

Quantitative Analysis of a Water-soluble Polymer Using the i-Raman EX Spectrometer

Vibrational spectroscopy is a well-established, powerful tool for polymer characterization.[1,2] Infrared and Raman spectroscopy are complementary techniques that provide a molecular fingerprint and are capable of both qualitative and quantitative analyses.[3] In general, infrared spectroscopy provides an excellent selection of functional groups associated with the polymeric composition, making it widely used in polymer identification and structural characterization. Raman spectroscopy also provides access to a selection of functional groups which enables determination of chemical composition and structure. Typically Raman spectroscopy probes non-polar chemical groups, such as multiple C-C backbone

vibrations and strong characteristic aromatic bands. These characteristic Raman bands provide detailed information on chain conformation as well as easy discrimination of aromatic-based polymeric systems.[3]

A simple, robust quantitative analysis is required to determine the extent of functionalization of polystyrene to create a water-soluble polymer. The new generation of Raman instrumentation provides a small, cost-effective, user-friendly high-performing system. The strong characteristic Raman bands of the aromatic group along with the negligible signal from water enables the simple measurement of the final aqueous polymer to determine the extent of reaction.

Initial evaluation of Raman spectroscopy to quantitate polymer functionalization reaction

The Raman spectra of the initial starting polymer and the fully functionalized polymer both show strong characteristic aromatic bands for the initial and fully functionalized polymers (see **Figure 1**). The strongest spectral features for the starting and fully functionalized polymers are observed as isolated

bands at 1002 and 1132 cm^{-1} , respectively. These unique spectral features demonstrate that Raman spectroscopy is well suited for the desired analysis and will support either a simple classic univariate band ratio method or a PLS-based (Partial Least Squares) quantitative method.

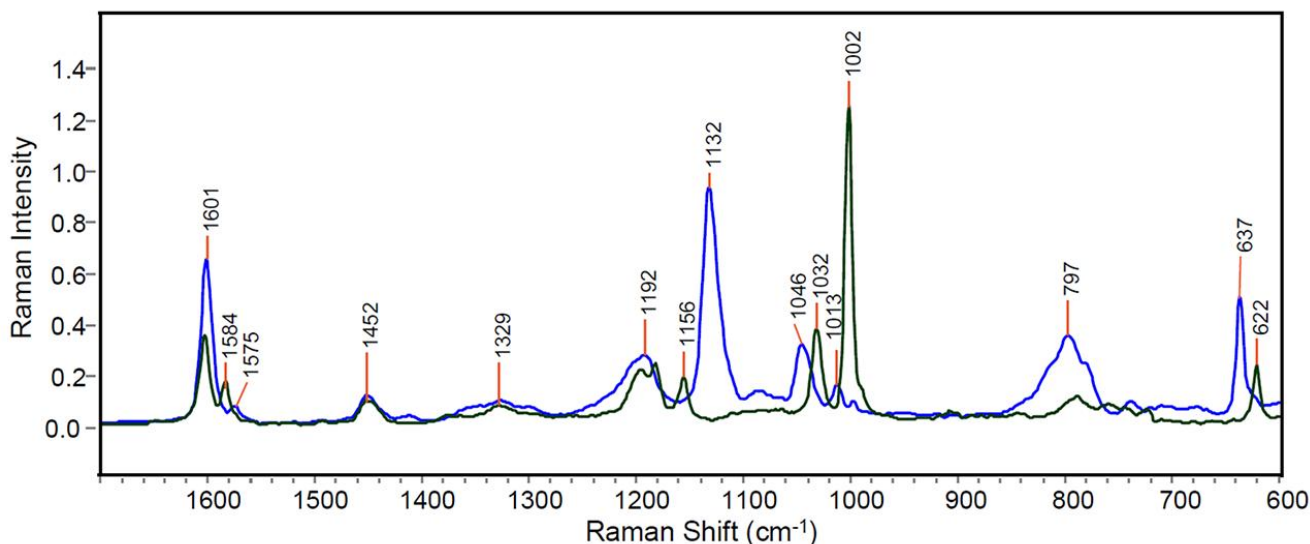


Figure 1. The FT-Raman spectra of the two pure polymer components: starting material polystyrene (green trace) and the fully functionalized polymer (blue trace) measured at 4 cm⁻¹ resolution with 1064 nm excitation

Analysis deliverables

Well-designed chemometric analysis methods are needed that are appropriate for a plant quality control (QC) laboratory to determine the extent of the final polymer functionalization. This requires an expert-based system in which the interface for the QC analyst is a simple push-button work flow.

