

1 Aqueous MAPA, DEEA and their blend as CO₂
2 absorbents: Interrelationship between NMR
3 speciation, pH and heat of absorption data

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11

12 **ABSTRACT**

13 Based on NMR speciation, heat of absorption and pH data, the reactions occurring in aqueous
14 MAPA (3-(Methylamino)propylamine), DEEA (2-(Diethylamino)ethanol) and their blend at
15 various carbon dioxide (CO₂) loadings were identified and discussed.

16 At increasing CO₂ loading, the basicity of the solutions decreased. In the MAPA solvent, this
17 led to the hydrolysis of the carbamate species, which corresponded to a drop in the heat of
18 absorption. In the blend, due to the activity of DEEA, such a drop in both pH and heat of
19 absorption was not noticeable. In the presence of MAPA, in the early loadings DEEA had not

1 yet an influence on the (bi)carbonate formation, but mainly on the MAPA activity. However,
2 as soon as the trend for (bi)carbonate formation went as in DEEA singly, the released heat
3 steadily decreased. Both in single and blended MAPA, the carbamate hydrolyses started at
4 similar pH values, i.e. about 9.8-9.5.

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1. INTRODUCTION

1.1. Background

Carbon dioxide (CO₂) from human activities has been recognized as the greenhouse gas making the largest contribution to the global warming, and actions to reduce its concentration in the atmosphere are solicited.¹ Amine scrubbing is one of the most common technology for removal of CO₂ from flue, natural and other exhaust gases of different industrial sources.² The benchmark absorbent is monoethanolamine (MEA, a primary amine) in an aqueous mean, which is known for its high reactivity and favorable reaction kinetics toward CO₂, but it suffers from degradation and unfavorable thermodynamics.³ Replacing MEA is a challenging task because the selection of any new absorbent should take into account several key criteria, like e.g. chemical and physical properties of the solvents, kinetics and thermodynamics of chemical reactions, environmental impacts, costs and so on.⁴

To reach ideal solvent properties, aqueous blended amines are receiving great attention, as the best features of two or more amines can be combined and/or even improved.⁵ The mixture composed of MAPA, 3-(Methylamino)propylamine, and DEEA, 2-(Diethylamino)ethanol, is a promising CO₂ absorbent. MAPA is a diamine, composed of a primary and a secondary amino functional group, which shows favorable kinetics of reaction but high heat of absorption, while for DEEA, a tertiary amine, the opposite is true.⁶ Several key properties, like e.g. heat of absorption, kinetics and volatility, of MAPA-DEEA blends both as single phase and/or as demixing solvent have been studied.⁷⁻⁹ The reaction mechanisms of MAPA and DEEA, singly and in blend, is also under investigation as it would allow the optimization of the existing solvents and the design of new ones with improved properties. Recently, Zhang et al.¹⁰ performed an NMR speciation analysis for CO₂ loaded aqueous MAPA and MEDA (*N*-Methylethylenediamine), two diamines with different chain lengths, to understand the competitive reactions between the intramolecular amino functional groups. Based on the

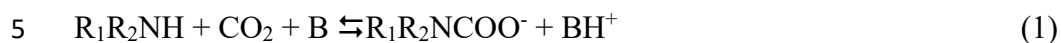
1 amount of each species as well as on the pH values at various addition of CO₂, three different
2 stages of reactions were identified for each amine solvent. However, in that study, MAPA was
3 investigated singly only. In 2018, Perinu et al.¹¹ performed by means of NMR spectroscopy a
4 chemical speciation analysis on MAPA, eight tertiary amines (including DEEA) and their
5 blends after CO₂ absorption (40 °C) and desorption (80 °C) tests in a screening apparatus. For
6 each amine system, the speciation analyses were performed on a unique sample withdrawn
7 after the absorption screening experiments, which terminated when the effluent reached 9.5
8 kPa partial pressure of CO₂.¹² Highlights on the role played by the tertiary amines in the
9 presence of MAPA were reported and it was hypothesized that the tertiary amines, which act
10 as Brønsted bases, contribute to reduce the protonation of MAPA species in favor of MAPA
11 carbamate formation. A list of the probable reactions was included in the study, but it was
12 possible neither to confirm them nor to define at which stage of the CO₂ addition they were
13 occurring.

14 During the absorption of CO₂ by an amine system in an aqueous mean, several acid-base
15 reactions are involved, and the species found at the equilibrium depend on the chemical
16 reactions occurred, which in turn depend on the chemical properties of the amines and reaction
17 conditions. Many factors may indeed influence the CO₂ absorption capacity of a solvent, like
18 e.g. type of amines and chemical structure, concentrations, basicity (pH of the solution and pK_a
19 of the amines), reaction energy and so on, and when selecting a solvent, the different features
20 should be taken into consideration.¹³ Moreover, when two or more amines are in mixture, the
21 CO₂ loaded amine systems become more complicated as each amine group react differently
22 and the ratio between the amines has an impact too.

