Research Article

Experimental and Thermodynamic Investigation on CO₂ Absorption in Aqueous MEA Solutions

Kai Zhu, 1 Changhai Yue, 1 Zhenhao Wei, 1 Jingjing Huang, 1 Meng Hu, 1 Yufan Ji, 1 Hanfei Liu, 1 Hao Zhu, 1 Wanxiao Guo, 1 Feng Zhou 2, 3 Chaoqun Yao, 3 and Yiping Huang 1

1 China Construction Industrial & Energy Engineering Group Co., Ltd., 6, Wenlan Road, Qixia District, Nanjing, 210023, China
2 National & Local Joint Engineering Research Center for Deep Utilization Technology of Rock-salt Resource, Faculty of Chemical Engineering, Huaiyin Institute of Technology, Huai’an 223003, China
3 Dalian National Laboratory for Clean Energy, Dalian Institute of Chemical Physics, Chinese Academy of Sciences, Dalian 116023, China

Correspondence should be addressed to Feng Zhou; zhoufeng@hyit.edu.cn and Yiping Huang; huangyiping@cscec.com

Received 24 June 2022; Accepted 23 August 2022; Published 13 October 2022

Academic Editor: Ghulam Rasool

Copyright © 2022 Kai Zhu et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The vapor-liquid equilibrium (VLE) of CO₂ in a reactive solvent is essential for the proper simulation and design of CO₂ absorption processes. This work presents a systematic investigation on CO₂ absorption in various aqueous monoethanolamine (MEA) solutions. CO₂ solubility in MEA was measured at 298, 313, 333, and 353 K with CO₂ partial pressure ranging from 34.5 to 78.0 kPa. A modified Kent-Eisenberg model was developed based on the measured solubility data, showing good predictions over the liquid phase speciation for the CO₂-H₂O-MEA system. We presented a new analysis based on the first-order difference curve of distribution profiles of the species. Based on the main reactions that occurred, the CO₂ absorption process was demonstrated to be divided into four regions with increasing CO₂ loading from 0 to 1. Accordingly, kinetic study was proposed to be conducted in the first region, whereas measuring of mass transfer in the first three regions. The findings in this work extend the existing knowledge of CO₂ absorption process in terms of speciation and can provide important guidance for further study of the process characteristics using aqueous amine absorbents.

1. Introduction

Since industrial activities have expanded over the past several decades, global warming and climate change have become increasingly important environmental issues in the twenty-first century (Reference [1]). Carbon dioxide (CO₂), as a greenhouse gas, mainly from large point sources, such as coal-based power plant and refinery off-gases, is the main contributor to the global warming and climate changes. Therefore, developing efficient technologies to capture and separate CO₂ is becoming increasingly popular. Chemical absorption with alkanolamines is the most suitable technique currently available for CO₂ removal [2–6]. Monoethanolamine (MEA), as a primary amine, is under intense research because of its various advantages, such as high chemical reactivity with CO₂, low cost of materials, and ease of production [7–11]. However, the application of this system is hindered by its large energy requirement in the regeneration process [12, 13], which reduces the output of a coal-fired power plant by 30-40% [14, 15].

Understanding the fundamentals of absorption/desorption deeply, including thermodynamics, kinetics, and mass transfer, is essential for dealing with the problem of extensive energy consumption; (Reference [16]). The thermodynamic properties play a pivotal role in the design and optimization of CO₂ capture processes, involving vapor-liquid equilibrium (VLE), solubility, enthalpy, as well as equilibrium speciation. As a result, a great deal of effort has been put into the determination of the thermodynamic properties of CO₂ absorption into MEA solution and the development of thermodynamic models to describe them. Mouhoubi et al. [17] developed a thermodynamic model for
the carbon capture process of DEEA, MAPA, and their mixtures, in which experimental data from the literature were compared favorably with the predicted and correlated data of vapor-liquid equilibrium (VLE) and heat of CO₂ absorption. Ma’ mun et al. [18] measured the protonation constant of MEA at a range of temperatures from 303 to 330 K and presented an empirical equation correlating the influence of temperature. Aronu et al. [19] reported a series of CO₂ solubility data for MEA, along with a rigorous equilibrium model that fits the experimental VLE data well. Ojala et al. [20] measured the absorption enthalpy and compared to predictive models in the literature, showing a high level of agreement. Additionally, as there are several reversible chemical reactions, it is necessary to study the speciation of the liquid mixture, which not only has an essential influence on the VLE [21] but is also indispensable for simulation models coupling mass transfer [19, 22].