The polystyrene functionalization reaction includes both an organic and aqueous phase where the desired water-soluble polymer is present in the aqueous phase. The analysis must include a

preliminary qualitative (classification) model to confirm the correct phase has been sampled for the final analysis. The subsequent quantitative analysis for the aqueous functionalized polymer will use a partial least squares for a single variable (PLS1) analysis to determine the percent functionalized polymer with well-defined accuracy and precision. This strategy ensures the analysis provides a simple, easy-to-use system appropriate for the multiuser plant analytical laboratory environment.

EXPERIMENTAL

Instrumentation

The Raman spectra are measured using 1064 nm excitation at ca. 495 mW, 180° backscattering geometry, and a spectral range from 250 to 2500 cm⁻¹ with a resolution of 9.5 cm⁻¹ at 1296 nm. The instrument components included a B&W Tek i-Raman EX spectrometer with a thermoelectrically cooled 512 element InGaAs detector array. A fiber optic is interfaced to an enclosed cuvette/vial sample compartment (NR-LVH, B&W Tek). The aqueous

polymer samples (ca. 10-20 weight percent solids) are transferred into borosilicate screw cap vials (height 4.5 cm, Outer Diameter 1.5 cm, Inner Diameter 1.1 cm) for the Raman measurements. Data acquisition is made using BWSpec v4.04 and includes a dark subtraction. Solutions are measured using an exposure time of 500 ms and 264 accumulations for a total measurement time of five minutes.

Software

A multivariable analysis software package, BWIQ version 3.0.6 (B&W Tek) was used to develop both the qualitative discrimination and quantitative analyses

methods. The software included standard pattern recognition, classification and quantitative chemometric tools.[4]

Data analysis

A classification model is used to ensure that the sample consisted of the aqueous layer containing the polymer rather than the aliphatic organic layer. The analysis used to develop the PCA-MD classification model included: sample partition (Kennard-Stone algorithm), spectral preprocessing (mean centering,

max value normalization), variable selection (spectral range) and classification (Principal Component Analysis-Mahalanobis Distance). The parameters selected for the final chemometric PCA-MD classification model for the measured Raman spectra are shown below (see **Table 1**):

Table 1. Parameters for the PCA-MD classification model for the measured Raman spectra.

Parameter	Component
Data Pretreatment	Center
Normalization	Max value normalization
Regression	PCA_MD
Principal components	3
Wavenumber	650 – 1700 cm ⁻¹

Once the measured Raman spectra are confirmed to match that of the water-soluble polymer, a partial least squares model is employed to determine the extent of the polymer functionalization. Sixty different water-soluble polymers with varying extents of polystyrene functionalization were selected. The

systems varied in the degree of functionalization from 65-98% as determined by ¹H NMR spectroscopy. The parameters selected for the final chemometric PLS1 % quantitation model for the measured Raman spectra of the functionalized water-soluble polymer are shown below (see **Table 2**).

Table 2. Parameters for the PLS1 quantitative model for % functionalized polymer based upon Raman spectra.

Parameter	Component
Data Pretreatment	Autoscale
Spectral derivative	Savitzky-Golay second derivative order = 3, window size = 5
Regression	PLS1
Principal components	6
Wavenumber region	995-1200 cm^{-1}

The developed methods were saved in chemometric modeling markup language (CMML) for easy transfer

to the manufacturing site.

RESULTS AND DISCUSSION

Initial evaluation of the functionalized polymer Raman data

The Raman spectra measured with the i-Raman EX spectrometer of the low, medium and highly functionalized polymer exhibits excellent specificity in the 900-1300 cm^{-1} region (see **Figure 2**). An additional feature observed at 981 cm^{-1} derives from an inorganic species. A second derivative spectral pretreatment provides a well-defined baseline and resolution of the desired aromatic peaks for the initial and functionalized polymer.

