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Vapor–Liquid–Liquid Equilibrium Measurements and Modeling of the Methanethiol + Methane + Water Ternary System at 304, 334, and 364 K

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ABSTRACT: New vapor–liquid–liquid equilibrium (VLLE) data for methanethiol (CH₃SH) + methane (CH₄) + water (H₂O) have been obtained at three temperatures (304, 334, and 364 K) and pressures up to 9 MPa. A “static-analytical” method was used to perform all of the measurements. The objective was to provide experimental VLLE data for CH₃SH with other natural gas contents at its crude form for which limited or no data are available in the open literature. Such kinds of data are required for the industrial modeling of sulfur emissions. It is observed from the experimental data that the solubility of CH₄ in the aqueous and organic phases increases with an increase of the total system pressure and decreases with an increase of the temperature. However, the solubility of CH₃SH in the aqueous and organic phases decreases slightly with an increase of the total system pressure and increases significantly with an increase of the temperature. The new VLLE data of this ternary system were compared with predictions of the cubic-plus-association equation of state. The model tends to underpredict the concentration of CH₃SH in all phases, particularly the vapor phase.

1. INTRODUCTION

Methanethiol (methyl mercaptan) is a common component encountered in natural gas, synthesis gas, and various refinery process streams. Its typical concentration in the host gas stream can range from several parts per million to 50% by volume.¹ Treatment processes need to be designed in order to remove not only H₂S and CO₂ but also organic sulfur species like methanethiol and other prohibited compounds. This is necessary because regulations for environmental protection force the petroleum industry to decrease the sulfur content in various petroleum fluids. Furthermore, any thiols/mercaptans (RSH), carbonyl sulfide (COS), and carbon disulfide (CS₂) not absorbed from the sour gas through the amine purification units complicate the process scheme for downstream liquid treatment units.² Knowledge of the phase equilibrium behavior and thermophysical properties of sulfur species mixtures with hydrocarbons/water is important to both process design and product specifications.

This work is a continuation of our previous work³–⁴ on the phase equilibrium measurements of systems containing thiols (mercaptans) with water and hydrocarbons. Such kinds of data are highly important for the design of new separation processes and the upgrade of existing processes. In this work, we provide new vapor–liquid–liquid equilibrium (VLLE) data for methanethiol (CH₃SH) + methane (CH₄) + water (H₂O) at three temperatures (304, 334, and 364 K) and pressures up to 9 MPa. The new VLLE data of these ternary systems were modeled with the cubic-plus-association (CPA) equation of state (EOS) using no adjustable parameters optimized in the ternary system data.

2. EXPERIMENTAL SECTION

Methanethiol (CH₃SH, CAS Registry No. 74-93-1) was obtained from Aldrich and has a purity of 0.99 mole fraction. Methane (CH₄, CAS Registry No. 74-82-8) was obtained from Messer and has a purity of 0.998 mole fraction. Ultrapure water (H₂O) was produced in the laboratory using commercial equipment (Millipore, model Direct-Q). The electrical conductivity of Direct-Q water (ultrapure H₂O) is 5.495 × 10⁻⁶ S·m⁻¹ at 298.15 K. No further purifications of the chemicals were made. The experimental work was carried at Mines ParisTech, where a “static-analytical” technique-based apparatus consisting of an equilibrium cell equipped with one moveable rapid online sampler injector was used. The equipment was the same as that used by Zehioua et al.,⁵ and the procedure was identical with that of Coquelet et al.⁶ The liquid and vapor samples were analyzed using a gas chromatograph (Varian model CP-3800), equipped with a thermal conductivity detector (TCD) and a flame ionization detector (FID).

In all experiments at first, H₂O was loaded into the equilibrium cell in the absence of air followed by CH₃SH. Then CH₄ was added to reach the desired pressure. The required temperature was obtained by putting the cell into a thermoregulated oil bath. Once equilibrium was achieved; the vapor, aqueous (liquid), and organic (liquid) samples from the equilibrium cell were directly introduced to the gas chromatograph through an isothermally heated transfer line. Two 100 Ω platinum
probes (Pt100) were used for temperature measurements: they were inserted inside thermowells drilled into the body of the equilibrium cell at two different levels and connected to an HP data acquisition unit. They were periodically calibrated against a 25 Ω reference platinum resistance thermometer (Tinsley Precision Instruments, France). The resulting accuracy in the temperature measurements was estimated to be within \( u(T, k = 2) \approx \pm 0.04 \) K. Pressures were measured by means of a Druck pressure transducer at 0.1−10 MPa, which was maintained at 353.15 K. The pressure transducer was calibrated against a dead-weight pressure balance (Desgranges & Huot S202S, CP 0.3−40 MPa, Aubervilliers, France). Accuracies in the pressure measurements have been estimated to be within \( u(P, k = 2) = \pm 0.003 \) MPa. The gas chromatograph detectors were calibrated using chromatographic syringes with maximum mole number uncertainties of 2% in the TCD and 1.5% in the FID; thus, the maximum uncertainty as mole fraction is \( u_{\text{max}} = 0.006 \).

