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Quantitative vibrational spectroscopy on liquid mixtures: concentration units matter

We have addressed the contrasting results reported in the literature about the difference between volume- and mass-based concentrations units used for quantitative vibrational spectroscopy on liquid solutions. Conclusive evidence is provided that spectroscopic intensities in both near-infrared, mid-infrared, and Raman spectroscopy correlate linearly with volume-based concentration units rather than mass-based units (such as wt%), the latter units being overwhelmingly used in the literature and industrial practice. Density differences between mixture components are a major source of non-linearity between volume- and mass-based concentration units.

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Quantitative vibrational spectroscopy on liquid mixtures: concentration units matter†

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Quantitative vibrational absorption spectroscopies rely on Beer's law relating spectroscopic intensities in a linear fashion to chemical concentrations. To address and clarify contrasting results in the literature about the difference between volume- and mass-based concentrations units used for quantitative spectroscopy on liquid solutions, we performed near-infrared, mid-infrared, and Raman spectroscopy measurements on four different binary solvent mixtures. Using classical least squares (CLS) and partial least squares (PLS) as multivariate analysis methods, we demonstrate that spectroscopic intensities are linearly related to volume-based concentration units rather than more widely used mass-based concentration units such as weight percent. The CLS results show that the difference in root mean square error of prediction (RMSEP) values between CLS models based on mass and volume fractions correlates strongly with the density difference between the two solvents in each binary mixture. This is explained by the fact that density differences are the source of non-linearity between mass and volume fractions in such mixtures. We also show that PLS calibration handles the non-linearity in mass-based models by the inclusion of additional latent variables that describe residual spectroscopic variation beyond the first latent variable (e.g., due to small peak shifts), as observed in the experimental data of all binary solvent mixtures. Using simulation studies, we have quantified the relative errors (up to 10–15%) that are made in PLS modeling when using mass fractions instead of volume fractions. Overall, our results provide conclusive evidence that concentration units based on volume should be preferred for optimal spectroscopic calibration results in academic and industrial practice.

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1 Introduction

Mid-infrared (MIR), near-infrared (NIR), and Raman spectroscopy are powerful and common techniques in both qualitative and quantitative analytical chemistry, with widespread applications in academic, medical, and industrial environments.¹ For absorption spectroscopies, successful quantitative applications of these vibrational techniques rely mostly on Beer's well-known law relating spectroscopic intensity (e.g., absorbance) to chemical concentration, and for Raman spectroscopy there is also a linear relation between Raman intensities and concentration.² The calibration process typically involves samples with known analyte concentrations, as

measured by a reference method, and the use of multivariate data analysis techniques such as partial least squares (PLS) or multiple linear regression (MLR) to establish the quantitative relationship between spectra and concentrations.^{3,4} These regression techniques are linear in nature and therefore work best when the assumed linear relationship between spectroscopic intensity and concentration holds.

By and large, weight percent (wt%) is the concentration unit that is most often used for quantitative spectroscopic calibration in industrial practice and the scientific literature. However, Mark *et al.* demonstrated that NIR absorbance spectra are in fact linearly related to volume fractions of the various components of a mixture.^{5,6} Because volume- and mass-based concentration units are not necessarily related to each other in a linear fashion, the use of wt% may lead to substantially larger calibration errors than volume-based units. Recent work by Yan *et al.* contradicted the results of Mark *et al.* by reporting similar performance between mass- and volume-based multivariate calibration models for quantitative NIR and Raman spectroscopy on ternary solvent mixtures.⁷

To address these contrasting conclusions and clarify this issue of broad importance to analytical chemists, we here

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provide conclusive evidence that NIR, MIR, and Raman spectroscopic intensities are linearly related to volume-based concentration units. Using both experimental data on four different binary solvent mixtures and simulations, classical least squares (CLS) and PLS analysis shows that the difference in calibration errors between models based on mass or volume fractions correlates strongly with the density difference between the two solvents in each binary mixture. Under certain conditions (e.g., large density differences between solvents and high analyte concentrations), relative calibration errors of 10–15% are introduced by using mass rather than volume fractions as concentration unit. PLS regression shows that non-linearity due to the use of mass fractions leads to the inclusion of additional latent variables (LVs) in the PLS models. Such additional LVs make use of the non-linear residual spectroscopic variation beyond LV1, originating mainly from small peak shifts present in all investigated binary solvent systems. Our results demonstrate that concentration units based on volume are preferred over mass-based units for accurate quantitative spectroscopy on liquid systems.