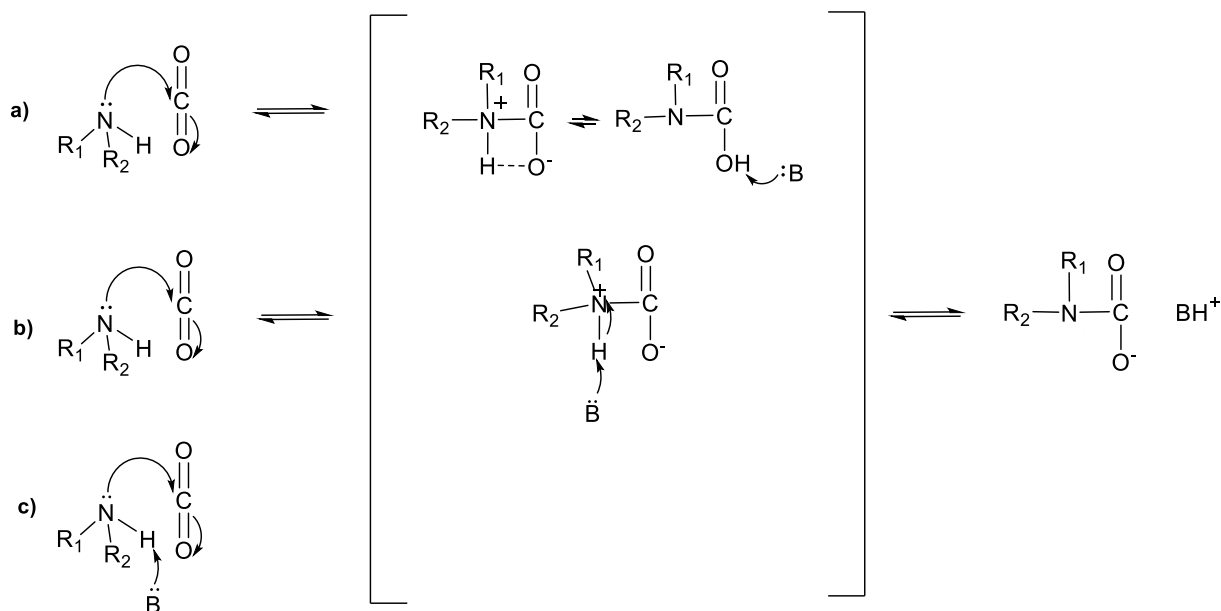
23 The primary reactions between amines and CO₂ in water are mainly known and largely
24 discussed in literature. As general, in an aqueous amine system in the presence of CO₂, amine

1 carbamate ($R_1R_2NCOO^-$) and bicarbonate/carbonate (HCO_3^-/CO_3^{2-}) are the two main reaction
 2 products, which strongly affects the total amount of CO_2 absorbed and released.

3 Amine carbamates are usually formed by the reaction of primary and/or secondary amines with
 4 CO_2 in the presence of a base (Reaction 1).



6 The base (B) can be an amine and/or water. The carbamate forming reaction is the reaction of
 7 the lone pair of the amine nitrogen (which is a Lewis base) with the electrophilic centre (Lewis
 8 acid) of CO_2 .¹⁴ Several mechanisms have been proposed in literature (Scheme 1) and, due to
 9 the fast kinetics and the wide range of amine conditions (such as type of amine, concentrations,
 10 pK_a), they are still under debate.^{15, 16} One possibility is that the amine carbamate is formed
 11 through an intermediate, either carbamic acid (Scheme 1a)¹⁶ or zwitterion (Scheme 1b)¹⁷, or
 12 through a single step (Scheme 1c)¹⁸, that is a termolecular reaction between CO_2 , the amine
 13 and a base (B).



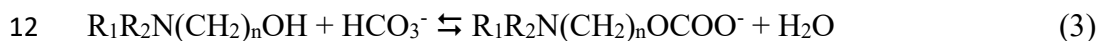
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16 **Scheme 1.** The main reactions mechanisms available in literature for amine carbamate
 17 formation. B represents a Brønsted base.

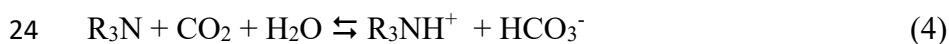
1 Secondary amines form amine carbamates less stable than those obtained by primary amines.
2 The carbamate hydrolysis constants of secondary amines are at least one order of magnitude
3 larger than those of corresponding primary amines.¹⁹ This lower stability may be because the
4 nitrogen (N-) atom is more hindered than in a primary amino group.

5 Tertiary amines cannot react with CO₂ directly, although a limited number of authors have
6 observed the formation of alkyl carbonate in aqueous tertiary alkanolamines.^{11, 20-23} Jørgensen
7 and Faurholt²⁰ suggested the reaction (2) in basic aqueous solutions of amino alcohols, and
8 proposed the overall reaction (3) (which represents the inverse of the alkyl carbonate
9 hydrolysis) as they observed the formation of the alkyl carbonate when mixing amines,
10 bicarbonate and carbonate.



13 At pH lower than 12, the rate of its formation is negligible and doesn't lead to relevant amount
14 at equilibrium. However, the amount of this species is documented to increase at increasing
15 CO₂ loading (decreasing pH), as reported in Behrens et al.²¹ in a study specific on
16 monolakylcarbonate formation in aqueous MDEA (Methyldiethanolamine). In that study, it
17 was concluded that, though formed in relatively small amount and very little knowledge are
18 available on it, those species must be taken into consideration in the models describing the
19 liquid phase reactions. In most cases, this species was indeed neglected, and in some NMR
20 studies the corresponding peaks were attributed to undefined side-products and/or impurities.

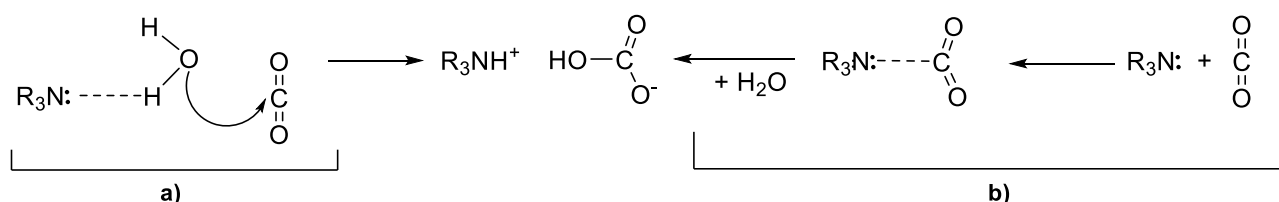
21 A primary reaction between CO₂ and tertiary amines in aqueous solutions is instead a kind of
22 base catalysis of the CO₂ hydration leading to the formation of bicarbonate (reaction 4), through
23 an acid-base buffer mechanism.



1 It is known that, in an aqueous mean, CO₂ can react with OH⁻ (reaction 5), followed by the
 2 reaction 6, and H₂O (reaction 7).²⁴ Reaction 7 is negligible at pH higher than 10, whereas at
 3 pH between 10 and 8 both reaction pathways (5 and 7) are possible.^{25, 26} It is observed that the
 4 loss of the proton from the H₂O molecules can cause an enormous increase in the affinity for
 5 CO₂²⁷ and, then, the reaction 7 can be catalysed by chemical constituents able to get hydrogen
 6 ions. On this regard, tertiary amines are good Brønsted bases, which by definition are
 7 substances able to accept a proton (reaction 8).^{24, 28} The pair of unshared electrons on the
 8 nitrogen indeed enables amines to act as general base catalysts.



