



Application Note AN-RS-056

Quantification of methanol in contaminated spirits

Protecting consumers from contaminated beverages

An alarming global trend highlights the serious harm that can result from ingesting illegal, improperly distilled alcohol. Home-distilled spirits prepared using industrial solvents (i.e., wood alcohol) and presented as legitimate alcoholic beverages often contain methanol. Methanol causes blindness and can lead to death when ingested. This has led to fatal consequences around the world [1–3].

The breaking point for the Czech Republic came in September 2012. The sale of hard liquor was

temporarily banned after 20 people died from the consumption of spirits with dangerous levels of methanol [2]. After an exhaustive study using various screening tools, the Czech Republic adopted Raman spectroscopy as the method of choice for identifying and quantifying methanol in contaminated spirits.

This Application Note demonstrates how Raman spectroscopy can be employed as an efficient and rapid screening method for samples of rum contaminated with methanol.

INTRODUCTION

Raman spectroscopy is a fast and easy analytical tool for quantifying the amount of methanol contamination present in alcoholic beverages. It is an ideal method for the discrimination of very similar molecules like ethanol ($\text{CH}_3\text{CH}_2\text{OH}$) and methanol (CH_3OH), as shown in **Figure 1**.

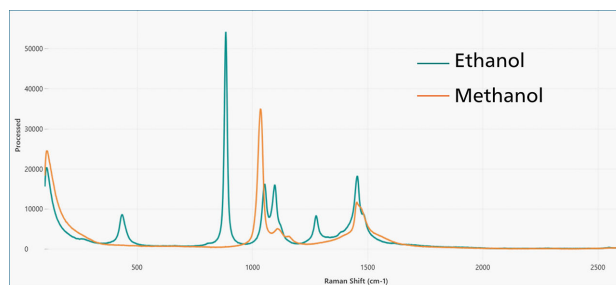


Figure 1. Raman spectra of pure ethanol (green) and pure methanol (orange).

The ability of Raman spectrometers to measure through containers and the lack of sensitivity to water make them better suited to measure methanol in beverage samples. These two key properties enable accurate detection of methanol down to approximately 1% by volume in the field with no need to open the bottles for testing. In the lab, the i-Raman NxG and SpecSuite software elevate the detection capabilities of Raman spectroscopy by adding the ability to quantify adulterants. (**Figure 2**)



Figure 2. The i-Raman NxG combined with SpecSuite software is ideally suited for rapid detection and quantitative screening of dangerous adulterants in alcoholic beverages.

EXPERIMENT

This example study measures commercially available coconut rum that is spiked with methanol in concentrations between 0.33% and 5.36%. The i-Raman NxG 785H with a fiber-optic probe is used to collect Raman spectra of the mixtures (**Figure 3**). **Table 1** lists the relevant equipment and instrument settings used for this application study.

The peak at around 1000 cm^{-1} (highlighted by the inset of **Figure 3**) visibly increases with increasing concentration of methanol, becoming significant at approximately 1%.

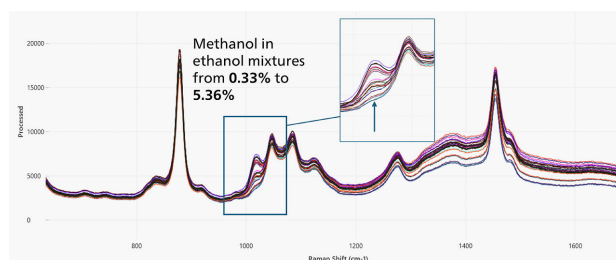
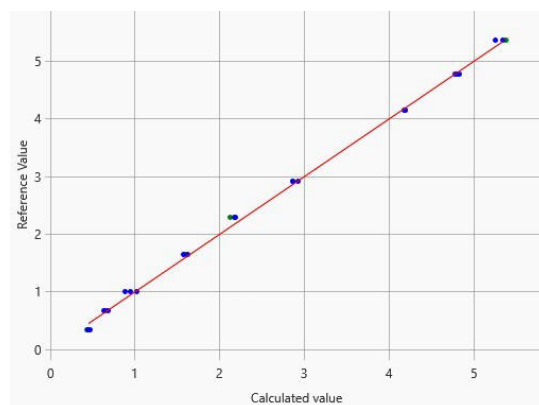


Figure 3. Raman spectra of methanol-laced rum with varying concentrations of methanol. Inlay: Peak intensities reflect changes in the methanol:ethanol ratio.

Table 1. Experimental parameters.

Equipment	Acquisition settings	
i-Raman NxG 785H	Laser Power	100
Vial holder	Int. time	1 s
SpecSuite Software	Average	1

This data is analyzed with SpecSuite software, and a partial least squares (PLS) regression model is developed on normalized data. The two-factor model developed over the range from 980–1040 cm^{-1} gives the calibration curve shown in **Figure 4**, which has an error of cross-validation (SECV) of 0.0794 (**Table 2**). The R^2 value of 0.9980 shown in **Table 2** means that the Raman method used here can be used to confidently quantify the amount of methanol in a mixed alcohol sample.

**Figure 4.** PLS regression model to predict the amount of methanol in rum.**Table 2.** Regression parameters used for the development of the PLS model to determine methanol in rum with the i-Raman NxG 785H.

Parameter	Value
Spectral processing	Mean centering Savitzky-Golay derivative
R^2	0.9980
SEC	0.0681
SECV	0.0794

CONCLUSION

These results validate that Raman spectroscopy can be used for rapid, quantitative screening of dangerous adulterants in alcoholic beverages. This technique can

be expanded to investigate adulteration in other media such as food, petroleum, and pharmaceutical drugs [4].

REFERENCES

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2. Spritzer, D.; Bilefsky, D. Czechs See Peril in a Bootleg Bottle. *The New York Times*. USA September 17, 2012.
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4. Gryniewicz-Ruzicka, C. M.; Arzhantsev, S.; Pelster, L. N.; et al. Multivariate Calibration and Instrument Standardization for the Rapid Detection of Diethylene Glycol in Glycerin by Raman Spectroscopy. *Appl Spectrosc* **2011**, *65* (3), 334–341. <https://doi.org/10.1366/10-05976>.

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CONFIGURATION



i-Raman NxG 785H

The i-Raman NxG 785H is ideal for routine quality control and process monitoring, especially where speed, stability, and reliability are essential. It offers a balanced combination of performance and efficiency to measure Raman scattering from 100 – 2800 cm^{-1} . Designed to support high signal throughput, this flexible system is ideal for monitoring chemical and polymer reactions, optimizing processes, and performing content uniformity testing of pharmaceutical tablets. The i-Raman NxG 785H can be easily adapted for see-through measurements even through opaque containers, adding to its versatility.

The i-Raman NxG 785H is the go-to solution for teams looking for reliable Raman analysis in challenging operational environments.

Discover why the i-Raman NxG is the perfect way to gain control of your quality control measurements:

- High-sensitivity spectrometers deliver results in seconds and can detect the faintest Raman signals
- Flexible fiber optic probe compatible with a wide array of accessories, including a vial holder, cuvette holder, immersion probe, and see-through adapter
- Powerful SpecSuite software for easy Raman data collection in addition to quantitative model building, identification with spectral libraries, and routine analysis
- Compact and stackable to save on valuable bench space.



Vial Holder Accessory

Vial holder accessory for use with lab-grade Raman probes with shaft diameter of 9.5 mm. Compatible with 15 mm-diameter vial. 6-pack of 15 mm borosilicate glass vials included.