2 Experimental

2.1 Sample preparation

Table 1 shows the binary liquid systems that were investigated in this study. For each system, 30 samples spanning the full compositional range (mass or volume fraction 0–1; see ESI† for compositional details) were prepared by gravimetric dosing of pure solvents into 40 mL glass vials using a high-throughput robotic system (Syntegon Technology GmbH, Waiblingen, Germany, formerly known as Bosch Packaging Technology), followed by vial capping and 60 s mixing in an orbital mixer. Of these 30 samples per binary system, 21 samples were used for development of calibration models and 9 samples were used as independent test samples for evaluation of model performance (e.g., root mean square error of prediction (RMSEP) values).

2.2 Instrumentation

NIR measurements were performed on a Bruker MPA instrument (Bruker Optics, Ettlingen, Germany) in transmission mode using sample vials (inner diameter 6.5 mm) kept at 27.5 ± 0.5 °C and the following acquisition settings: resolution 8 cm^{-1} , 16 scans per spectrum. A background spectrum was

recorded before each series of 30 samples per binary system. Raman measurements were performed on a Kaiser RXN-4 instrument (Kaiser Optical Systems, Inc., Ann Arbor, MI) by using a fiber-optic probe head coupled to a microscope objective (Leica HCX PL Fluotar, $50\times/0.55\text{ NA}$, long working distance of 8 mm) to focus the excitation beam ($\lambda_{\text{exc}} = 785\text{ nm}$) through the glass wall of the sample vials. Acquisition settings: 10 s acquisition time per spectrum (5 accumulations of each 2 s); cosmic ray filtering and dark subtraction were both applied. The NIR and Raman measurements were automated by integration of the above-mentioned spectrometers into a high-throughput Lipos plat-form (Zinsser Analytic GmbH, Frankfurt, Germany) programmed by WinLissy 8.1 software. MIR measurements were performed manually on a Spectrum 100 instrument (PerkinElmer, Groningen, The Netherlands) in ATR mode (diamond crystal) with the following settings: resolution 4 cm^{-1} (spectral point spacing 1 cm^{-1}), 16 scans per spectrum. A background measurement was recorded before each series of 30 samples per binary system. Evaporation of liquid on the ATR crystal was prevented by positioning a spacer around the ATR crystal and filling the resulting well with enough liquid sample. MIR ATR spectra were corrected for refractive index-dependent optical penetration depths using the “extended ATR correction” in Bruker OPUS software 7.5, with known refractive index values for the pure components and using the Arago-Biot equation for calculating the refractive index values of the mixtures.

2.3 Software

All data analyses were performed using MATLAB R2019b (The MathWorks, Natick, MA). CLS regression and 2D correlation spectroscopy (2D-COS) were performed with MATLAB routines developed in-house. Principal component analysis (PCA) and PLS regression were performed using PLS_Toolbox 8.8 software (Eigenvector Research, Inc., Manson, WA) running under MATLAB.

2.4 Data analysis

The following spectral regions were used for analysis: NIR, $6400\text{--}10\,200\text{ cm}^{-1}$; Raman, $250\text{--}1150\text{ cm}^{-1}$ (with slight variations per binary system); MIR, $600\text{--}1700\text{ cm}^{-1}$ (with slight variations per binary system), except for the chloroform–heptane system which was analyzed in the spectral region $1150\text{--}1550\text{ cm}^{-1}$. CLS coefficients were calculated for each binary component by using the pure component spectra and a

