Abstract

With its adverse effects on the health of consumers and the global economy, food adulteration is considered to be one of the pressing issues of our times. Beef Protein powders (BPP) are particularly among the most fraud-prone products mainly due to their ever-growing consumption. For the purpose of our study, FT-NIR spectroscopy was applied to quantify four commonly used adulterants: Melamine (M), Urea (U), Glycine (G) and Taurine (T) in BPP. Analysis with chemometric tools proved the efficiency of the aforementioned technique as a rapid nondestructive analytical tool for the detection of Beef protein powder adulteration.

Keywords

Beef Protein Powder, FT-NIR spectroscopy, adulteration, Chemometrics
melamine has the highest nitrogen content among the studied adulterants, the latter were added in melamine concentration equivalent to account for any differences.

The total number of mixtures corresponded to 90 samples (15 mixtures × 6 adulteration levels per mixture). Each of these was prepared in triplicates including a control resulting in a total number of samples of 271.

Acquisition of NIR spectra

Prior to scanning, each mixture was poured into a 2 cm depth cuvette. The vials were then shaken for an approximate duration of 10 seconds and taped five times for a better homogenization. The samples were scanned in a random order and three consecutives were taken for each sample. The NIR spectra of pure and adulterated protein powder samples were measured over the range [800 to 2500] nm on a multipurpose analyser Fourier Transform Near Infrared Spectrometer (MPA, Bruker Optics, Ettlingen, Germany) using the OPUS software (v.7.2, Bruker Optics, Ettlingen, Germany). The data was collected by diffuse reflectance using the Integrating Sphere module (sphere macro sample) allowing the spectra of the materials to be collected through the bottom of the vials with a resolution of 8 cm⁻¹. Each spectrum was the average of 32 scans. Background spectrum was acquired before each series of measurements.

Data analysis

The analysis of the generated spectra was performed with R-project software (v. 3.3.2, The R Foundation for Statistical Computing, Vienna, Austria) using aquap2 package. To extract the significant information contained in the spectral data, several pre-treatments were applied: Savitzky-Golay (SG) filter (second-order polynomial with 7, 11 or 21 points), second derivatives and Multiplicative Scatter Correction (MSC). (SG) parameters were optimized and the optimal window length that offered the best signal-to-noise ratio of the original spectra was set to 21 points. To minimize the effect of overlapping peaks and baseline shifts, second derivatives of pure, adulterated beef protein powder spectra as well as pure adulterants were calculated based on Savitzky–Golay derivation using 9 points. Last but not least, Multiplicative Scatter Correction was performed on smoothed spectra in order to reduce any possible baseline variations.

Multivariate data analysis

The data set was examined for its overall distribution using principal component analysis (PCA) and also, to detect outliers before building linear discriminant models (LDA). The latter were based on the type of adulteration and a three-fold cross validation was employed to ensure the inclusion of the samples at least once in the evaluation set resulting in three LDA models [1][4]. Partial Least squares Regression (PLSR) was used to develop chemometric models that could predict the concentration of the studied adulterants in adulterated beef protein powders.

The precision and accuracy of the developed PLSR models were evaluated by the coefficient of determination (R2) and root mean square error (RMSE) of both cross-validation and independent prediction. Latent variables (LV) were selected based on the lowest RMSECV (Root Mean Squared Error of Cross-validation) [5].
Results and discussion

Visual inspection of the NIR spectra

From figures (2.a and 2.b), the spectra of adulterated samples revealed some features similar to that of pure adulterants where distinctive peaks were observed at both of the spectra. When inspecting the typical absorbance of melamine, the first overtone region with N-H stretches was the most distinctive, including the absorbance peaks at 1523 nm, which was visible in mixtures containing melamine (figure 2.a). This confirms other results obtained by Tang, Cai, Shi, & Xu, 2016 [6] when quantifying melamine in different brands of milk powders using Near Infrared Spectroscopy.

The most prominent absorbance peak for taurine was centred at 1590 nm, which was also present in samples containing taurine (Figure 2.a). This is in accordance with studies conducted by Lukacs et al., (2018) regarding the absorbance of Taurine in NIR spectral range. Glycine, on the other hand, presented a broad band in the region 1590 nm-1630nm which contains the N-H stretches (first overtone). In the case of urea, the first overtone region with N-H stretches had the most distinctive peaks at 1475 and 1530 nm. These findings are in agreement with the previous studies of [7]. The obtained results are quite informative regarding the typical peaks of adulterants and their effect on the original spectra of pure beef protein powder. However, for more conclusive results, the data was further evaluated by PCA.

Qualitative Data evaluation of the NIR spectroscopy results with PCA and LDA

Results of PCA analysis showed a pattern of separation of the different tested mixtures where the samples containing individual adulterants (U, G, T, and M) were more distant from those containing a combination of the adulterants. Such pattern was also confirmed when each mixture was colored by adulteration level and the most notable separation was obtained for samples adulterated by UM mixture. The pattern is recognizable in the scores plot shown in Figure.3 where PC1 represent 56.76%, while PC2 40.81 % of the total variance and an increasing pattern of adulteration could be seen based on PC2. There is a clear separation between samples containing 3% (level 6) and 2.5% (level 5) of UM compared to the rest of adulteration levels. However, what is most noteworthy is that NIRS was sensitive enough to separate even the lowest concentrations of 0.5% (level 1) and 1% (level 2) in PCA. This further confirms the efficiency of NIRS for the simultaneous detection of adulteration in protein-rich supplements. This is in agreement with what was reported by [8].

