	MSC	9	99.0	0.343	10.0	0.0142	99.2	0.395	11.6	-0.1140
HMF (mg/kg)	1500–1063 + 336–190	SLS	10	99.9	0.187	26.8	-0.0083	99.0	0.169	10.4	-0.0419
Proline (mg/kg)	1500–1210 + 1065–919 + 774–480 + 336–190	SLS	10	99.2	1.71	11.4	0.1050	99.4	2.03	12.8	-0.1770
Diastase index (Schade units/g)	1500–1063 + 774–627 + 336–190	SLS	10	99.6	0.186	15.8	0.0095	99.3	0.211	12.8	0.0706
Apparent sucrose (%)	1520–80	MSC	9	94.2	0.232	4.2	0.0062	97.3	0.180	6.4	0.0551
Total flavonoids content (mg/100 g)	1500–1063 + 774–627 + 483–190	MSC	10	98.9	0.257	6.9	0.0023	99.2	0.228	11.2	-0.0413
Total phenol content (mg/100 g)	1500–1063 + 919–627 + 336–190	SLS	10	99.9	1.37	26.0	-0.0168	99.9	1.44	31.7	0.0238

VecNor – Vector normalization; SLS - Straight line elimination; MSC - Multiplicative scatter correction; Rk – Rank; r<sup>2</sup> - coefficient of determination; RMSECV - root mean square error of cross-validation; RMSEP - root mean square error of prediction; RPD - residual prediction deviation; Bias - mean value of deviation, also called systematic error.

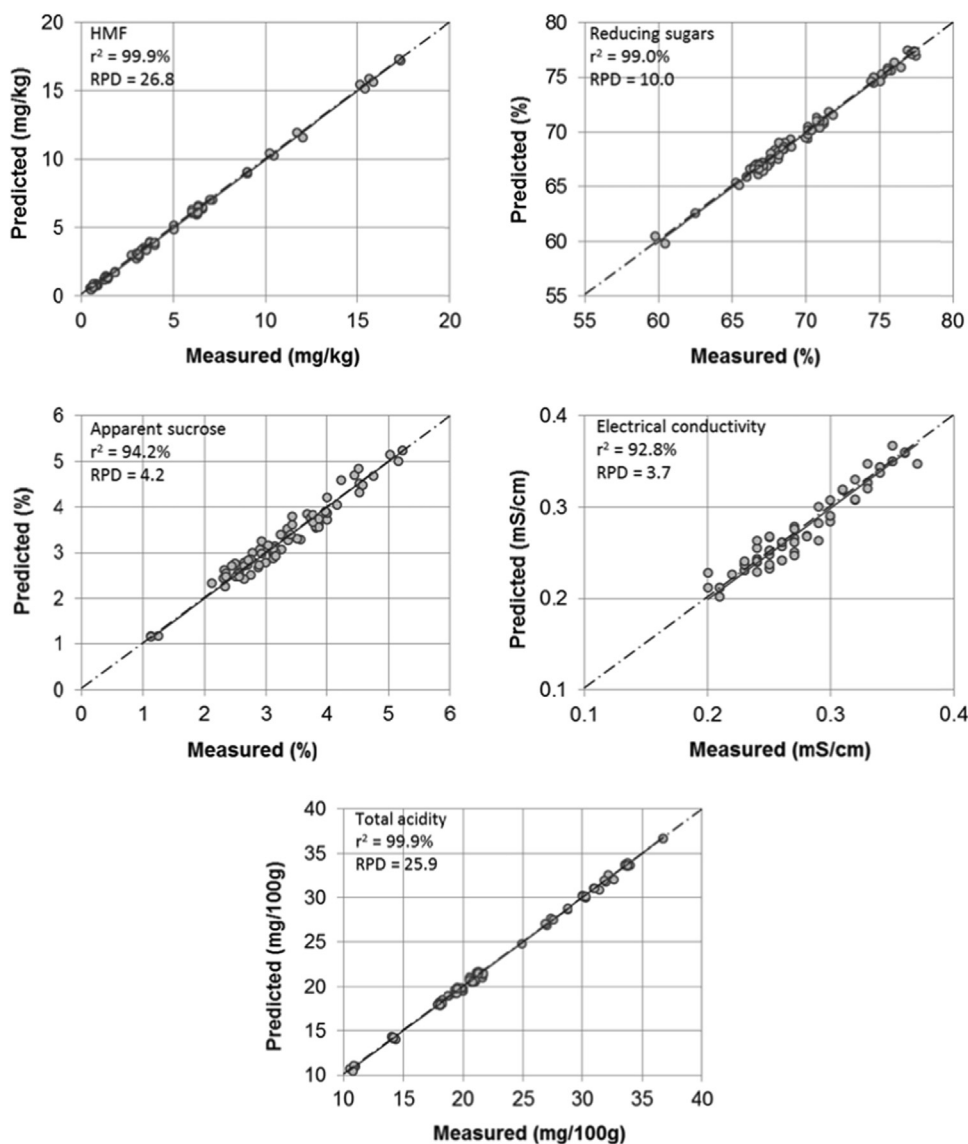


Fig. 2. Cross validation correlation plots of predicted and measured values of HMF, reducing sugars, apparent sucrose, electrical conductivity and total acidity using PLS-R models with FT-Raman spectra.