Table 1 Binary liquid systems investigated in this study, their corresponding differences in component densities, and their reported excess molar volumes

| System | Comp. 1 density ^a (kg L^{-1}) | Comp. 2 density ^a (kg L^{-1}) | $\Delta\rho$ (kg L^{-1}) | V^{E} at $x_1 = 0.5$ ($\text{cm}^3\text{ mol}^{-1}$) |
|--------|---|---|-------------------------------------|---|
| 1 | Chloroform 1.4832 | Heptane 0.6837 | 0.80 | +0.50 (ref. 8) |
| 2 | Chloroform 1.4832 | Toluene 0.8669 | 0.62 | +0.06 (ref. 9) |
| 3 | Toluene 0.8669 | Heptane 0.6837 | 0.18 | +0.15 (ref. 10) |
| 4 | MEK 0.8054 | Heptane 0.6837 | 0.12 | +0.80 (ref. 11) |

^a CRC Handbook of Chemistry and Physics, 72nd ed.; Boca Raton, FL, 1991–1992.



variable 0th order baseline offset to fit the mixture spectra of each binary system. The resulting coefficients of the 21 calibration samples were subsequently regressed against the known mass and volume fractions of each component to calculate RMSECV values (using 7-fold Venetian blinds as cross-validation method). The regression models were then used to calculate RMSEP values for the 9 independent samples. For PCA, PLS, and 2D-COS, all spectra (NIR, Raman, MIR) were pre-processed by Savitzky–Golay 1st derivative (2nd order, filter width 15 points for NIR and 9 points for Raman and MIR) followed by mean centering.

3 Results and discussion

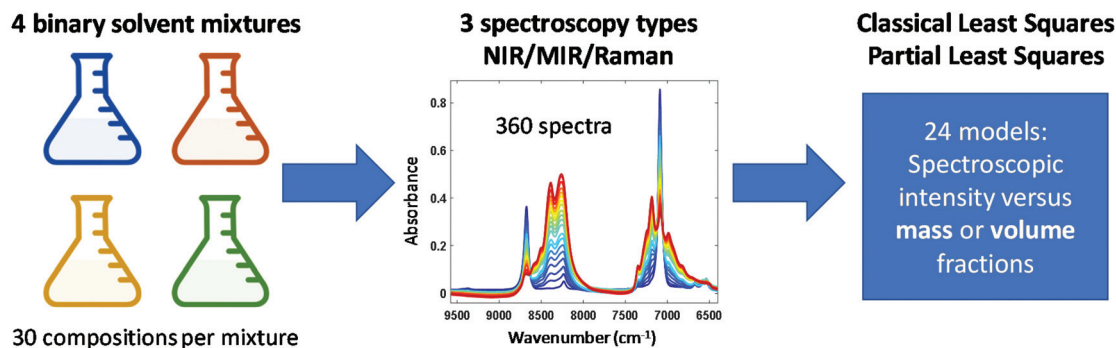
3.1 Selection of binary systems and their spectroscopic analysis

Scheme 1 shows the experimental design that we used to address the contrasting conclusions reached by Mark *et al.*^{5,6} and Yan *et al.*⁷ with respect to the difference between volume- and mass-based concentration units as reference data for multivariate calibration. The components of the four binary liquid mixtures shown in Table 1 were selected to provide a

broad range of density differences and therefore a varying degree of non-linearity between mass and volume fractions.

Moreover, the selected components are apolar and lack protic moieties such as OH groups, thereby preventing strong non-covalent interactions such as hydrogen bonds with pronounced and complex effects on vibrational spectra,^{12–14} which is undesirable for this study. Furthermore, by taking literature values for the excess molar volumes of these systems (Table 1), we calculated a relative volume change due to non-ideality of $0.33 \pm 0.29\%$ (average ± 1 s for the 4 systems) at equimolar concentrations. The degree of volume non-ideality is therefore small for these binary systems, hence this effect was not further taken into account in this study.