Chemical speciation in the absorption system can be obtained by experimental measurement [23–26], thermodynamic model prediction [12, 19, 21, 27, 28], or molecular simulations [29]. Jakobsen et al. [23] and Bottinger et al. [24] showed that nuclear magnetic resonance (NMR) spectroscopy was capable of acquiring quantitative information on the species distribution, but unable to distinguish the protonated and unprotonated amines. Wong et al. [25], however, showed that Raman spectroscopy could give a clear distribution of all species in CO₂-loaded MEA solutions. In Li’s research, for the CO₂ capture system using aqueous HEPZ, the cyclic capacity, speciation with loading, and heat of reaction are predicted and analyzed using the developed thermodynamic model. [30] A simple VLE model based on Henry’s law and equilibrium of eight reversible reactions was proposed by Kent and Eisenberg [28] ignoring the activity coefficients to predict equilibrium compositions. As compared with NMR data, Hessen et al. [12] demonstrated that the refined electrolyte nonrandom two liquid (NRTL) model could give a good prediction of the speciation. Using Helmholtz energy equations of state combined with excess Gibbs energy models, Neumann et al. [31] formulated a model for describing reactive mixtures that reproduced experimental data of vapor-liquid equilibrium, homogeneous density, and speciation of partial binary/ternary systems involving monoethanolamine well. As reported by Balaji et al. [29], accurate species concentrations could only be obtained through molecular simulations in which bi-carbonate hydrolysis is included in the reaction pathway when loadings exceed 0.5 mol CO₂/mol MEA, but it can be neglected in cases where CO₂ loadings are lower than 0.5 mol CO₂/mol MEA.

In spite of the relatively large number of publications concerned with chemical speciation in the MEA system, all of these studies only reported species distribution profiles and did not provide detailed analyses of the evolution rate of the species under varying CO₂ loads. In our work, CO₂ solubility in MEA aqueous solution was examined at temperatures ranging from 298 to 353 K and CO₂ partial pressures ranging from 34.5 to 78.0 kPa. Based on Kent-Eisenberg model framework [28], a modified model was developed using measured solubility data. Based on the obtained thermodynamic model, the species distributions of a 20 wt.% MEA system under different CO₂ loadings are well predicted. The first-order difference curve of species distribution profiles was derived as a new approach to analyzing the absorption process. Therefore, four speciation regions were first proposed based on the usage of a first-order difference method dealing the equilibrium concentration. It is noteworthy that the results obtained in this work not only contribute to the existing knowledge regarding speciation of CO₂ absorption process but also provide a basis for further study of the characteristics of aqueous amine absorption processes.

2. Materials and Methods

2.1. Reagents. Monoethanolamine (MEA) with a mass fraction purity of >99% was supplied by Kernel Co. Ltd. 20 wt.% (3.285 mol/L) aqueous MEA solution was prepared by mixing amine and deionized water. 30 vol.% carbon dioxide (CO₂) balanced with nitrogen (N₂) was supplied by Dalian Special Gas Co. Ltd.

2.2. Measurement of Solubility of CO₂. The solubility of CO₂ was measured using the experimental setup shown in Figure 1. The equilibrium cell with an inner volume of 130 ml was made up of stainless steel. A water circulating bath was used to control the temperature of the equilibrium cell. A magnetic stirrer was placed under the equilibrium cell to maintain a constant stirring speed of the absorption solution. A mass flow controller (0–500 mL/min of N₂ at 0°C, 101.325 kPa) was used to deliver the gas mixture at a constant flow rate. A back pressure valve and pressure sensor were used to maintain the outlet pressure of the equilibrium cell at the required pressure. A gas chromatography (GC) was used to measure the CO₂ concentration of the outlet gas after absorption as shown in Figure 1. A CarboPLOT-P7 column (12.5 m × 530 μm × 20 μm, Agilent) and thermal conductivity detector (TCD) were installed in the GC. The
temperature of the column and TCD were set at 303 and 523K, respectively. All the experiments were carried out over a temperature range of 298 to 353 K and a CO₂ partial pressure range of 34.5 to 78.0 kPa. The total amount of CO₂ dissolved in the absorbent was calculated by integrating differences between the concentrations of injected CO₂ and emitted CO₂. The amounts of CO₂ injected into the tank and emitted from the tank were calculated by the ideal gas equation. Thus, the CO₂ dissolved in the absorbent can be determined through equation (1). CO₂ loading (α), defined as the ratio of mole of absorbed CO₂ to mole of absorbent, can be calculated using equation (2).