A simple peak height ratio analysis is used to evaluate the quality of the Raman data to determine polymer functionalization. Twelve samples that varied in the extent of polymer functionalization as determined by

^1H NMR spectroscopy were selected. Next the single point peak height intensities of the second derivative spectra at 1126 and 1005 cm^{-1} are determined. The peak height analysis of the two components (i = initial, f = final) utilized the ratio of these two unique bands for component band f divided by the sum of the two component bands ($i + f$). The expected range of this ratio will be 0 to 1, where 1 corresponds to a 100% functionalized polymer. This simple univariate band ratio analysis resulted in very good linearity (see **Figure 2**) confirming the suitability of the Raman spectral data to develop a robust quantitative PLS1 model.



Figure 2. The spectral overlay on the left shows the Raman spectra and their second derivative spectra of three different polymers representative of low, medium and highly functionalized systems. The spectral feature at 981 cm⁻¹ (red star) derives from varying amounts of an inorganic species. The highly diagnostic bands at 1002 and 1132 cm⁻¹ from the starting and functionalized polymer species are identified with a green star. The correlation plot for the simple peak height ratio using the two diagnostic bands versus the ¹H NMR determined extent of functionalized polymer is shown on the right.

Standard selection

A well-designed calibration set includes the expected possible variation of future samples. This includes concentration ranges of constituents, as well as process-induced and environmental variability. For the PCA-MD classification of the organic and aqueous phase samples, this included multiple production

samples from both the organic and aqueous phases. For the final PLS1 quantitative analysis this included over seventy samples from initial research lab design-of-experiments, pilot plant batches, and production lots that were fully characterized by the ¹H NMR spectroscopy.

PCA-MD Discrimination analysis of organic and aqueous phases

Principal component analysis (PCA) is a chemometric method in which the original spectral data is transformed into a new coordinate system, which condenses the information found in the original inter-correlated variables into a few uncorrelated variables, called principal components. The PCA can be used to reduce the dimensionality by zeroing out one or more of the smallest principal components. This results in a lower-dimensional projection of the data that

preserves the maximal data variance.

This multivariate analysis reveals the internal structure of the data in a way that enables a simple explanation of the variance in the data. As such, it is a useful tool for visualization of datasets. In any principal component rotation, the first component accounts for the maximum proportion of the variance and subsequent components account for the remaining variance.

PCA clearly enables grouping or clustering the Raman spectra into different classes. The cumulative explained ratio plot quantifies how much variance is explained in the first N components. In this application, three (N) variables explain 98% of the variance. Cluster analysis methods aggregate the Raman spectra stepwise, based upon the similarity of their spectral features (variables). A simple distance measurement of the individual spectrum relative to the various clusters provides the pattern recognition. A standard distance metric includes the Mahalanobis

distance. Here, the Mahalanobis distance is a statistical tool that is coupled to PCA and can be considered as the distance of a point from the center of a principal component cluster. The smaller the Mahalanobis distance, the higher probability that the point belongs to the cluster.

A simple PCA-MD classification model was successfully developed using 37 calibration and 34 validation Raman spectra. The model was 100% accurate in classification of the training set Raman spectra for both the organic and aqueous phases.

PLS1-based quantitative determination of % functionalized polymer

The PLS regression is one of the most popular multivariate techniques presently used in vibrational spectroscopic methods for quantitative analyses. It is used to develop a linear model between two matrices, the Raman spectral data and the reference values. The PLS model finds the empirical relationship between the two matrices (X and Y) and enables prediction of a continuous property, such as the % polymer functionalization, from the training set. In the case of the Raman spectra of the standards, the variation in the 1002 cm^{-1} and the 1127 cm^{-1} spectral features are highly correlated with the % polymer functionalization.

