

Detection of Sugar Solution Adulteration of Fresh Orange Juice by Near Infrared Spectroscopy

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Received April 10, 2015; Revised April 20, 2015; Accepted April 20, 2015; Published April 24, 2015

ABSTRACT

Orange juices are a popular refreshment beverage and nutrition source. However, the adulteration of pure juice has now become a common practice. In this study, near infrared spectroscopy was applied as a means to determine whether different orange juice samples had been adulterated with sugar solution or not. For this purpose, 10 samples of 100% fresh juice and 80 samples with different concentrations of sugar solutions were created in a laboratory setting. Each sample was scanned with an FT-NIR spectrometer. A PLS regression model tested by validation set to predict the sugar-added content of juice samples provided R^2 , RMSEP and a bias of 92.06%, 0.0361 g/ml, and -0.0041 g/ml, respectively and for water-added 87.68%, 7.37%, and -1.26%, respectively. These results confirm that an NIR-based protocol could be applied for sugar adulteration in orange juice.

Keywords: Orange juice; Adulteration; Sugar solution; Near infrared spectroscopy; PLS regression.

INTRODUCTION

Due to increasing temperatures and the desire for healthier diets, cool fruit juices have become popular particularly in Bangkok and near-by provinces. Orange juice is rich in phenolic compounds and ascorbic acid [1]. Because of its phenolic content, orange juice has known antioxidant properties [2]. In addition, significant amounts of L-ascorbic acid or vitamin C is contained in oranges, making them an important source of nutrition. Indeed, the nutrition content found of oranges is commonly used as a nutritional index.

Due to the size and value of the commercial fruit juice market, the adulteration of the product has been widely practiced, especially in locations along busy roads and intersections in Thailand. Orange juice vendors are found scattered throughout Bangkok and near-by provinces. It is a widely held belief that adulteration of the orange juice by the addition of sugar solution is a common practice. Therefore the ability to detect for the adulteration of orange juice with sugar solution is clearly needed to protect consumers.

Near infrared spectroscopy (NIR) is a non-destructive technique that can be used to rapidly evaluate chemical constituents of materials. Rodriguez-Saona et al. [3] developed an FT-NIR model to predict the sugar levels in fruit juices (apple and orange from department stores) and compared it with HPLC and standard enzymatic techniques.

The models generated from transmittance spectra gave the best performance with a standard error of prediction (SEP) < 0.10% and an R^2 of 99.9%, with the ability to accurately and precisely predict the sugar levels in juices.

Hong and Wang [4] studied the authentication of fresh cherry tomato juices adulterated with different levels of overripe tomato juices: 0–30%. Their study indicated that the prediction performances by fusion approaches were better than by the sole usage of an e-nose or e-tongue method; yet classification and prediction performances based on different fusion approaches vary. Boggia et al. [5] proposed a strategy based on UV-VIS spectroscopy for the detection of filler juices and water added to pomegranate juices.

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Citation: Nawayon J & Sirisomboon P (2015) Detection of Sugar Adulteration of Fresh Orange Juice by Near Infrared Spectroscopy. International Journal of Bioprocess and Biotechnological Advancements, 1(1):57-62.

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Visible and NIR spectroscopy have also been used to determine the levels of important nutrients in orange juice including citric acid and tartaric acid. The correlation coefficients (*r*) and root mean squares error of prediction (RMSEP) in the best model were 0.944 and 0.596 for citric acid and 0.930 and 0.013 for tartaric acid [6]. For soluble solids content (SSC) and pH, the *r*, standard error of prediction and RMSEP for SSC were 0.98, 0.68, and 0.73 and for pH were 0.96, 0.06, and 0.06, respectively [7]. Determination of glucose, fructose, sucrose, citric and malic acids in orange juices were carried out using dry extract samples and the ability of calibration models was acceptable in comparison with the reference methods [8]. The classification accuracy for prochloraz residue was found to be 100% [9].