\[ n_{\text{CO}_2, \text{abs}} = \frac{p}{RT} \int (Q_{\text{in}}C_{\text{in}} - Q_{\text{out}}C_{\text{out}}) \, dt \]  
\[ \alpha = \frac{n_{\text{CO}_2, \text{abs}}}{n_{\text{MEA}}} \]

2.3. Thermodynamic Framework. In order to understand the chemical speciation in the solution with increasing CO₂ loading, the bulk concentrations of species are required. Thus, the chemical reaction process, the vapor-liquid equilibrium model (VLE), and some model parameters are required as follows.

2.3.1. Chemical Reaction Process. In the absorption process of CO₂ into MEA [32–34], some species exist in the systems, including CO₂, H₂O, RNH₂, RNH+ 3, RNHCOO−, H₃O⁺, OH⁻, HCO₃−, and CO₃²−. Reactions as follows may occur in the liquid phase [23, 24, 34].

Carbamate formation reaction:

\[ \text{CO}_2 + 2\text{RNH}_2 \rightleftharpoons K_1 \text{RNH}_3^+ + \text{RNHCOO}^- \]  
\[ \text{CO}_2 + \text{RNH}_2 + \text{H}_2\text{O} \rightleftharpoons K_2 \text{H}_3\text{O}^+ + \text{RNHCOO}^- \]  
\[ \text{CO}_2 + \text{RNH}_2 + \text{OH}^- \rightleftharpoons K_3 \text{H}_2\text{O} + \text{RNHCOO}^- \]  
\[ \text{CO}_2 + \text{RNH}_2 + \text{CO}_3^{2-} \rightleftharpoons K_4 \text{HCO}_3^- + \text{RNHCOO}^- \]  

Water ionization reaction:

\[ 2\text{H}_2\text{O} \rightleftharpoons K_5 \text{H}_3\text{O}^+ + \text{OH}^- \]  

Dissociation reaction of dissolved CO₂:

\[ \text{CO}_2 + 2\text{H}_2\text{O} \rightleftharpoons K_6 \text{H}_3\text{O}^+ + \text{HCO}_3^- \]  

Dissociation reaction of bicarbonate:

\[ \text{HCO}_3^- + \text{H}_2\text{O} \rightleftharpoons K_7 \text{CO}_3^{2-} + \text{H}_3\text{O}^+ \]  

Dissociation reaction of RNH₃⁺:

\[ \text{RNH}_3^+ + \text{H}_2\text{O} \rightleftharpoons K_8 \text{RNH}_2 + \text{H}_3\text{O}^+ \]  

With the increase of CO₂ loading, the concentration of MEA in the solution gradually decreases, and the solution gradually becomes acidic. Meanwhile, the concentration of the protonated MEA gradually increases. Thus, the carbamate hydrolysis reaction, which was adopted by several research studies to form the equilibrium framework [27, 28, 33], is not involved to reflect the consumption of carbamate for relatively high CO₂ loading. Instead, equation (8) was used as a main modification to the model of Kent and Eisenberg [28]. A similar treatment was also adopted by Abouheir [33] for the consumption of the zwitterion.

Carbamate consumption reaction:

\[ \text{RNHCOO}^- + \text{H}_3\text{O}^+ \rightleftharpoons K_9 \text{RNH}_3^+ + \text{HCO}_3^- \]  

The above process consisting of some reactions allows us to understand the whole absorption process of CO₂ into the aqueous MEA solution.