The gas chromatograph generated peaks of the individual components [CH₄ (FID), CH₃SH (FID), and H₂O (TCD)] at specific retention times, which were used to calibrate the detectors against a 25 Ω reference platinum resistance thermometer (Tinsley Precision Instruments, France). The resulting accuracy in the temperature measurements was estimated to be within \( u(T, k = 2) = \pm 0.04 \) K. Pressures were measured by means of a Druck pressure transducer at 0.1−10 MPa, which was maintained at 353.15 K. The pressure transducer was calibrated against a dead-weight pressure balance (Desgranges & Huot S202S, CP 0.3−40 MPa, Aubervilliers, France). Accuracies in the pressure measurements have been estimated to be within \( u(P, k = 2) = \pm 0.003 \) MPa. The gas chromatograph detectors were calibrated using chromatographic syringes with maximum mole number uncertainties of 2% in the TCD and 1.5% in the FID; thus, the maximum uncertainty as mole fraction is \( u_{\text{max}} = 0.006 \).

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### 3. MODELING SECTION

The experimental data were modeled using the CPA EOS. The CPA EOS is given in terms of the pressure as a sum of the SRK EOS and an associating contribution in eq 1. The association term in this equation is a simpler but mathematically identical version of the term used in statistical association fluid theory.

This form was proposed by Michelsen and Hendriks. The CPA EOS proposed by Kontogeorgis et al. can be expressed for mixtures in terms of pressure as

\[
P = \frac{RT}{V_m - b} - \frac{\alpha(T)}{V_m(V_m + b)} - \frac{1}{2} \frac{RT}{V_m} \left(1 + \rho \frac{\partial \ln g}{\partial \rho} \right)
\]

where \( V_m \) is the molar volume. The key element of the association term is \( X_{iA} \), which represents the fraction of sites (type A) on molecule \( i \) not bonded with other active sites, while \( x_i \) is the mole fraction of component \( i \). \( \alpha(T) \) and \( b \) are the energy and covolume parameters of the SRK term. \( X_{iA} \) is related to the association strength \( \Delta_{AB} \) between two sites belonging to two different molecules, e.g., site A on molecule \( i \) and site B on molecule \( j \); more details were presented in previous studies.

The CPA EOS for thiols (mercaptan-containing systems) was described in our previous publications. The 4C association scheme was used for H₂O according to the terminology from Huang and Radosz. Furthermore, CH₃SH was considered to be a non-self-associating fluid but capable of cross associating with H₂O (solvation). For this reason, one negative association site was assumed on every CH₃SH molecule. The cross-association parameters were calculated using the

<table>
<thead>
<tr>
<th>Pressure</th>
<th>Temperature</th>
<th>( x_{CH_4} )</th>
<th>( x_{CH_3SH} )</th>
<th>( x_{H_2O} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.0 MPa</td>
<td>303.15 K</td>
<td>0.1</td>
<td>0.1</td>
<td>0.8</td>
</tr>
<tr>
<td>2.0 MPa</td>
<td>303.15 K</td>
<td>0.2</td>
<td>0.2</td>
<td>0.6</td>
</tr>
<tr>
<td>3.0 MPa</td>
<td>303.15 K</td>
<td>0.3</td>
<td>0.3</td>
<td>0.4</td>
</tr>
</tbody>
</table>

\( \sigma \) stands for the standard deviation on each mole fraction.
modified CR-1 rule. The CPA EOS pure-component parameters \((b/L\cdot mol^{-1}, \Gamma/K, \text{and} \ c_i)\) for \(CH_3SH\), \(CH_4\) and \(H_2O\) are presented in Table 2. The \(a_0\) parameter is tabulated as \(\Gamma = a_0/Rb\). The binary interaction parameters for \(CH_4 - H_2O\), \(CH_3SH - CH_4\) and \(CH_3SH - H_2O\) are presented in Table 3. The parameters were taken from the literature and applied to predict the VLLE of \(CH_3SH + CH_4 + H_2O\).

4. RESULTS AND DISCUSSION

In our previous work, we performed CPA calculations for Henry’s law constant of methanethiol (MM) in \(H_2O\) as a function of the temperature. It has been observed that, by taking into account the cross-associating effect in the \(CH_3SH + H_2O\) system, the CPA results were improved. The \(CH_3SH + H_2O\) system has also been studied and discussed in the literature by Gillespie and Wilson. The authors observed a drastic change in the solubility of \(CH_3SH\) at higher temperature \((523-673 K)\) compared to the lower temperature \((313-423 K)\) data. We did not find any further data in the open literature for the ternary system investigated in this work, so a direct comparison is not possible. It is observed that the solubility of \(CH_4\) in the aqueous and organic phases increases with an increase in the total system pressure and decreases with an increase in the temperature. Furthermore, the solubility of \(CH_3SH\) in the aqueous and organic phases decreases slightly with an increase in the pressure and increases significantly with an increase in the temperature. It has also been observed that the solubility of \(CH_3SH\) is higher than that of \(CH_4\) in the aqueous phase in some cases. The solubilities of \(H_2O\) in the organic and gas phases were not detected by gas chromatography because they were under the limit of detection. The detection limit of the gas chromatograph detectors in terms of mole numbers are as follows: \(CH_3SH = 9.78 \times 10^{-9}\), \(H_2O = 3.32 \times 10^{-5}\), and \(CH_4 = 4.51 \times 10^{-8}\). However, Gillespie and Wilson showed that the solubility of \(H_2O\) in the gas phase varies inversely with the system pressure except at temperatures above 422 K. These authors also conclude that the solubility of \(H_2O\) in the liquid phase is slightly lower than the solubility in the vapor phase at three-phase saturation conditions.