Before discussing the CLS and PLS calibration results in the following sections, we first present results from analysis of the mixture spectra themselves. Despite the relatively strong ideality of the selected systems, small but significant spectral non-ideality was observed in the NIR, Raman, and MIR spectra of all mixtures. This is exemplified in Fig. 1, which shows the results of PCA and 2D correlation spectroscopy (2D-COS) analysis^{15,16} for the NIR calibration spectra of the chloroform–toluene system. The plots of PCA scores (showing a concave curve) and loadings, as well as the characteristic pattern observed in the asynchronous 2D-COS plot,¹⁷ are indicative of



Scheme 1 Experimental design of our study.

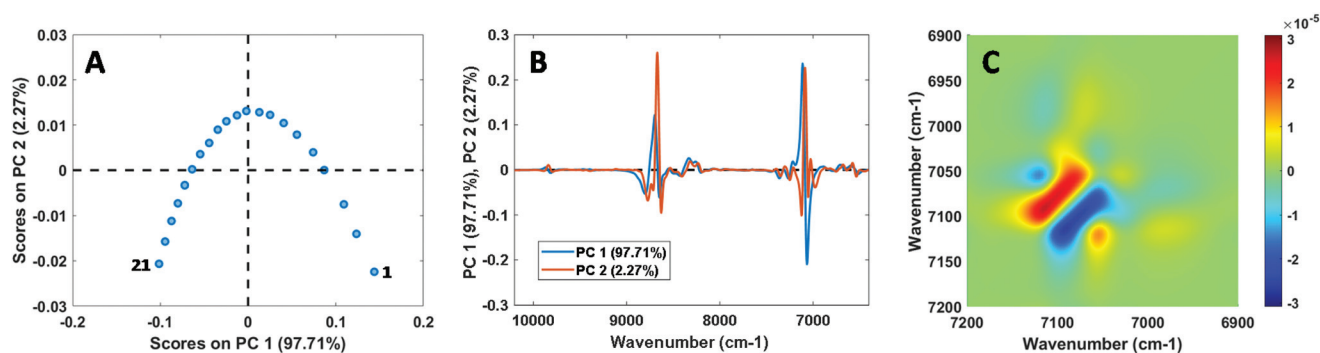


Fig. 1 Scores (A) and loadings (B) of the first 2 principal components (together explaining 99.98% of the variance) obtained by PCA on the 21 NIR calibration spectra of the chloroform–toluene binary system. The labels “1” and “21” in panel A correspond to the spectra of pure chloroform and toluene, respectively. (C) Asynchronous plot for the 7200–6900 cm^{−1} region obtained by 2D-COS of the chloroform–toluene system.



NIR peak shifts as a function of chloroform–toluene composition. Using PCA, PC1 and PC2 were found to explain $98.67 \pm 0.89\%$ and $1.20 \pm 0.85\%$ (average ± 1 s) of the spectral variance, respectively, for all systems combined excluding the MIR data on the chloroform–toluene system (which showed 90.60% and 5.92% variance explained by PC1 and PC2, respectively). Similar to these results, spectroscopic non-ideality is the rule rather than the exception for reported vibrational studies on liquid mixtures, even in seemingly ideal systems such as methanol–ethanol^{18,19} and binary aliphatic or aromatic hydrocarbon systems (*e.g.*, benzene–toluene and *n*-hexane–*n*-heptane),²⁰ and PCA has recently been proposed as a powerful tool for unsupervised screening of non-ideal mixtures.²¹ Spectral non-ideality results are generally interpreted as providing evidence for the existence of microheterogeneity (*i.e.*, homo- and heteroclusters) at the molecular level in mixtures.^{18–20}

Excess spectroscopy¹³ is another powerful tool to reveal non-ideal features in spectroscopic data on liquid mixtures. An excess spectrum ϵ^E (expressed as an absorption coefficient spectrum) is the difference between the measured spectrum of a mixture and that of the respective ideal mixture under identical conditions, and can be calculated as follows:¹³

$$\epsilon^E = \frac{A}{d(C_1 + C_2)} - (x_1\epsilon_1^* + x_2\epsilon_2^*) \quad (1)$$

where A is the spectroscopic intensity (*e.g.*, absorbance) of the mixture, d is the optical path length, C_1 and C_2 are the molarities of the two components, x_1 and x_2 are the mole fractions of components 1 and 2, and ϵ_1^* and ϵ_2^* are the molar absorption coefficients of the two components in their pure states, respectively.