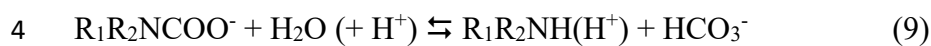
13 In literature, for (bi)carbonate formation in the presence of a tertiary amine, several
 14 mechanisms have been proposed,²⁸ and for all of them the overall pathway is represented by
 15 reaction 4 (overall reaction, 5+8).^{22, 28, 29} A possible chemical path for reaction (4) is assumed
 16 to be a termolecular reaction (scheme 2a) where a hydrogen bonding between the free amine
 17 and water increases the nucleophilic reactivity of the water towards CO₂.^{30, 31} Depending on
 18 the basicity of the amines, a two-step cascade reaction is also possible, that is reaction 8
 19 followed by reaction 5.³² Moreover, it was also suggested the formation of an intermediate in
 20 the catalytic path, as shown in the Scheme 2b.²⁸



21

22 **Scheme 2.** Possible reaction mechanisms for CO₂ hydration catalyzed by a tertiary amine.

1 In addition, bicarbonate is also formed through the hydrolysis of the amine carbamate. Due to
2 steric hindrance or weakly alkalinity, the amine carbamate can become unstable, leading to the
3 formation of HCO_3^- , as in reaction 9.^{33, 34}



6 **1.2. Aim of the work**

7 An aqueous amine system in the presence of water and CO_2 leads to the occurrence of several
8 parallel and/or consecutive reversible reactions. Knowing the structure and the amount of the
9 reaction products is fundamental, as well as the measurement of the pH may be valuable to
10 understand the dynamics leading to specific reaction pathways and the position of the acid-
11 base reaction equilibria. By measuring pH, the status of equilibrium of all acid-base containing
12 reactions is indeed measured. Moreover, the heat released during the addition of CO_2 in an
13 amine solution is directly related to the steam consumption and gives information on how
14 sensitive the system is when subject to a change in temperature.³⁵ The heat of absorption is
15 given by the combination of the heat of physical dissolution of CO_2 into the amine solvent and
16 the heat of reaction between CO_2 and the amine solvent. The formation of carbamate is an
17 exothermic reaction, whose released heat is higher than that for the formation of bicarbonate.⁷
18 The present study aims to understand the reaction mechanisms of the absorption of CO_2 in
19 aqueous MAPA and DEEA, singly and in mixture, and the factors influencing them. Speciation
20 analyses, as given by NMR (Nuclear Magnetic Resonance) spectroscopy, pH and heat of
21 absorption measurements were performed at various addition of CO_2 in the amine solvents.
22 The heat of absorption and pH measurements were combined to the NMR speciation data to be
23 able to explain them in a more fundamental level. Insights on the chemical reactions are given,
24 and reactions mechanisms are discussed.

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1 **2. EXPERIMENTAL SECTION**

2 **2.1 Chemicals**

3 The chemicals 3-(Methylamino)propylamine ($\geq 97\%$, CAS: 6291-84-5) and 2-
4 (Diethylamino)ethanol ($\geq 99.5\%$, CAS: 100-37-8) were purchased from Sigma-Aldrich, and
5 Carbon Dioxide (99.999%) was purchased from AGA. The chemicals were used without
6 further purification.

7

8 **2.2 Sample preparation and density**

9 Aqueous MAPA 1M, DEEA 3M and MAPA1M-DEEA3M were prepared in a volumetric
10 flask: the required amounts of amines were weighed on a Mettler Toledo balance (model
11 MS6002S, $d = 0.01$ g), and distilled water was added up to a predetermined volume and
12 weighed. The density of the unloaded solutions was measured at 25 °C using an Anton Paar
13 DMA 4500 M densitometer. The nominal repeatability, as given by the manufacturer, was 0.01
14 kg/m^3 and 0.01 °C. A detailed description of the experimental procedure is given by Pinto et
15 al.³⁶ The density of the loaded solutions was measured, as reported in Perinu et al. 2018,¹¹ by
16 weighing on a Mettler Toledo analytical balance (model ME204, $d = 0.0001$ g) 1 mL of sample
17 (as measured by Hamilton syringes, model 1001 TLL) at room temperature. The uncertainty
18 of this procedure was determined to be $<0.6\%$. In Tables S1 and S2 of the Supporting
19 Information, the densities of unloaded and loaded solutions are reported, respectively.

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21 **2.3 CO₂ loading**

22 CO₂ loaded MAPA 1M, DEEA 3M and DEEA 3M-MAPA 1M solutions were prepared by
23 bubbling pure CO₂ gas into each amine solution at room temperature. To prevent an excessive
24 overheating of the solution, and thus water and amine loss, CO₂ was slowly added into the
25 solutions. Furthermore, the concentration of the amine solutions before and after CO₂ bubbling

1 was measured and the error percentage was lower than 1.5 %, i.e. within the analytical error.
2 The highly CO₂-loaded amine solutions were mixed with respective unloaded amine solution
3 to obtain the predetermined CO₂ loading. To determine the CO₂ loading, the amine and CO₂
4 concentrations were analyzed using the titration and the precipitation methods, described in
5 Ma'mun et al.³⁷

7 **2.4 NMR experiments**

8 NMR is a powerful non-invasive analytical technique for chemical analyses and allows direct
9 access to the chemical composition of CO₂ capture solvents (speciation). Chemical structures
10 of the molecules, including unknown products and/or side-products, can be defined, and in
11 proper performed NMR experiments the species can also be quantified.³⁸