2.3.2. Vapor-Liquid Equilibrium Model. In the present work, the vapor-liquid equilibrium is based on the physicochemical Kent-Eisenberg model [28]. Compared to the refined electrolyte NRTL model [12] and molecular simulations [29], the model assumes all activity coefficients and fugacity coefficient to be unity in the solutions is relatively simple and has good performance in predicting the CO₂ solubility [28, 35–38]. It was essential to adjust it to the thermodynamics of our systems, in terms of balance equations and equilibrium constants. The chemical equilibrium in the system is governed by equations (4)-(8). The pseudo-equilibrium constants representing the reactions in the CO₂/MEA/H₂O system are given by equations (9)-(13).

\[ K_5 = [\text{H}_3\text{O}^+][\text{OH}^-] \]  
\[ K_6 = [\text{H}_3\text{O}^+][\text{HCO}_3^-]/[\text{CO}_2] \]  
\[ K_7 = [\text{H}_3\text{O}^+][\text{CO}_3^{2-}]/[\text{HCO}_3^-] \]  
\[ K_8 = [\text{RNH}_3^+][\text{H}_3\text{O}^+]/[\text{RNHCOO}^-][\text{H}_3\text{O}^+] \]  

In addition to the above equilibrium equations, the mass balance equations of amine and carbon dioxide (equations (14) and (15)) and charge balance equation (16) must be satisfied.

\[ [\text{RNH}_2]_t = [\text{RNH}_2]_o + [\text{RNH}_3^+] + [\text{RNHCOO}^-] \]  
\[ [\text{CO}_2]_t = \alpha [\text{RNH}_2]_t \]
\[ = [\text{CO}_2] + [\text{HCO}_3^-] + [\text{CO}_3^{2-}] + [\text{RNHCOO}^-]. \]
\[
[ \text{RNH}_2^+ ] + [ H_2O^- ] = [ \text{OH}^- ] + [ \text{HCO}_3^- ] + 2[ \text{CO}_2^- ] + [\text{RNHCOO}^- ].
\] (16)

The physically dissolved CO\textsubscript{2} concentration is related to the CO\textsubscript{2} equilibrium partial pressure in the solvent by Henry’s law.

\[
[\text{CO}_2] = \frac{P_{\text{CO}_2}}{H_{\text{CO}_2}}.
\] (17)

The nine equations, equations (9)-(17), may be solved to obtain the value of CO\textsubscript{2} loading over MEA solutions of a given CO\textsubscript{2} partial pressure at a particular temperature. Equations (9)-(17) can be reduced, and expressions about the CO\textsubscript{2} loading, \( \alpha \), and [H\textsuperscript{+}] can be obtained consequently, shown in equations (18) and (19).

\[
\alpha = \left( \frac{P_{\text{CO}_2}/H_{\text{CO}_2}}{[\text{RNH}_2^+] + (K_6P_{\text{CO}_2}/H_{\text{CO}_2}[\text{H}^+] + (K_6P_{\text{CO}_2}/H_{\text{CO}_2}[\text{H}^+]^2 + ((K_8[\text{RNH}_2^+]P_{\text{CO}_2}/H_{\text{CO}_2})/(K_8[\text{RNH}_2^+]P_{\text{CO}_2}/H_{\text{CO}_2} + K_9[\text{H}^+]^2))} \right).
\] (18)

\[
[H^+] = \frac{K_5}{[H^+] + \frac{K_6P_{\text{CO}_2}}{H_{\text{CO}_2}[H^+] + \frac{2K_6K_7P_{\text{CO}_2}}{H_{\text{CO}_2}[H^+]^2} + \frac{K_6[\text{RNH}_2^+]P_{\text{CO}_2}/H_{\text{CO}_2} - K_9[\text{RNH}_2^+]^2}{K_8K_9[H^+] + K_8P_{\text{CO}_2}/H_{\text{CO}_2} + K_9[H^+]^2}}.
\] (19)

For the above equation, values of \( K_5, K_6 \), and \( K_7 \) used in the calculation are obtained from the published literature [33], summarized in Table 1. \( H_{\text{CO}_2} \) is the Henry’s coefficients, which can be calculated by the correlation from Tsai et al. [39, 40], summarized in Table 2. Therefore, there are three parameters (\( K_8, K_9 \), and [H\textsuperscript{+}]) remaining unsolved. By fitting the equations to the experimental values of CO\textsubscript{2} loading and partial pressure for each temperature condition using Levenberg–Marquardt algorithm, we could obtain the parameters of \( K_8 \) and \( K_9 \). During fitting, the range of [H\textsuperscript{+}] was limited between 10\textsuperscript{-6} and 10\textsuperscript{-12} mol/L [41]. After \( K_8 \) and \( K_9 \) were obtained, they were substituted into equations (9)-(17) to calculate the species equilibrium concentration, including [H\textsuperscript{+}].