To compare the degree of spectroscopic non-ideality for different liquid mixtures, Wrzeszcz *et al.*¹⁹ have defined the excess parameter ER as:

$$ER = \frac{\int |\text{mean}(\text{excess spectrum})|}{\int \text{mean}(\text{spectrum})} \times 100\% \quad (2)$$

where “mean” indicates the composition-averaged excess or normal spectrum. The ER values calculated in this way for the systems investigated in this study are shown in Table 2. It is clear that ER values strongly depend on the type of spectroscopy, with the average ER values increasing in the order NIR < Raman < MIR. The NIR-based ER value for toluene–heptane (5.75) is similar to the value reported for the analogous system benzene–hexane (6.69).²⁰ The ER values correlate

fairly well with the variance explained by PC2 ($R^2 = 0.55$) as obtained from the PCA results, confirming the report by Kiefer and Eisen that analysis of higher PCs can be used as screening tool for non-ideal mixture systems.²¹ The intensity of excess spectra, from which ER values are calculated, depends strongly on the magnitude of peak shifts as a function of composition. For example, the strong anti-symmetric CCl_3 stretching band at 744 cm^{-1} in the MIR spectrum of chloroform shifts to 758 cm^{-1} upon mixing with toluene (ESI Fig. S5†). Using spectral simulations, we calculated that the shift of this band alone produces an ER value of 47% for the MIR data of the chloroform–toluene system. Other bands in the MIR spectra with less pronounced peak shifts have lower ER values, leading to an average ER of 22.73% for this case (Table 2). Similarly, other large ER values in Table 2 are explained to a large extent by peak shifts.

3.2 CLS calibration results

CLS calibration was performed on the 12 investigated data sets (4 binary systems, each measured by NIR, Raman, and MIR spectroscopy) in order to directly compare our results with those obtained by Mark *et al.*^{5,6} and by Yan *et al.*,⁷ who also reported CLS results in their studies (see Data analysis section 2.4 for details about the CLS analyses). The clear difference between mass- and volume-based models obtained in this fashion is exemplified by the Raman results for the chloroform–heptane system shown in Fig. 2. Very strong non-linearity in the predicted *versus* measured plot is observed for the mass-based model (Fig. 2A), leading to a large RMSEP value for predicted mass fraction, whereas the volume-based model is much more linear in nature and therefore exhibits a much lower RMSEP value for predicted volume fraction (Fig. 2B). This confirms the NIR results reported by Mark *et al.*^{5,6} and proves that Raman spectroscopic intensities in liquid solutions are also linearly sensitive to volume-based concentration units. An overview of all CLS results, expressed as RMSEP values, is shown in Fig. 3. We observed that all 3 spectroscopic techniques show deviations from Beer's law when using mass fractions as concentration unit. However, the non-linearity and therefore RMSEP difference between mass- and volume-based models was found to decrease in going from systems with a high density difference between both components (*e.g.*, chloroform–heptane) to systems with a low density difference such as MEK–heptane (see also Table 1). In fact, when averaging the RMSEP difference results ($\text{RMSEP}_{\text{mass}} - \text{RMSEP}_{\text{volume}}$) over the 3 spectroscopic techniques, we found a strong dependence of the RMSEP on the density difference between solvents, as shown in Fig. 4, where the estimated RMSEP converges to zero for solvents of equal density.