12 A liquid sample of each solution at different loadings was taken for NMR analysis. NMR
13 experiments were performed at 26.85 °C on a Bruker 600 MHz Avance III HD equipped with
14 a 5-mm cryogenic CP-TCI z-gradient probe. Qualitative 1D and 2D NMR experiments were
15 performed to identify the species in solution, whereas quantitative ¹³C NMR spectroscopy was
16 applied to measure the corresponding concentration. The ¹³C NMR spectra were acquired with
17 the inverse gated decoupling acquisition sequence, using a recycle delay time of 120 s, a pulse
18 width of 11.4 μs (90° pulse angle) and 256 scans. Deuterated water was used as “lock” solvent
19 and was inside a coaxial insert, whereas Acetonitrile was chosen as internal reference standard.
20 Further details on the method are available in previous studies.^{11,39}

22 **2.5 pH measurements**

23 The pH of each unloaded and loaded sample was measured using an InLab NMR pH electrode
24 by Mettler Toledo connected to a pH meter (SevenEasy by Mettler Toledo). The calibration of
25 the pH was performed using technical pH buffer solutions at pH 7.00, 9.21 and 11.00 provided

1 by Mettler Toledo. Each solution was inserted in an NMR tube and immersed in a thermostatic
2 water-bath (Lauda, Ecoline E100) at $T = 25.0 \pm 0.1$ °C. Before and after the measurements of
3 the CO₂-loaded amine solutions, the pH of the standard solutions was also measured as they
4 were samples, giving a pH variation of ± 0.01 . In Table S3 of the Supporting Information, the
5 pH values of the solutions are reported.

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7 **2.6 Calorimeter**

8 Differential heats of absorption of CO₂ in MAPA 1M, DEEA 3M and DEEA 3M-MAPA 1M
9 aqueous solutions were measured at 25 °C in a reaction calorimeter CPA-122 (Chemisens). To
10 validate the data, heat of absorption of CO₂ at 40 °C for MAPA 1M and DEEA 3M was
11 measured and compared to available literature data. The calorimeter is a stainless steel
12 mechanically agitated reactor with a volume of 2000 cm³. The reactor was operated under
13 isothermal conditions and flow of heat, pressure, temperature and other parameters were
14 measured as a function of time. CO₂ was fed from two storage cylinders through a mass flow
15 controller. A detailed description of the experimental procedure is given in Kim and Svendsen⁴⁰
16 with a small modification given in Kim et al.⁴¹

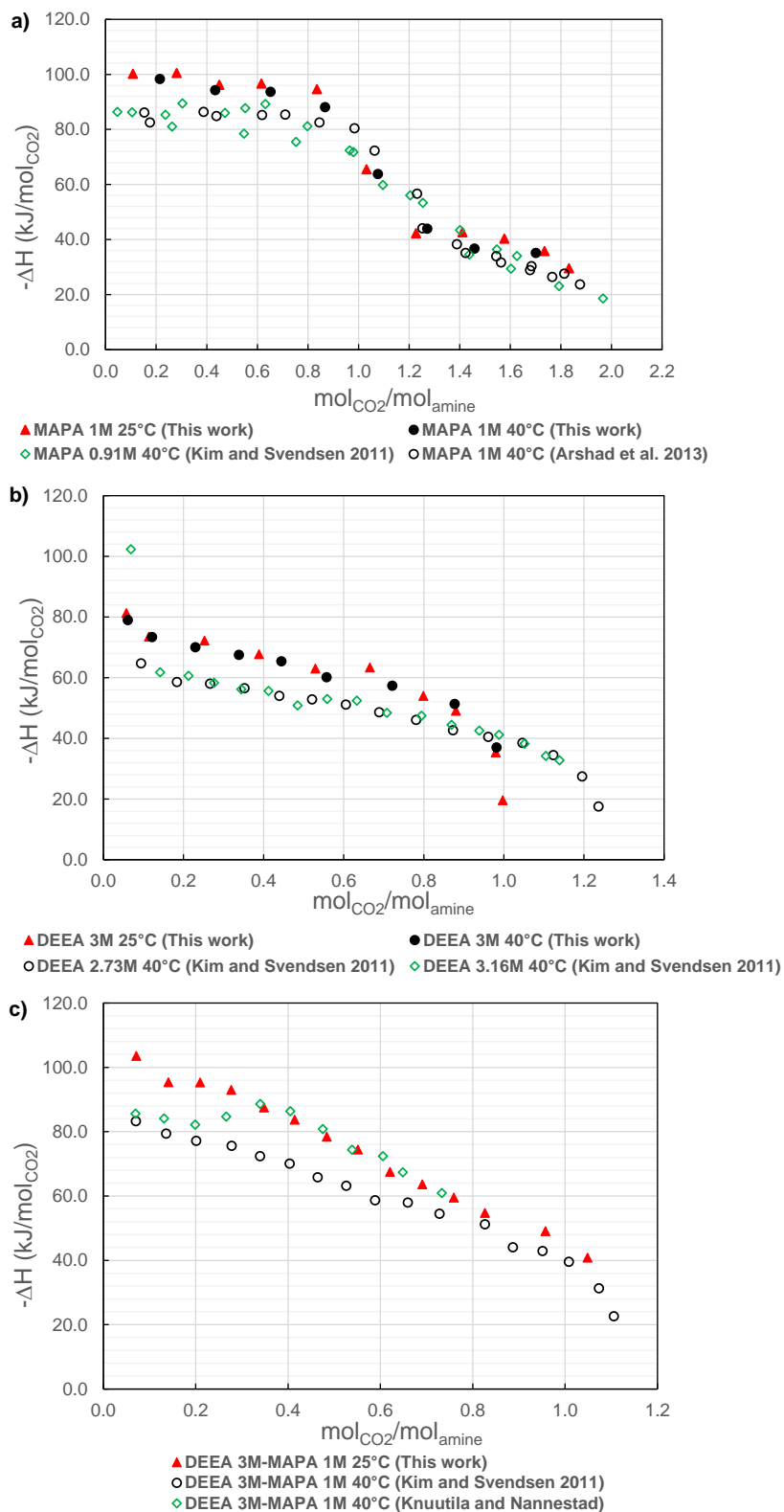
17 The experiments were initiated by evacuating the reactor to ~ 0.04 bar. Subsequently, around
18 1.1 - 1.2L of amine solution was fed to the reactor, and the reactor was once again evacuated
19 to degas the solution. When the system reached equilibrium, under stirring at the experimental
20 temperature, CO₂ was added to the reactor through a bottom valve. The system was left to reach
21 equilibrium before a new batch of CO₂ was fed into the solution. Equilibrium was considered
22 to be reached when there was no change in heat flow, temperature and pressure. This procedure
23 was repeated until the pressure in the reactor reached around 6 bar. At the end of the
24 experiment, a liquid sample was taken for CO₂ analysis using the precipitation and titration
25 method.³⁷