### 3. Result and Discussion

#### 3.1. CO\textsubscript{2} Absorption Curve in MEA Absorbent

Figure 2 shows the absorption curves of CO\textsubscript{2} in an MEA aqueous solution at different temperatures. The absorption curve presents the ratios of CO\textsubscript{2} concentrations at the outlet gas \( C_{\text{out}} \) to the initial CO\textsubscript{2} concentration \( C_{\text{in}} \) as a function of time [42]. From the absorption curve at 298 K, it can be observed that \( C_{\text{out}}/C_{\text{in}} \) is close to zero in the first 35 minutes of the absorption process, and then increases rapidly thereafter. It can be speculated that most MEA has been consumed in the first 35 minutes. Due to the significantly lower MEA concentration, absorption rates decreased in the later phase. Absorption curves shift to the left as absorption temperature increases, indicating a faster saturation. Based on the absorption curve, the absorption load of MEA at different temperatures can be achieved through equation (1).

#### 3.2. CO\textsubscript{2} Solubility

The solubility of CO\textsubscript{2} in 20 wt.% aqueous MEA solutions was investigated to determine the dissociation equilibrium constants (\( K_a \)) of the RNH\textsuperscript{+} 3 and the carbamate consumption equilibrium constants (\( K_9 \)) at 298, 313, 333, and 353 K. The experimentally measured CO\textsubscript{2} loadings were plotted in Figure 3 for aqueous MEA solutions. The values of \( K_8 \) and \( K_9 \), which were obtained by using the modified Kent-Eisenberg model as discussed above and experimentally measured CO\textsubscript{2} loadings, are summarized in Table 3. According to the correlation equation of Table 1, \( K_8 \) and \( K_9 \) were modified to the modified Kent-Eisenberg model for various CO\textsubscript{2} loadings and temperatures. To facilitate comparison, experimental data from Wong et al. [25] and Lee et al. [43] are also included. It is evident that the new model is capable of correlating all the CO\textsubscript{2} solubility data well over the temperature range.
considered. Figure 4 illustrates the overall comparison of experimental equilibrium solubility data and values correlated from the modified Kent-Eisenberg model. As shown in the parity plot, all data points fall within 5% relative deviation, indicating a high degree of accuracy and reliability.

3.3. Chemical Species Distribution. The concentration distribution of liquid-phase species was analyzed with the modified Kent-Eisenberg model. Figure 5(a) shows a typical distribution profile at 313 K as a function of CO₂ loading, which is similar to the widely reported profile of CO₂ in primary amines. It is noteworthy that the equilibrium concentrations of CO₂⁻, OH⁻, and H₃O⁺ are very low and almost unchanged with the increase of loading from 0 to 1.0 mol CO₂/mol MEA. Thus, CO₂⁻, OH⁻, and H₃O⁺ were neglected in the whole absorption process, which is consistent with a previous study [24]. The modeled speciation results are compared with experimental results of NMR-spectroscopic measurements from Bottinger et al. [24] and Raman-spectroscopic measurements from Wong et al. [25]. Since protonated and unprotonated amines cannot be distinguished in NMR experiments, only HCO₃⁻ and RNHCOO⁻ from Bottinger et al. [24] are shown for comparison. As can be seen in Figure 5(a), model predictions are in a good agreement with literature NMR and Raman data at low loading levels, whereas slight deviations are observed at high loading levels, likely due to the increased experimental uncertainty at these levels [21]. Similar deviations are observed in the literature as well [24, 25]. In general, the modified Kent-Eisenberg model developed in this work generally gives better predictions than the previously reported Deshmukh–Mather model especially at high loadings [25, 44]. In general, the developed model displays a very good agreement with the experimental data obtained, either from NMR or Raman spectroscopy measurements, demonstrating its robustness.

For better understanding of the absorption process, the first-order difference curve of the equilibrium concentrations of all the species as a function of CO₂ loading with a difference step of 0.02 was further derived, as shown in Figure 5(b). Four regions are distinguished:
Figure 5: (a) Equilibrium speciation in MEA aqueous solution as a function of CO2 loading. (b) First-order difference curve of equilibrium speciation as a function of CO2 loading. Lines: data from modified Kent-Eisenberg model prediction; full symbols: experimental data of Wong et al. [25]; and empty symbols: experimental data of Bottinger et al. [24].