This is not a surprising result, since the degree of non-linearity between mass and volume fractions depends on the density difference between the 2 components in a binary mixture. The contrasting results by Yan *et al.*,⁷ who reported similar CLS performance for mass- and volume-based units in quantitative NIR and Raman studies on ternary mixtures, are explained by the small density differences (about 0.1 and

Table 2 ER values calculated for the 4 binary liquid systems investigated in this study

| System | ER (%) NIR | ER (%) Raman | ER (%) MIR |
|--------------------|------------|--------------|------------|
| Chloroform–heptane | 2.16 | 20.26 | 16.38 |
| Chloroform–toluene | 19.85 | 6.58 | 22.73 |
| Toluene–heptane | 5.75 | 7.97 | 15.67 |
| MEK–heptane | 6.63 | 10.53 | 16.88 |



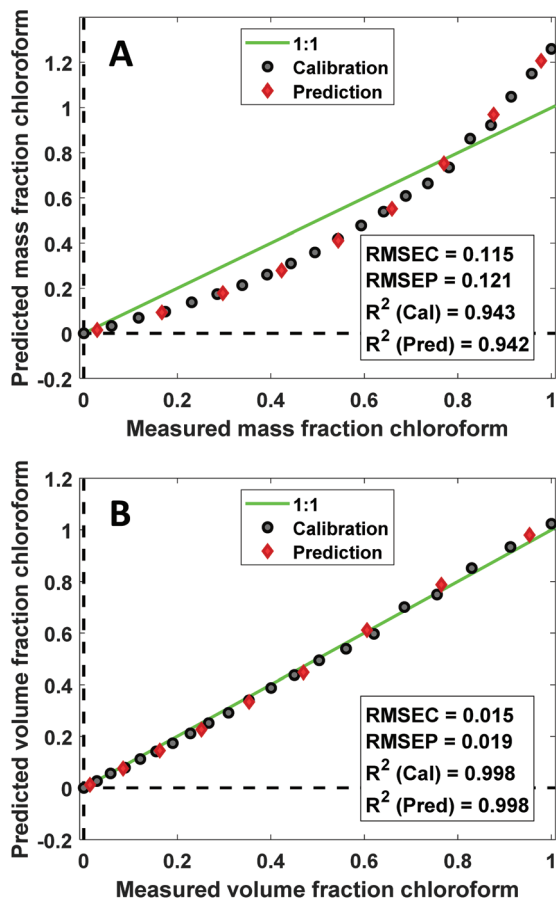


Fig. 2 Predicted vs. measured plots obtained from linear regression of Raman CLS coefficients against mass (A) and volume (B) fraction of chloroform for the chloroform–heptane binary system.

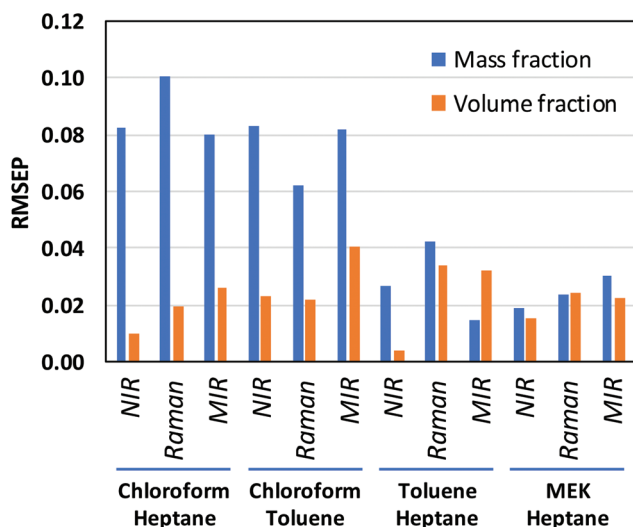


Fig. 3 RMSEP values obtained from linear regression models of CLS coefficients against mass and volume fractions. RMSEP values were averaged over the 2 components of each binary mixture.