1 For each injection of CO₂, equilibrium CO₂ loading (mol_{CO2}/mol_{amine}) was calculated by
2 subtracting moles of CO₂ in the gas phase from moles of CO₂ fed into the reactor, and then
3 divided by moles of amine. Moles of CO₂ fed from the CO₂ vessel and present in the gas phase
4 were calculated by Peng-Robinson equation of state, using information about pressure
5 difference and temperature in the cylinder and the reactor, respectively. The last CO₂ loading
6 deviated in average 2.4% from the CO₂ analysis. Heat of absorption was calculated by
7 integrating the heat flow curves and it was corrected for the compression-work and
8 cooling/heating effect of the incoming CO₂ gas.

9 **3. RESULTS AND DISCUSSION**

10 **3.1 Heat of absorption: comparison to literature data**

11 The differential heat of absorption of CO₂ (-ΔH, kJ/mol_{CO2}) in MAPA 1M, DEEA 3M and
12 DEEA 3M-MAPA 1M solutions, measured in this work (Tables S4-S8 of the Supporting
13 Information) and in the literature, are presented in Figure 1. The results show no significant
14 difference in the heat of absorption measured at 25 and 40 °C. Overall, for MAPA and DEEA,
15 the measured heat of absorption at 40 °C as a function of CO₂ loading follows the same
16 behavior as reported in the literature.^{6, 7, 40, 42} For MAPA 1M, the heat of absorption was slightly
17 higher at low CO₂ loadings, but similar after a loading of 0.6 mol_{CO2}/mol_{amine}. For DEEA, the
18 measured values were slightly higher throughout the CO₂ loading range than reported in Kim
19 and Svendsen⁴³ and Kim.⁴² Nevertheless, the heat of absorption at 40 °C reported by Kim and
20 Svendsen⁴³ were also lower than those reported by Knuutila and Nannestad⁶ for the blended
21 DEEA-MAPA system measured at the same temperature.



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2

3 **Figure 1.** Differential heat of absorption of CO₂ in a) MAPA, b) DEEA and c) DEEA-MAPA

4 solutions.

3.2 Chemistry of the system and NMR analysis

In the current study, the species identified and quantified at the equilibrium in the amine systems under investigation at the different CO₂ loadings are represented in Figure 2. They are in line with the typical reaction products expected by a primary/secondary diamine and a tertiary amine in an aqueous medium in the presence of CO₂.^{11, 44, 45}

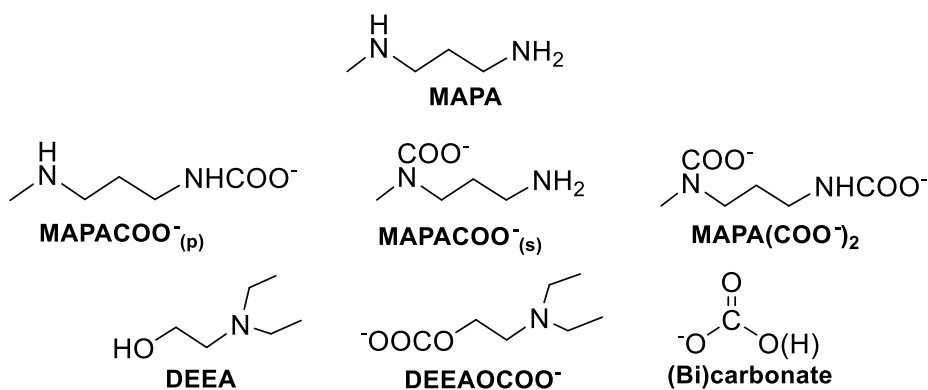
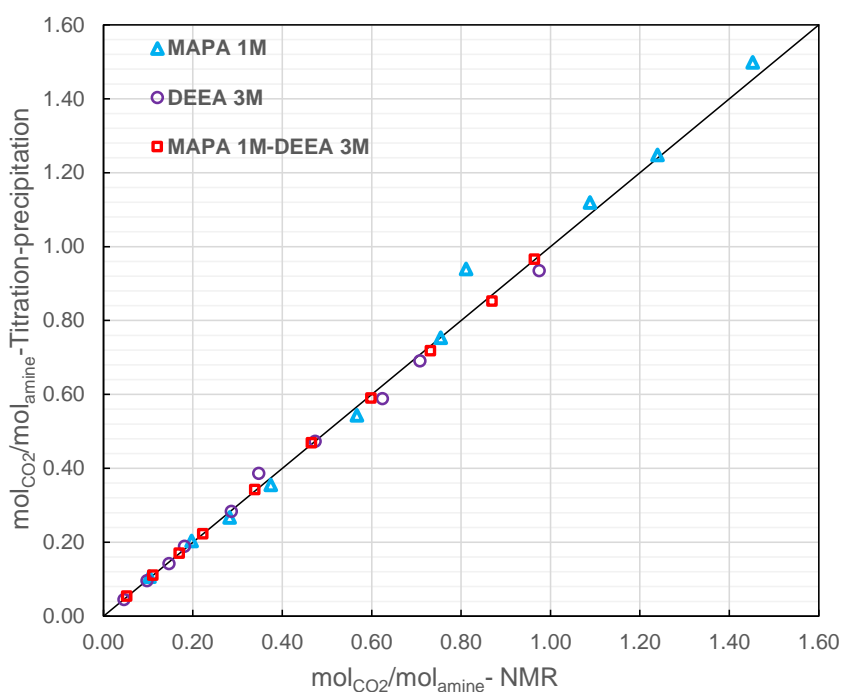


Figure 2. Chemical structure of MAPA, DEEA, their derivatives and (bi)carbonate in the aqueous solutions in the presence of CO₂. Each species is in equilibrium with the corresponding protonated form.