Table 4: Four regions of CO2 absorption in MEA with CO2 loading from 0 to 1 at 313 K.

<table>
<thead>
<tr>
<th>Region</th>
<th>CO2 loading (mol/mol)</th>
<th>Main reaction processes</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>0-0.32</td>
<td>(CO_2 + 2H_2O \rightleftharpoons K_K H_2O^+ + HCO_3^-) (3)</td>
</tr>
<tr>
<td>II*</td>
<td>0.32-0.5</td>
<td>(CO_2 + 2RNH_3+ \rightleftharpoons K_{K_1} RNH_3^+ + RNHCOO^-) (3)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(CO_2 + 2H_2O \rightleftharpoons Kg H_2O^+ + HCO_3^-) (5)</td>
</tr>
<tr>
<td>IIIb</td>
<td>0.5-0.66</td>
<td>(RNHCOO^- + H_2O \rightleftharpoons K_g RNH_3^+ + HCO_3^-) (8)</td>
</tr>
<tr>
<td>IV</td>
<td>0.66-1</td>
<td>(CO_2 + 2H_2O \rightleftharpoons K_g H_2O^+ + HCO_3^-) (5)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(RNHCOO^- + H_2O \rightleftharpoons K_g RNH_3^+ + HCO_3^-) (8)</td>
</tr>
</tbody>
</table>

*In region II, equation (3) is the dominant reaction. In region III, equations (5) and (8) are the dominant reactions.

(1) When CO2 loading is less than 0.32 mol CO2/mol MEA, CO2 dissolved in the aqueous MEA solution is all converted into carbamate. The ratio of the difference values of MEA, RNH+ 3, and RNHCOO– (−2 : 1 : 1) obeys the stoichiometric ratio of Reaction (3), and this reaction is the major process while other reactions can be neglected in the absorption process.

(2) For CO2 loading in the range of 0.32-0.5 mol CO2/mol MEA, the difference value of RNHCOO– concentration decreases significantly. It implies that Reaction (8) begins to take place and consumes the RNHCOO– formed by Reaction (3). As the difference value of RNHCOO– is still positive, Reaction (3) dominates over Reaction (8). In this region, CO2 hydrolysis (Reaction (5)) also occurs since it provides H2O+ for RNHCOO– consumption.

(3) For the CO2 loading between 0.5 and 0.66 mol CO2/mol MEA, the difference value of RNHCOO– begins to be negative, namely, the equilibrium concentration of RNHCOO– begins to decrease. Meanwhile, MEA is still present in the solution as shown in Figure 5(a) and RNHCOO– is formed continuously. Thus, in this region, Reactions (3), (5), and (8) all occur, but the latter two reactions are dominant. Due to that both Reactions (3) and (8) generate RNH3+, and the difference value of RNH3+ equilibrium concentration keeps almost unchanged in the first three regions from 0 to 0.66 mol CO2/mol MEA.

(4) For the loading from 0.66 to 1 mol CO2/mol MEA, only the consumption reaction of RNHCOO– (Reaction (8)) and CO2 hydrolysis reaction take place. MEA has been completely consumed and no RNHCOO– forms any more, leading to a minimum difference value of RNHCOO– at the CO2 loading of 0.66 CO2/mol MEA. Meanwhile, the difference value of CO2 concentration begins to increase significantly in this region, indicating a declined CO2 hydrolysis reaction. This then leads to declined RNHCOO– consumption, that is, decreased...
evolution rates (absolute difference values) of RNHCOO$^-$ and RNH$_3^+$ equilibrium concentration, as shown in Figure 5(b). This is also consistent with the decreased difference value of HCO$_3^-$ as the product of Reaction (8).

The four regions and main reactions are summarized in Table 4. In most of the previous studies, the process was simply divided into two regions at the point of 0.5 CO$_2$/mol MEA [24, 25, 29], which is the first time the process has been divided into four regions based on analyzing the difference curve of speciation. The first-order difference curve provides a better understanding over the equilibrium reactions. As can be seen from Table 4, the first region involves only one reaction, whereas the other three regions involve multiple reactions. Based on this, kinetic study of CO$_2$ absorption process should be conducted in the first region. In light of

Table 5: Four regions of CO$_2$ absorption in MEA with CO$_2$ loading from 0 to 1 at different temperatures.