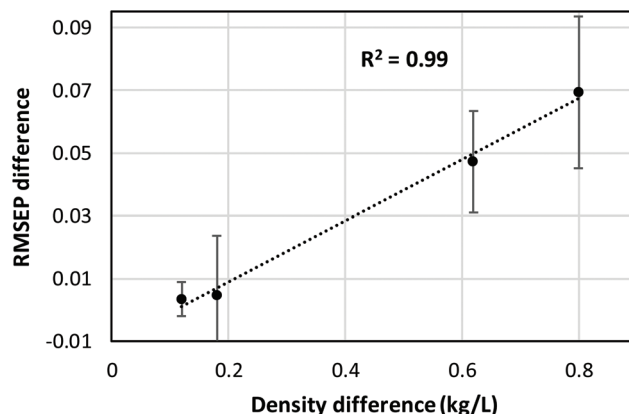


Fig. 4 Correlation between average RMSEP difference values ($RMSEP_{mass} - RMSEP_{volume}$) and density difference of the 2 components of each mixture. RMSEP difference values were averaged over the 3 spectroscopic techniques and over the 2 components of each binary mixture. Error bars represent 1 s.

0.2 kg L⁻¹) between the components in the ternary mixtures (benzene–cyclohexane–ethylbenzene and ethyl acetate–1-heptanol–1,4-dioxane, respectively) that they studied. At such low density differences between solvents, the degree of non-linearity between mass and volume fractions is only small and therefore only small differences between $RMSEP_{mass}$ and $RMSEP_{volume}$ are observed (Fig. 4).

3.3 PLS calibration results

PLS regression is nowadays by far the most often used multivariate calibration method for quantitative vibrational spectroscopy. It is therefore of interest to investigate how PLS deals with the non-linear relation between spectral intensities and mass fractions. Fig. 5 shows PLS results for the NIR data of the chloroform–toluene and toluene–heptane systems.

In the chloroform–toluene system with a large density difference (Fig. 5A), volume-based models only require 2 latent variables (LVs) to reach low RMSECV and RMSEP values, whereas mass-based models require 3 LVs to obtain such values. Thus, the non-linearity between NIR absorbance and mass-fraction leads to an increased number of LVs. This is expected due to the bilinear nature of the PLS method. In contrast, in the toluene–heptane system with much lower density difference (Fig. 5B), the RMSECV and RMSEP values at 1 and 2 LVs are very similar for mass- and volume-based models. These results are in agreement with the CLS results (*vide supra*) and prove that the use of mass-based concentration units in PLS regression leads to increased model complexity, which can be avoided by using volume-based units instead. Inspection of the PLS model loading weights (data not shown) reveals very similar spectral differences (*i.e.*, peak shifts) between the loading weights as observed in the principal components from PCA (Fig. 1B), indicating that PLS uses such spectral non-ideality in higher LVs to decrease model error. The contribution of this effect is more prominent in mass-



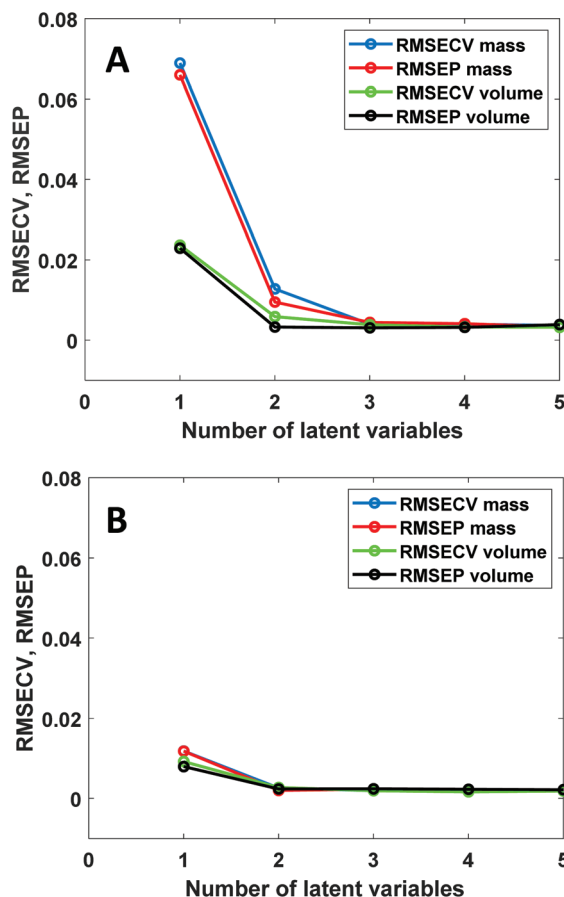


Fig. 5 RMSECV and RMSEP values as a function of the number of latent variables for PLS models built with mass and volume fractions as reference data. (A) NIR data of the chloroform-toluene system. (B) NIR data of the toluene-heptane system.