MAPA/MAPAH⁺/MAPA(H⁺)₂ (simply reported as MAPA/MAPA(H⁺)₂ along the manuscript and in the graphs), primary MAPA carbamate (MAPACOO⁻(p)/MAPAH⁺COO⁻(p)), secondary MAPA carbamate (MAPACOO⁻(s)/MAPAH⁺COO⁻(s)) and MAPA(COO⁻)₂ are the species identified in the amine systems containing MAPA; DEEA/DEEAH⁺ and tertiary amino carbonate (DEEAOCOO⁻/DEEAH⁺OCOO⁻) are identified in single and blended DEEA; and (bi)carbonate species (HCO₃⁻/CO₃²⁻) is found in all the systems. Details on the corresponding concentrations (mol/L) can be found in the Supporting Information (Tables S9-S11).

In the ¹³C NMR spectra, the signals corresponding to carbons of the protonated and neutral form of each species (free/protonated amines and/or HCO₃⁻/CO₃²⁻) appear with a common signal because, at the selected temperature, the proton exchange is faster than the NMR time

1 scale. The chemical shifts (expressed in ppm) depend on their ratio and, in the case of HCO_3^-
2 and CO_3^{2-} , they can span from about 161.1 ppm for 100% HCO_3^- to about 168.7 ppm for 100%
3 CO_3^{2-} .³⁹ In this study, an estimation of the HCO_3^- and CO_3^{2-} content was performed by applying
4 the method developed by Perinu et al.³⁹
5 To verify the reliability of the NMR data, the CO_2 loadings as measured by NMR and by
6 titration-precipitation method were compared (Figure 3). The measurements obtained by both
7 methods were fitting quite well, except for one point in a loaded solution of MAPA 1M.
8 Specifically, at the 7th CO_2 loading ($0.94 \text{ mol}_{\text{CO}_2}/\text{mol}_{\text{amine}}$), as measured by using the titration-
9 precipitation method, and at $0.81 \text{ mol}_{\text{CO}_2}/\text{mol}_{\text{amine}}$, as measured by NMR analysis, there is a
10 mismatch. The reason may be related to the fact that the peak corresponding to the HCO_3^-
11 $/\text{CO}_3^{2-}$ species was not identified in the NMR spectra, although this species might be present.
12 It may be possible that the peak corresponding to the (bi)carbonate species is overlapped with
13 the carbon-peak of the carboxylate ion of a MAPA carbamate species.



14
15 **Figure 3.** Comparison of $\text{mol}_{\text{CO}_2}/\text{mol}_{\text{amine}}$ as measured by NMR and by titration-precipitation
16 analyses.

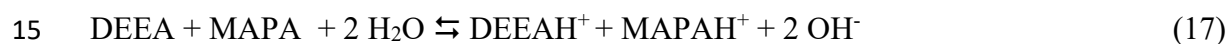
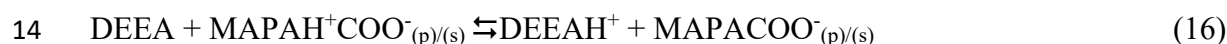
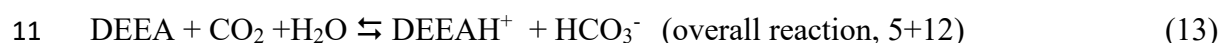
1 The identification and quantification of the species and the interrelation with pH and heat of
 2 absorption data allowed defining a list of reactions occurring in the solvents under
 3 investigation. For sake of clarity, all the reactions discussed in each section are here listed, but
 4 they are not in the order they may occur in the different systems:

5 **CO₂-H₂O reactions**

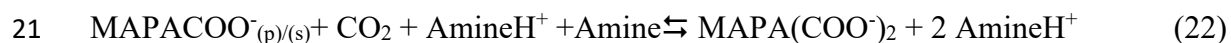
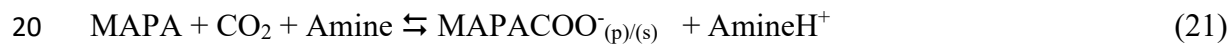
6 Reactions (5), (6), (7) (see section 1.1)



9 **DEEA reactions**



16 **MAPA reactions**



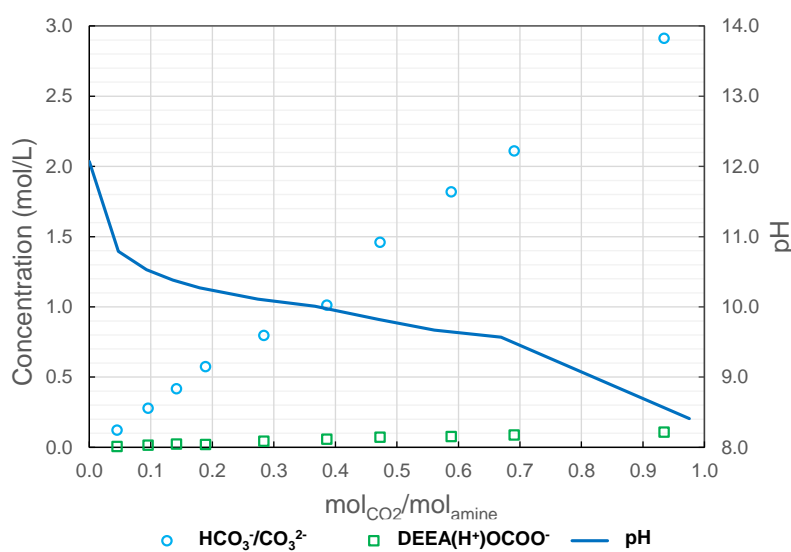
1 Reaction 10 is neither discussed nor mentioned during the discussion of the results, as it is
2 considered to be a default reaction. In the reactions 21 and 22, the “Amine” acting as counter-
3 ion (stronger Brønsted base) may change as a function of the solvents investigated. In reaction
4 20, the nitrogen of $\text{MAPAH}^+\text{COO}^-_{(p)/(s)}$ not bound to CO_2 is protonated and acts as counterion
5 of the $-\text{COO}^-$ group.