The developed quantitative model for the % functionalized polymer employs a PLS1-based (partial least squares for a single variable) algorithm. The Raman spectrum uses the $995\text{-}1200\text{ cm}^{-1}$ spectral region which includes diagnostic bands from both the

starting and reacted polymer. Data pretreatment included a second derivative. The overall performance of the final quantitative model is summarized in the plot of the predicted (PLS) versus known ($^1\text{H NMR}$) % functionalized polymer (see **Figure 3**). 29 calibration (blue symbols) and 31 validation (red symbols) standards were used. Excellent linearity was observed with an R^2 of 0.95 for the calibration standards and 0.87 for the validation standards. The RMSE (root mean square error) is based upon the residuals of the calibration data and provides a simple measure of the predicted values. This was calculated for both the calibration (RMSEC) and validation (RMSEP) standards, with an RMSEC of 1.22 and an RMSEP of 1.43. The RME less than two (<2) demonstrates the model exhibits good predictive performance.

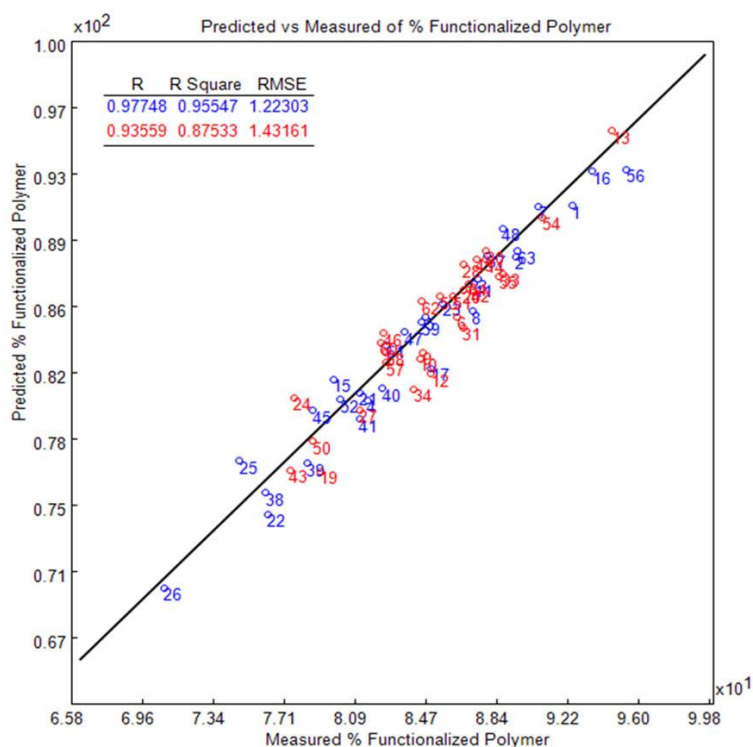


Figure 3. Plot of the predicted (PLS) versus the measured (1H NMR) % functionalized polymer. The blue symbols depict the calibration and the red the validation standards.

PLS1 quantitative Raman model accuracy and precision

The final developed model provides good linearity between the predicted (PLS) and assayed (¹H NMR) values for both the calibration and validation standards. **Table 3** summarizes the precision and accuracy performance of the quantitative PLS Raman model. This includes the RMSEP from the validation standards (see **Figure 4**) a standard single-day and lot

precision measurement and finally the average RMSE calculated for 16 subsequent manufactured lots (see **Figure 4**). The observed precision of 0.49% and the RMSE of 1.31 to 1.43 demonstrates the good statistical performance of this quantitative Raman method.

Table 3. Selected statistical measures of merit (precision, accuracy) for the PLS1 quantitative Raman model to determine polymer % functionalization.

Statistical merit	Value	Comments
RMSEP	1.43%	Precision: Figure 3 PLS model ¹
Precision	0.49%	Single day, single lot precision Raman
(Predicted - Assay) Avg. RMSE	1.31%	Accuracy: Figure 4 for 16 additional lots ¹



Figure 4. External validation of second manufacturing campaign batches. Raman predicted % functionalized polymer versus NMR determined value.