based models because the variance explained by LV1 in the concentration data is lower in that case than in models using volume-based concentration units. Note that unlike PLS, CLS cannot deal with peak shifts because in CLS each mixture spectrum is fitted with a combination of fixed pure component spectra. RMSEP values obtained by CLS are therefore larger than those obtained from PLS using 2 or more LVs.

To quantify the relative errors that are made in PLS modeling when using mass fractions instead of volume fractions, we simulated mixture spectra as noise-free Gaussian peaks scaling in intensity with volume fraction for 15 binary solvent systems with varying density differences (ESI†). PLS models were subsequently developed using mass fractions as reference data. Only 1 LV can be obtained for this error-free system because the first LV already accounts for 100% of the variance in the spectral data. Fig. 6 shows the resulting RMSECV errors expressed relative to the mean of the relevant mass fraction range. It is clear from these results that large errors up to 10–15% are made when both the density difference between the 2 components and the fraction ranges are high.

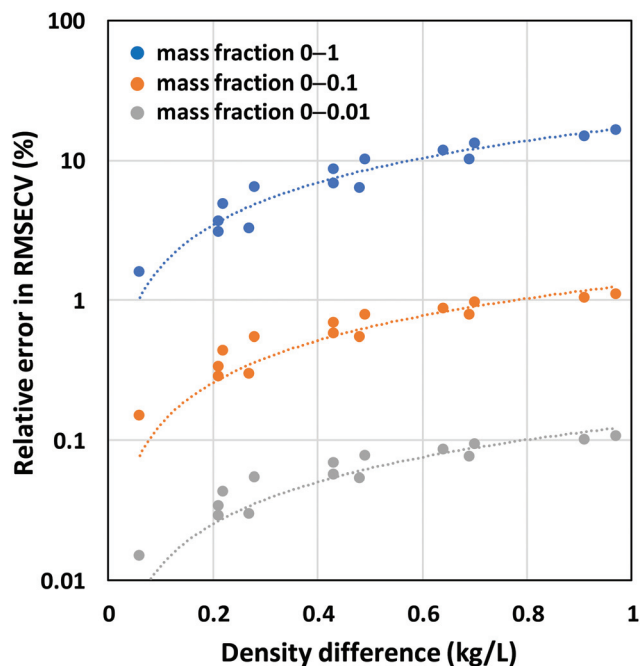


Fig. 6 Relative errors obtained by PLS using mass fractions as reference data and simulated Gaussian peaks scaling in intensity with volume fractions as spectral data.

4 Conclusions

We have demonstrated that NIR, Raman, and MIR spectroscopy are linearly related to volume-based concentration units, confirming the previous report by Mark *et al.*^{5,6} and extending their conclusion about NIR spectroscopy to MIR and Raman spectroscopy. Unnecessary calibration errors (extending up to 15% at high density differences and high analyte concentrations) are made when using mass-based units for quantitative vibrational spectroscopy, due to non-linearity between mass- and volume-based units, caused by density differences between mixture components. The implication of using mass-based units for PLS regression is that additional LVs will be required, which increases the risk of overfitting and reduces model robustness. Based on our results, it is recommended to use volume-based concentration units for optimal spectroscopic calibration of liquid solutions.

Author contributions

Henk-Jan van Manen: conceptualization, methodology, data curation, investigation, formal analysis, visualization, supervision, project administration, writing – original draft. Jan Gerretzen: conceptualization, methodology, data curation, investigation, formal analysis, writing – review and editing. Martijn Smout: methodology, investigation, resources, writing – review and editing. Geert Postma: writing – review and editing. Jeroen J. Jansen: methodology, supervision, writing – review and editing.



Conflicts of interest

There are no conflicts to declare.

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