6

7 3.3 Analysis of NMR speciation, pH and ΔH data

8 3.3.1 DEEA 3M- CO_2 - H_2O system

9 The concentration (mol/L) of all the species formed in aqueous DEEA 3M, in the presence of
10 different addition of CO_2 , are plotted together with pH data (Figure 4).



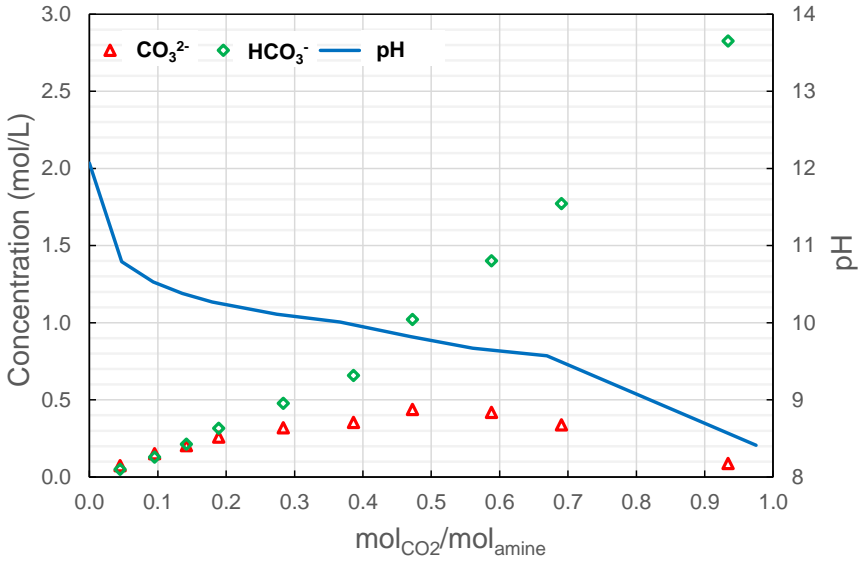
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12 **Figure 4.** NMR speciation and pH data in aqueous DEEA 3 M at increasing CO_2 loading (25
13 $^{\circ}\text{C}$).

14

15 (Bi)carbonate ($\text{HCO}_3^-/\text{CO}_3^{2-}$) and DEEA OCCO^- are the two CO_2 -containing species present
16 at the equilibrium at the different loadings.

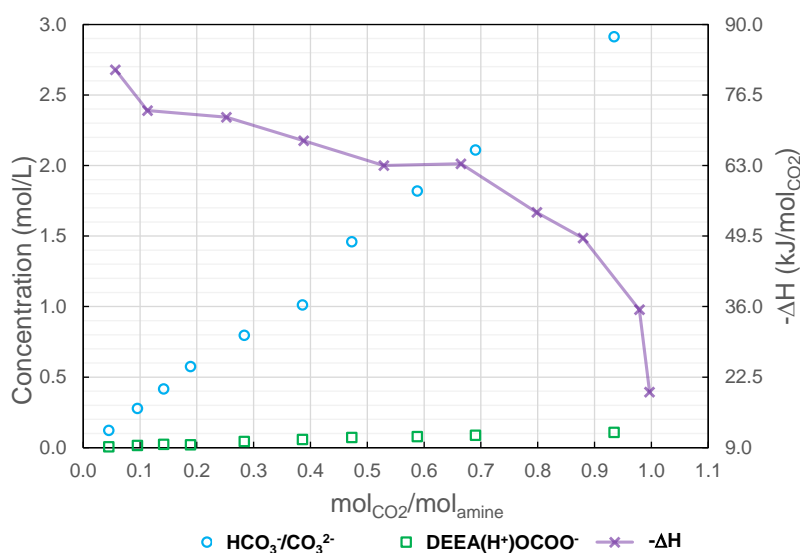
1 DEEAOCOO⁻ is found in negligible amount and it appears to have the tendency of increasing
 2 at increasing addition of CO₂, as for MDEAOCOO⁻ in the study on CO₂ loaded MDEA in
 3 water by Behrens et al.²¹
 4 As expected, (bi)carbonate species are the main product of the CO₂ absorption and the total
 5 concentration of HCO₃⁻ and CO₃²⁻ constantly increases since the beginning of the loading. In
 6 this system, from 0.04 molCO₂/mol_{amine} to about 0.59-0.69 molCO₂/mol_{amine}, the chemical
 7 shifts were ranging from 165.7 ppm to 162.8-162.3 ppm, respectively, and so were consistent
 8 with the presence of both HCO₃⁻ and CO₃²⁻, with the HCO₃⁻ content increasing at increasing
 9 loading. At the maximum loading, the chemical shift value (161.3 ppm) is instead consistent
 10 with the predominant presence of HCO₃⁻ (the chemical shifts corresponding to HCO₃⁻/CO₃²⁻
 11 signal are available in the Table S12 of the Supporting Information). In terms of concentrations,
 12 the Figure 5 shows that, at increasing loading, the amount of HCO₃⁻ constantly increases,
 13 whereas the amount of CO₃²⁻, mainly in concentrations lower than HCO₃⁻, increases until a
 14 loading of around 0.47 molCO₂/mol_{amine} before it starts being consumed.



15
 16 **Figure 5.** Estimation of the HCO₃⁻ and CO₃²⁻ content in aqueous DEEA 3M.