<table>
<thead>
<tr>
<th>T/K</th>
<th>Region</th>
<th>I</th>
<th>II</th>
<th>III</th>
<th>IV</th>
</tr>
</thead>
<tbody>
<tr>
<td>298</td>
<td>0-0.34</td>
<td>0.34-0.5</td>
<td>0.5-0.66</td>
<td>0.66-1</td>
<td></td>
</tr>
<tr>
<td>313</td>
<td>0-0.32</td>
<td>0.32-0.5</td>
<td>0.5-0.66</td>
<td>0.66-1</td>
<td></td>
</tr>
<tr>
<td>333</td>
<td>0-0.26</td>
<td>0.26-0.5</td>
<td>0.5-0.66</td>
<td>0.66-1</td>
<td></td>
</tr>
<tr>
<td>353</td>
<td>0-0.2</td>
<td>0.2-0.5</td>
<td>0.5-0.66</td>
<td>0.66-1</td>
<td></td>
</tr>
</tbody>
</table>

Figure 6: First-order difference curve of equilibrium speciation as a function of CO$_2$ loading at different temperatures (a) T = 298 K; (b) T = 333 K; and (c) T = 353 K.
the fact that the bulk CO₂ equilibrium concentration increases very significantly in the fourth region, mass transfer measurement should be conducted in the first three regions rather than in the fourth.

The equilibrium species profile and the corresponding first-order difference curve of CO₂/MEA system at different temperatures of 298, 333, and 353K are also investigated, as shown in Figure 6. It can be observed that an increase in temperature does not significantly affect the trends described above, which is in consistent with the literature. Bottinger et al. [24] compared the results of the species concentrations at 293 and 333K, showing a very weak dependence with temperature. Balaji et al. [29] also found a similar phenomenon for results of the speciation at 333 and 353K. For all studied temperatures, CO₂ absorption process can also be divided into four regions at different temperatures. Whereas, the critical point between regions I and II changes notably with temperature. As shown in Table 5, the critical CO₂ loading between regions I and II becomes lower at higher temperatures. However, other critical points show no obvious change with temperature.

### 3.4. Effect of Temperature on Equilibrium Concentrations

Considering carbamate to be the most important species in CO₂ absorption process in amine solutions, the effect of temperature on its equilibrium concentration was investigated. Figure 7(a) shows the equilibrium concentration of RNHCOO⁻ as a function of CO₂ loading at different temperatures, and Figure 7(b) is the derived first-order difference curve. As can be seen, RNHCOO⁻ concentration curves in Figure 7(a) all present a parabolic relationship in the investigated temperature range. RNHCOO⁻ equilibrium concentration increases with CO₂ loading in regions I and II, whereas decreases in regions III and IV. The increasing rate of RNHCOO⁻ concentration in regions I and II shows a distinct decline with increasing temperature, as illustrated in Figure 7(b). The lower RNHCOO⁻ equilibrium concentration at higher temperatures with CO₂ loading lower than 0.5 CO₂/mol MEA in Figure 7(a) also illustrates this. As we discussed above, the dominant process in regions I and II is RNHCOO⁻ formation reaction (Reaction (3)), which is an exothermic process. According to Le Chatelier’s principle (the equilibrium law), increasing temperature causes equilibrium to shift to the left side of Reaction (3), namely, the lower increasing rate of RNHCOO⁻ concentration.

In contrast, RNHCOO⁻ consumption reaction (Reaction (8)) becomes dominant in regions III and IV, which is also an exothermic reaction because \( K_9 \) is negatively related with temperature as shown in Table 3. Therefore, the decreasing rates of RNHCOO⁻ concentration become lower at higher temperatures in regions III and IV, as shown in Figure 7(b). Finally, RNHCOO⁻ equilibrium concentration at higher temperatures exceeds that at lower temperatures at the point of CO₂ loading of 0.66 CO₂/mol MEA, as shown in Figure 7(a).