The CPA EOS predictions for the VLLE of the \(CH_3SH + CH_4 + H_2O\) ternary system from 1 to 9 MPa pressure at 304, 334, and 364 K are shown in Figures 1–3, respectively. It is observed that the deviation between the experimental data and the CPA predictions is sometimes higher than 40%. Especially, the CPA EOS predictions for the vapor-phase composition are rather poor. However, one should keep in mind that the CPA results are pure predictions because no parameters were adjusted to the experimental data. From this point of view, the CPA rather satisfactorily predicts the vapor–liquid and vapor–liquid–liquid regions for the \(CH_3SH + CH_4 + H_2O\) ternary system at 363.68 K and 8.966 MPa, as plotted in Figure 4, which shows that model can qualitatively predict the complicated three-phase equilibria of this system.

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Table 2. CPA Pure-Component Parameters Used in This Work

<table>
<thead>
<tr>
<th>component</th>
<th>(T_c/K)</th>
<th>(b/L\cdot mol^{-1})</th>
<th>(\Gamma/K)</th>
<th>(c_i)</th>
<th>(\epsilon_{AB}/bar\cdot L^{-1}\cdot mol^{-1})</th>
<th>(\beta_{AB})</th>
<th>% AAD in (p^{\text{exp}})</th>
<th>% AAD in (p^{\text{cal}})</th>
</tr>
</thead>
<tbody>
<tr>
<td>CH3SH(^3)</td>
<td>469.95</td>
<td>0.0437</td>
<td>2266.2</td>
<td>0.8007</td>
<td>166.55</td>
<td>0.0692</td>
<td>0.69</td>
<td>0.47</td>
</tr>
<tr>
<td>CH4(^12)</td>
<td>190.56</td>
<td>0.0291</td>
<td>959.1</td>
<td>0.4472</td>
<td>363.3</td>
<td>0.079</td>
<td>0.35</td>
<td>1.97</td>
</tr>
<tr>
<td>H2O(^13)</td>
<td>647.29</td>
<td>0.0145</td>
<td>1017.3</td>
<td>0.6736</td>
<td>8.168</td>
<td>0.017</td>
<td>0.91</td>
<td>0.98</td>
</tr>
</tbody>
</table>

\(^a\)Average absolute deviation (AAD) is defined as \% AAD = \((1/n)\sum_{i=1}^{n}(X_i^{\text{cal}} - X_i^{\text{exp}})/X_i^{\text{exp}}\) × 100, where \(X\) stands for \(p^{\text{cal}}\) or \(p^{\text{exp}}\) and \(n\) is the number of experimental data points.

\(^b\)Cross-association between \(CH_3SH\) and \(H_2O\), \(CH_4\) and \(H_2O\), \(CH_3SH\) and \(CH_4\), and \(CH_3SH\) and \(H_2O\) has been considered using the modified CR-1 rule.

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Figure 1. VLLE of the \(CH_3SH + CH_4 + H_2O\) ternary system from 1 to 9 MPa pressure at 304 K: (○) \(CH_3SH\) in the organic phase; (△) \(CH_3SH\) in the vapor phase; (□) \(CH_3SH\) in the aqueous phase. Solid lines: CPA EOS predictions.

Figure 2. VLLE of the \(CH_3SH + CH_4 + H_2O\) ternary system from 1 to 9 MPa pressure at 334 K: (○) \(CH_3SH\) in the organic phase; (△) \(CH_3SH\) in the vapor phase; (□) \(CH_3SH\) in the aqueous phase. Solid lines: CPA EOS predictions.
adjustment of any parameter to the ternary experimental data.

5. CONCLUSION

New VLLE data for CH$_3$SH + CH$_4$ + H$_2$O were performed at three temperatures (303, 334, and 364 K) in a pressure range from 1 to 9 MPa. A "static-analytical" method was used successfully to perform all of the measurements. It is concluded that the gas solubility of CH$_3$SH is higher than that of CH$_4$ in the aqueous phase under similar temperature and pressure conditions. The CPA EOS has been applied for the representation of ternary systems containing CH$_3$SH. These modeling results for the ternary mixture of CH$_4$ + CH$_3$SH + H$_2$O are satisfactory considering that they are predictions without the adjustment of any parameter to the ternary experimental data.

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Notes
The authors declare no competing financial interest.

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