In addition, 3D-front-face fluorescence spectroscopy has been used to assess the adulteration of orange juice by grapefruit juice at percentages as low as 1% [10]. Principal Component Analysis (PCA) was applied to a set of physicochemical variables and the addition of sugar were investigated. Detecting adulterations started from approximately 15%. The rapid automated screening technique Curie-point pyrolysis mass spectrometry (PyMS) was used to detect a 10% (w/v) beet sucrose solution adulterated with freshly squeezed orange juice over the range 0-20% (or 0-20 g/l of added sucrose) and provided calibration models which gave excellent predictions for sucrose adulteration levels below 1% [12].

The applicability of rapid analytical methods, such as NIR, for fraud detection in fruit juice, and in particular orange juice is of most interest to researchers and government officials. Therefore, the objective of this research was to assess the application FT-NIR spectroscopy for the determination of the adulteration of orange juice with sugar solution at different concentrations.

MATERIALS AND METHODS

Table 1. Different concentration of orange juice adulterated with sugar solution.

Levels	Juice: Sugar Solution*: Water (by volume)	Added Sugar (g/ml juice)	Added Water (% of juice)
1	90:10:00	0.06	6.00
2	80:20:00	0.12	11.93
3	70:30:00	0.18	17.90
4	60:40:00	0.24	23.87
5	50:50:00	0.30	29.83
6	40:60:00	0.36	35.80
7	50:10:40	0.12	45.97
8	50:20:30	0.24	41.93
9	50:30:20	0.36	37.90
10	40:10:50	0.10	55.97
11	40:20:40	0.20	51.93
12	40:30:30	0.30	47.90
13	30:20:50	0.17	61.93
14	30:30:40	0.26	57.90
15	30:40:30	0.34	53.87
16	30:50:20	0.43	49.83

*Sugar solution concentration was 50% w/w.

Samples and Adulteration of Orange Juice with Sugar Solution

There were 10 samples of 100% fresh juices squeezed from oranges (*Citrus tangerina*) bought from a main agricultural distribution market in Pathumthani province, Thailand. *Citrus tangerina*, referred to as “Kiew Wan” in Thailand, is a popular variety used for making juice for sale. Before squeezing, the fruit was cut in half, with 2-3 halves being squeezed at a time without peeling or seed removal. Eighty 200 ml samples of mixed pure juice, sugar solutions and water with different concentrations by volume were prepared (Table 1). There were 5 samples for each level of adulteration. The sugar solution concentration used was 50% w/w.

Near Infrared Scanning

Each sample was transferred into a glass vial of 22 mm diameter and covered with a stainless steel transreflection plate. This provided a 2 mm optical path length, Samples were then scanned between 12500-3600 cm⁻¹ with a nominal resolution of 8 cm⁻¹, accumulating 32 scans per spectrum using a background of the gold. The scanning was performed at room temperature (25 ± 1 °C) using a Multi-Purpose Analyzer (MPA, FT-NIR spectrometer, Bruker, Bremen, Germany).

Spectrum Pre-treatment and NIR Spectroscopy Model Establishment

The NIR calibration models for the pure orange juice and adulterated samples were developed for determination of the amount of added sugar and added water using partial least squares regression (PLS). After the reference data and spectral data were merged and run in ascending order of the reference data. The calibration and validation set were randomly selected using OPUS, v.7.0.129 software with the

condition that calibration set covered the full measurement range. Therefore, all types of juice, 100% fresh juice and adulterated juice, were in both calibration set and validation set. The calibration set consisted of approximately 70% with the remainder used as a validation set. OPUS, v.7.0.129 was used in both spectrum pre-treatment and model development. The NIR spectra used for model development were pre-treated using the following methods in the software; no pre-treatment, constant offset elimination, straight line subtraction, vector normalization (SNV), min-max normalization, multiplicative scatter correction (MSC), first derivatives, second derivatives, first derivatives+straight line subtraction, first derivatives+SNV and first derivatives+MSC. The combination of the following wavenumber ranges including approximately 9000-7500 cm^{-1} (Region A), 7500-6100 cm^{-1} (Region B), 6100-5450 cm^{-1} (Region C), 5450-4600 cm^{-1} (Region D) and 4600-4250 cm^{-1} (Region E), and pre-treatment method was used for model development. The number of latent variables (PLS factors) in a calibration model was optimized by minimum root mean square error of cross validation (RMSECV). After that the optimal model was tested by validation set and the coefficient of determination (R^2), root mean square error of prediction (RMSEP) and the prediction bias were reported.