However, the separation of the different adulteration levels by PCA was not as conclusive for
other studied mixtures where some overlapping of the different levels was observed. Further qualitative and quantitative modelling was thus performed to ensure the adequacy of NIRS for the detection of Urea, Melamine, Taurine and Glycine.

Results of LDA classification showed that, at the lowest adulteration level corresponding to 0.5%, the model presented low accuracy and different mixtures were misclassified (Figure 4.a). The most distant points correspond to the samples adulterated with each of the individual adulterants. These are quite well separated as opposed to those containing a combination of adulterants. What is equally noticeable is that samples containing amino acid adulterants such as Glycine (maroon) and Taurine (blue) or a combination like GT are close to each other while those containing adulterants of different nature like urea (light blue), melamine or a combination of these adulterants like UM (yellow) are far (Figure 4.b).

Figure 3. PCA scores plot of UM adulterated beef protein powder on the MPA smoothed (Savitzky-Golay filter; 21 points) and MSC pretreated spectra in the wavelength range 1400-1630 nm

Figure 4. Results of LDA classification at level 1 (0.5%) (a) and level 6 (3%) (b) of Beef Protein Powder
When the adulterated beef protein samples were subjected to LDA at the adulteration level 6 corresponding to 3% adulteration, the different mixtures were perfectly classified based on the root 1 and 2 presenting 89.89% of the between group variance. The validation set (x) and training set (●) of each group of mixtures were perfectly fitted (figure 4.b). The different recognition and prediction performances at different levels are shown in Table 1.

### Table 1. LDA recognition and prediction rates at different adulteration levels in 1400-1630nm range

| Level  | Recognition | Prediction |
|--------|-------------|------------|
| 1      | 93.71%      | 70.87%     |
| 2      | 98.5%       | 87.24%     |
| 3      | 99.63%      | 97.79%     |
| 4      | 100%        | 99.17%     |
| 5      | 100%        | 100%       |
| 6      | 100%        | 100%       |

It was observed that the accuracy of the applied method is dependent on the level of adulteration where a decrease of prediction rate from 100% to 70.87% when the adulteration level decreased is observable.

### PLSR Prediction of beef protein powder adulteration:

The prediction accuracy depended on the tested adulterant and the best performances were obtained for the quantification of urea with coefficient of prediction (R²Pr) of 0.89 and Root mean squared error of prediction RMSEP of 0.39%. The corresponding Y-Fit graph is illustrated in Figure 5 where test points are quite fitted to the training set points and concentrations as low as 1.5% were determined.

In case of melamine adulterated samples, the best predictive model was built using the raw spectra acquired in the region 1400-1630 nm by MPA. It was based on six latent variables and presented a coefficient of determination for the test set of 0.95 and an error of prediction (RMSEP) of 0.1961 % (Figure 6) suggesting the efficiency of NIRS when it comes to determining the concentration of melamine in BPP.

![Figure 5. Y-fit graph of the prediction of urea concentration in adulterated BPP samples](image)

In the case of adulterations with a combination of adulterants, the prediction models were also good with coefficients of prediction ranging from 0.88 to 0.96 and errors of prediction from 0.22% to 0.84%. What is notable is that the efficiency of predicting the content of adulterants in mixtures containing adulterants of similar chemical structure such as Glycine and Taurine (GT) was lower than those containing adulterants of different nature such as Urea and Glycine (UG) or Glycine and Melamine (GM).
These findings further confirm the reported results of LDA where misclassification was more probable to occur when the adulterants were similar in nature which was also reflected in the between classes distances. This also accentuates the importance of melamine and urea in conferring a certain recognizable pattern to adulterated samples when tested by PCA.

Figure 6. Y-fit graph of the prediction of melamine concentration in melamine adulterated samples

Conclusions

The study confirmed the presence of an identifiable pattern among adulterated samples, mainly due to the presence of Melamine and Taurine. The results obtained by Principal Component Analysis (PCA) could separate adulterants with concentrations as low as 0.5%. These findings were confirmed by the qualitative analysis with Linear Discriminant Analysis (LDA) where different groups were identified with performances that depended both on the adulterants present (separately or in combination) and the level of adulteration. The samples with higher levels of adulteration (level 6) corresponding to higher concentration of adulterants (3%) were easily classified compared to those adulterated with relatively lower amounts (0.5%) with LDA prediction rates of 100% and 70.87% respectively. According to adulteration type, beef protein powder samples with similar chemical structure (amino-acidic structure in case of glycine and taurine) tended to belong to groups with minor between group variance leading to some misclassifications. Partial Least Squares Regression (PLSR) had prediction coefficients (R²Pr) ranging from 0.86 to 0.96 and errors of prediction from 0.2% to 1.01%. The results accentuate the efficiency of NIR spectroscopy as a rapid non-destructive analytical tool for the detection of Beef protein powder adulteration where levels corresponding to a concentration of adulterant as low as 0.5% can be successfully identified, classified and quantified.

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