CONCLUSION

High-quality Raman spectra of an aqueous based water-soluble polymer (10-20% solids) are rapidly and easily measured using the i-Raman EX spectrometer. The Raman spectrum provides strong, unique bands for both the initial and fully reacted polymer. This enables development of a simple, robust quantitative analysis of the percent polymer

functionalization using a PLS1 chemometric model with the vendor-supplied software (BWIQ). The final Raman method exhibits excellent linearity, accuracy and precision. This Raman method is now used routinely in a multiuser plant quality control (QC) laboratory.

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CONFIGURATION



Spectromètre Raman portable i-Raman EX

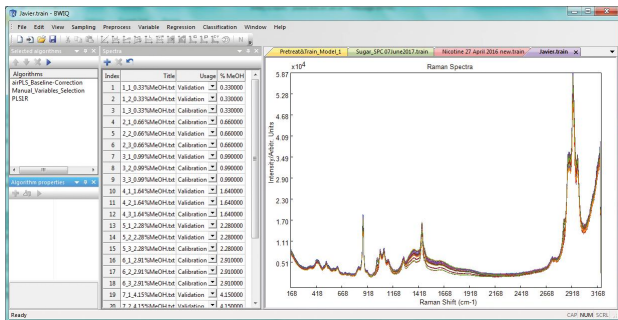
L'i-Raman[®] EX appartient à notre série primée de spectromètres Raman portables « i-Raman », équipée de notre laser breveté CleanLaze[®] avec excitation à 1064 nm. Ce spectromètre Raman portable utilise un détecteur à barrette InGaAs à haute sensibilité, refroidi par cryogénie thermoélectrique et large gamme dynamique et un spectrographe conçu pour un haut débit. Il fournit un rapport signal-bruit élevé sans induire d'autofluorescence, ce qui permet de mesurer une large gamme de produits naturels, d'échantillons biologiques (par exemple, cultures cellulaires) et d'échantillons colorés.

L'i-Raman EX couvre une plage spectrale de 100 cm^{-1} à 2500 cm^{-1} , ce qui autorise des mesures dans toute la région de l'empreinte digitale. Le faible encombrement et la légèreté de structure de ce système peu énergivore lui permettent d'effectuer partout des analyses Raman de qualité recherche. L'i-Raman EX est équipé d'une sonde à fibre optique et d'une table de translation XYZ avec support de sonde. Il peut être utilisé avec toute une série d'accessoires d'échantillonnage pour faciliter la mesure des échantillons les plus divers. Pour élargir le champ des possibilités d'analyse, il peut être utilisé avec notre logiciel d'analyse à variables multiples BWIQ[®] ou notre logiciel d'identification BWID[®]. Avec le i-Raman EX, vous avez une solution Raman pérenne de haute fidélité pour l'analyse qualitative et quantitative sans fluorescence.



Adaptateur de support de flacon

Adaptateur de support de flacon pour une utilisation avec la sonde Raman BAC100/BAC102 de qualité laboratoire, d'un diamètre de tige de 9,5 mm. Compatible avec des flacons d'un diamètre de 15 mm. Lot de 6 flacons en verre de borosilicate (15 mm).



BWIQ : logiciel d'analyse à variables multiples

BWIQ[®] est un progiciel très complet d'analyse multivariée de données spectrales, y compris des analyses exploratoires et qualitatives ainsi que des méthodes de régression quantitative. BWIQ supporte les méthodes chimiométriques classiques de classification et de régression, dont la régression des moindres carrés partiels (PLS), l'analyse en composantes principales (ACP) et l'analyse discriminante avec algorithmes machines à vecteurs de support (Support Vector Machine ou SVM) pour les blocs de données non linéaires. Il propose un flux de travail intuitif, de l'importation des données à l'évaluation, avec des graphiques complets pour étayer l'analyse et l'interprétation des modèles. BWIQ dispose d'outils de prétraitement qui permettent l'analyse optimale des blocs de données spectroscopiques Raman, NIR, LIBS et autres. Le modèle chimiométrique de BWIQ est utilisé avec QTRam[®]. C'est le progiciel chimiométrique idéal pour une utilisation en ligne avec des appareils de la série i-Raman[®] pour évaluer et analyser en temps réel des données de spectroscopie haute résolution.