RESULTS AND DISCUSSION

Table 2 shows the minimum (Min), maximum (Max), mean, and standard deviation (SD) of sugar-added and water-added in fresh orange juice samples in calibration set and validation set.

The scatter plot in Figure 1 shows the prediction data and the reference data for the sugar adulterated samples. The best model was developed using the vector normalization (SNV) method in the range of 6102-5446.3 cm^{-1} leading to a coefficient of determination (R^2), root mean square error of prediction (RMSEP), a bias and residual predictive deviation (RPD) of 92.06%, 0.0361%, -0.004% and 3.57, respectively.

Table 2. Minimum (Min), maximum (Max), mean, and standard deviation (SD) for sugar-added and water-added in fresh orange juice samples in calibration set and validation set.

Type of Set	Addition	No. Samples	No. Spectra	Mean	Max	Min	SD
Calibration set	Sugar (g/ml)	61	122	0.22	0.43	0.00	0.11
Validation set	Sugar (g/ml)	25	50	0.17	0.43	0.00	0.13
Calibration set	Water (%)	64	128	38.66	61.93	0.00	18.13
Validation set	Water (%)	26	51	26.50	61.93	0.00	21.45

Table 3. Prediction statistics of sugar-added adulterated of orange juice by PLS model.

Pre-treatment	Wavenumber Range (cm^{-1})	PLS Factor	Calibration set			Validation set			
			R^2	RMSEE	RPD	R^2	RMSEP	RPD	Bias
Vector Normalization (SNV)	6102- 5446.3	3	88.21	0.0396	2.91	92.06	0.0361	3.57	0.004

Table 3 shows the prediction statistics associated with the PLS model of the adulterated samples.

In addition to a very low error of prediction, the R^2 of between 92-96% implies that the model is sufficiently predictive to be used in applications including quality assurance [13]. The RPD is calculated from the ratio between the standard deviation of reference values in the validation set and the standard error of prediction. The RPD of 3.1-4.9 implies that the model is sufficient for screening [13].

Figure 2 shows a plot of the regression coefficients for the model generated on the sugar adulterated samples while Figure 3 shows the corresponding X-loadings. The large regression coefficients and X-loadings indicate molecular vibration bands that are important in defining the PLS model.

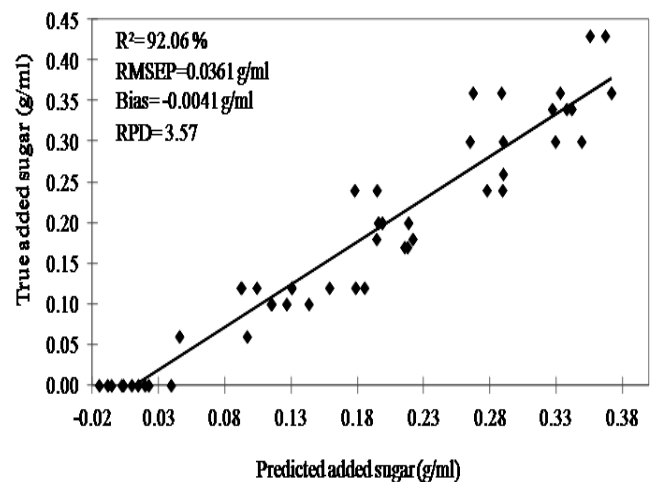


Figure 1. Plot of the observed versus predicted added sugar concentrations for the optimal NIR model generated on sugar adulterated juice samples of the validation set.