In addition, we also studied the effect of temperature on CO₂ bulk equilibrium concentration. As shown in Figure 8, the bulk CO₂ concentration is zero in regions I and II and gradually increases with CO₂ loading in regions III and IV. In the first two regions, CO₂ dissolved in MEA solution is all converted into carbamate by MEA. Whereas when CO₂ loading is higher than 0.5 CO₂/mol MEA (regions III and IV), MEA is completely consumed. The dissolved CO₂ can be converted into HCO₃⁻ by the hydrolysis reaction or exists in the form of molecular state. As CO₂ hydrolysis is an
exothermic reaction, it consequently results in more CO₂ existing in the form of molecular state at higher temperature, as illustrated in Figure 8. Furthermore, as the interfacial concentration of CO₂ (equilibrium to the gas phase) will decrease with the increase in temperature at fixed CO₂ partial pressure, the bulk CO₂ concentration (equals to the interfacial concentration at equilibrium) also decreases at higher temperature. From Figure 8, a higher bulk CO₂ concentration in the curve of higher temperatures corresponds to a lower CO₂ loading. Therefore, it can be concluded that CO₂ solubility will decrease with increasing temperature, which is in agreement with the experimental results presented in Section 3.2.

4. Conclusion

This work has systematically investigated CO₂ absorption equilibrium in MEA aqueous solution by experimental measurement combined with thermodynamic model. The data of CO₂ solubility in MEA were measured at 298, 313, 333, and 353 K over a CO₂ partial pressure range of 34.5 to 78.0 kPa. C_{out}/C_{in} is close to zero in the first 35 minutes of the absorption process at 298 K, and then increases rapidly. The absorption curve moves leftwards with increasing temperature, indicating a faster saturation. A modified Kent-Eisenberg model was developed using the measured solubility data based on the Kent-Eisenberg model framework. The obtained thermodynamic model shows a good prediction of the species distribution of 20 wt.% MEA system under different CO₂ loadings.

In the first attempts to analyze the absorption process, a difference curve of the equilibrium concentrations of liquid-phase species as a function of CO₂ loading has been used. According to the change of reaction processes, CO₂ absorption process has been divided into four regions with CO₂ loadings ranging from 0 to 1. CO₂ dissolved in the aqueous MEA solution is all converted into carbamate in the first region. The difference value of RNHCOO⁻ concentration decreases significantly in the second region. The difference value of RNHCOO⁻ begins to be negative, namely, the equilibrium concentration of RNHCOO⁻ begins to decrease in the third region. Only the consumption reaction of RNHCOO⁻ and CO₂ hydrolysis reaction take place in the fourth region. Furthermore, based on the characteristics of each region, we propose that further kinetic study for measuring reaction rates and kinetic constants should be conducted in the first region, whereas measuring of mass transfer rates could be conducted in the first three regions. The innovative division of CO₂ absorption into four regions contributes to a deeper understanding of the process, as well as providing guidance for further kinetic studies. In addition, this novel method can be further extended to more CO₂ absorption processes utilizing different absorbents at various conditions. The CO₂ absorption processes using microreactors can also be easily applied to a larger-scale reaction by simply numbering-up.

Nomenclature

$\alpha$: CO₂ loading;
$C$: Concentration, mol/m³;
$H$: Henry’s law constant, kPa/(kmol/m³);
$K$: Equilibrium constant;
$n$: Mole number, mol;
p: Pressure, Pa;
$Q$: Gas flow, m³/s;
$R$: Gas constant, J/(K mol);
t: Time, s;
$T$: Temperature, K;
in: Inlet;
abs: Absorption;
exp: Experimental;
cal: Calculative;
gas: Gas phase;
$L$: Liquid phase;
t: Total.

Data Availability

All data used to support the findings of this study are currently under embargo while the research findings are commercialized. Requests for data, 12 months after publication of this article, will be considered by the corresponding author.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Acknowledgments

The authors gratefully acknowledge the financial supports for this project from the China State Construction Engineering Corp. (CSCEC-2021-Z-51 and CSCEC-PT-007-02), the China Construction Industrial & Energy Engineering Group Co. Ltd. (AZ-2020-02), Nanjing Municipal Commission of Urban-Rural Development (Ks2253), General Project of Natural Science Research in Colleges and Universities of Jiangsu Province (No. 21KJB530009), and Opening Fund of National & Local Joint Engineering Research Center for Deep Utilization Technology of Rock-Salt Resource (Grant no. SF202007).
References