range between 2.12 and 6.54 but the median is 3.5 and 6.54 is an outlier in this group of data, but a possible value founded in honey. In this case more data with higher variability would be needed in order to try to improve the calibration model, similarly to what other authors found for other spectroscopic techniques [34,35].

The RPD for cross-validation and test set validation was higher than 3.7 and ranged from 3.7 (cross-validation for electrical conductivity) to 31.7 (Test-Set for total phenol content). These results are available in Table 3.

Regarding the parameters considered as “basic standards” for honey’s commercialization (electrical conductivity, ash, total acidity, pH, HMF, reducing sugar and apparent sucrose), this work provides very good calibration models with  $r^2$  values varying from  $r^2 = 92.8\%$  for electrical conductivity to  $r^2 = 99.9\%$  for HMF and Total acidity, all with high RPD values. Concerning the remaining parameters, total phenol content presented the better calibration models, considering cross-validation and test set validation.

Tahir et al. [17] have also found good calibration models for phenolic compounds and antioxidant activity in honey.

In Fig. 2, provides scatter plots of the parameters referenced in the legislation predicted by PLS-R calibration models based on the FT-Raman spectra. All of the plots show significant correlation between the measured and predicted values. The accuracy of each prediction model

is quantified in Table 3.

As previously mentioned HMF content is a very important quality parameter because it is related to the honey freshness and/or overheating. Given the results obtained in this work, the calibration models for this parameter, for *Lavandula* honey, are very good:  $r^2 = 99.9\%$  and RPD = 26.8 for cross-validation and  $r^2 = 99.0\%$  and RPD = 10.4 for Test-Set. This supports the importance of a spectroscopy model in quality control laboratory to quickly point out a low quality honey sample.

Lichtenberg-Kraag et al. [34] also found good calibration models for the analytical parameter of honey (sugars, proline, free acids, invertase, moisture, HMF, pH and electrical conductivity) analysed using Fourier-transform infrared spectroscopy (FT-IR) and partial least-square regression. The coefficient of determination found by this author ranging from 84% to 98%.

Using the Near infrared reflectance spectroscopy (NIRS) technique, Cozzolino and Corbella [35] evaluated the chemical composition of fresh honey samples from different locations across Uruguay. Their models, with PLS-R, for water content, pH, electrical conductivity, colour and HMF present a coefficient of determination in calibration ranging between 67% and 96%, and concluded that NIRS is a useful method to evaluate chemical composition of honey.

With Raman technique, some honey analytical parameters had

already been measured with good accuracy [15,36,37], but the studies with the parameter referenced in the European legislation for honey quality control are scarce.

Our research reconfirms the ability of the FT-Raman technique to measure and evaluate the quality of a specific type of honey available in the market. However, more studies are needed prior to the implementation of this technique for the routine laboratorial assessment, particularly regarding the external calibration. Also, it will be important to analyse what spectroscopic technique (NIR, FTIR or Raman) is more accurate and at a lower cost.

#### 4. Conclusion

The results obtained in this study suggest that the proposed methodology has an acceptable accuracy for being applied in the quality control of *Lavandula* spp. Monofloral honey. Also, it appears to be useful for the assessment of other chemical parameters important to support the high quality of this product, namely the content of total phenols and total flavonoids.

Indeed, from the residual prediction deviation value as well as from the determination coefficient, the models demonstrate to be a powerful tool. This technique may be used on honey analysis, especially for supporting a professional extraction unit where the beekeeper needs to have their honey quality evaluated in an easy and fast way to commercialize his product.

More studies are required in order to validate these results with more samples and applying different methodologies. In addition, it would be interesting to test its applicability for the measurement of other health-promoting compounds available in the honey samples.

#### Acknowledgments

Acknowledgements: CEF is a research unit funded by Fundação para a Ciência e a Tecnologia I.P. (FCT), Portugal (UID/AGR/00239/2013).

Centre of Molecular and Environmental Biology is a research unit funded by FCT, UID/BIA/04050/2013 (POCI-01–0145-FEDER-007569) and by the ERDF through the COMPETE2020 - Programa Operacional Competitividade e Internacionalização (POCI).

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