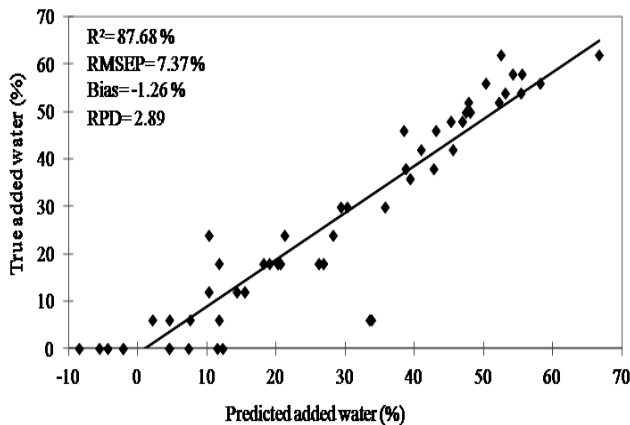


Figure 4. Plot of the observed versus predicted sugar concentrations for the optimal NIR model generated on the water adulterated juice samples of validation set.

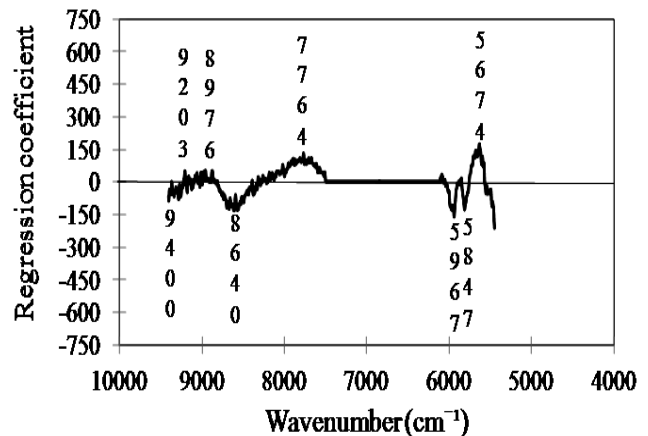


Figure 5. Plot of the regression coefficients versus NIR wavenumber for the optimal NIR model generated on the water adulterated juice samples.

Table 6. Wavenumber and bond vibration that illustrated high regression coefficients and X-loading of optimal prediction model of water added adulterated of orange juice.

Wavenumber (cm ⁻¹)	Wavelength (nm)	Wavelength (nm) [14]	Bond Vibration [14]	Structure [14]	Source (Fig. 5 & Fig. 6)
9400	1064	1064	N-H str. second overtone	RNH ₂	Regression coefficient
9280	1078	1080	2 x C-H str.+ 2xC-C str.	benzene	Factor 3
8640	1157	1152	C-H str. second overtone	CH ₃	Regression coefficient, Factor 3
5936	1685	1685	C-H str. first overtone	aromatic	Factor 3
5847	1710	1705	C-H str. first overtone	CH ₃	Regression coefficient
5674	1762	1765	C-H str. first overtone	CH ₂	Regression coefficient
5628	1777	1780	C-H str. first overtone	cellulose	Factor 3

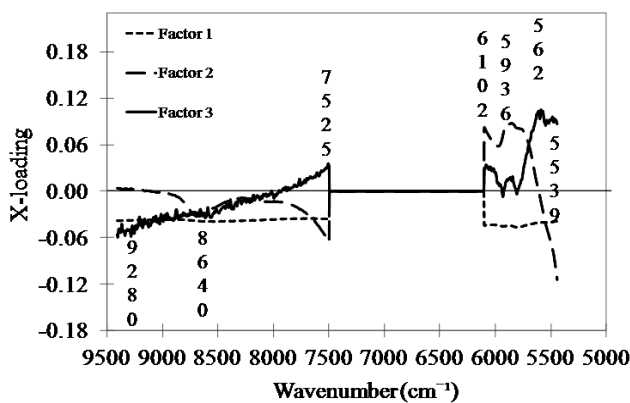


Figure 6. Plot of the X-loading versus NIR wavenumber for the optimal NIR model generated on the water adulterated juice samples.

ACKNOWLEDGEMENTS

The authors would like to express their gratitude to the Faculty of Engineering, King Mongkut’s Institute of Technology Ladkrabang for financial support for this research and to the Near Infrared Spectroscopy Research

Center for Agricultural Product and Food (www.nirsresearch.com) for providing the instruments.

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