17 The pH values span from pH 12.07 at 0.00 molCO₂/mol_{amine} to pH 8.41 at maximum loading,
 18 and it shows a constant decrease, with some changes in the slope of the curve. We observe a

1 drop of $\Delta\text{pH}=1.28$ in the range between the $0.0 \text{ molCO}_2/\text{mol}_{\text{amine}}$ and $0.05 \text{ molCO}_2/\text{mol}_{\text{amine}}$,
 2 due to the first addition of the acid CO_2 , and another of $\Delta\text{pH}=1.16$ in the last loadings-region,
 3 between 0.67 and $0.98 \text{ molCO}_2/\text{mol}_{\text{amine}}$, where the basicity of the solution greatly decreases as
 4 the majority of the amines are protonated. DEEA/DEEAH⁺ is a buffer system made of a weak
 5 acid (DEEAH⁺) and its conjugate base (DEEA). When the pH of the solution is lower than the
 6 pK_a value of an amine, the larger part of the amine is protonated and cannot act as a catalyst of
 7 CO_2 hydration.³⁰ In this study, at about $0.47 \text{ molCO}_2/\text{mol}_{\text{amine}}$, the pH of the solution is 9.8 and,
 8 from here onward, it becomes lower than the pK_a value of DEEA (9.84). Hence, the amine
 9 capacity, in terms of amount of conjugate base (DEEA), is reduced, i.e. the basicity of the
 10 amine solution is weaker, and since the concentration of CO_3^{2-} starts to be diminished, it can
 11 be concluded that CO_3^{2-} also reacts as a conjugate base,^{24, 46} as reported in reaction 11 (similarly
 12 to the generic reaction 4 or the DEEA-specific reaction 13).
 13 Concerning the heat of absorption (Figure 6), as general tendency, at increasing loading a
 14 steady and slight decrease is observed.



15
 16 **Figure 6.** NMR speciation and heat of absorption data in aqueous DEEA 3 M at increasing
 17 CO_2 loading (25 °C).

1 However, similarly to the pH curve, a rapid drop off occurs in the last-loading region, from
2 about 0.67 mol_{CO2}/mol_{amine} onward, which can hence be defined as the saturation loading
3 region.⁴⁷ Due to the reduction of the amine capability as conjugate base, alternative pathways
4 for the bicarbonate formation and/or CO₂ capturing (like e.g. reaction 11) may go along and
5 show to release less heat than the reactions occurring before the saturation region. Accordingly,
6 the DEEAOCOO⁻ tendency of increasing at increasing loading may also be attributable to the
7 saturation of the amine nitrogen capabilities. In this study, its formation is generically
8 represented by the reaction 14 since, with the data available, it was not possible to investigate
9 the reaction mechanisms of this species.

10 In summary, alongside the addition of CO₂, the reactions 13 and 14 are mainly occurring.
11 Reaction 13 represents the catalysis of the reaction 7 and it sums up the reactions 5 and 12.
12 Moreover, until about 0.5 mol_{CO2}/mol_{amine}, which corresponds to pH_≥9.8, the reaction 6,
13 leading to CO₃²⁻ species, also takes place; from here to the maximum loading (pH<9.8), the
14 CO₃²⁻ is instead consumed as represented in the reaction 11.

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3.3.2 MAPA 1M-CO₂-H₂O system

In Figure 7, the speciation of the aqueous diamine MAPA 1M at increasing CO₂ addition is reported together with the pH measurements.

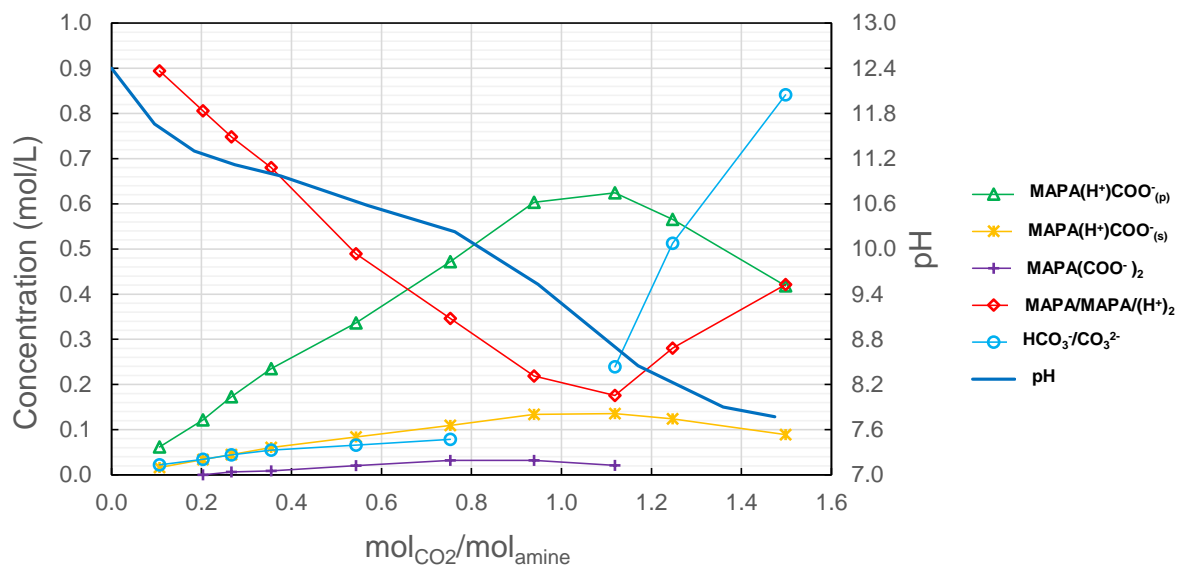


Figure 7. NMR speciation and pH data in aqueous MAPA 1M at increasing CO₂ loading (25 °C).

As soon as the solution is loaded with CO₂, MAPA/MAPA(H⁺)₂ starts decreasing to form primary and secondary MAPA carbamate (MAPA(H⁺)COO_(p)⁻ and MAPA(H⁺)COO_(s)⁻, respectively), as described in the reactions 20 and 21. The constant increase of MAPA(H⁺)COO_(p)⁻ and MAPA(H⁺)COO_(s)⁻ at increasing loading is evident until around 0.94 mol_{CO2}/mol_{amine}. Around this point, a sort of stabilization in the increase of these two species is observed, and from around 1.12 mol_{CO2}/mol_{amine} onward, their concentration decreases, i.e. hydrolysis reactions 24-25 occur.

MAPA(COO⁻)₂ is formed (reaction 22) in amounts in the order of the second decimal number and much lower than MAPA(H⁺)COO_{(p)/(s)}⁻. At beginning of CO₂ loading, the corresponding peaks in the ¹³C NMR spectra are not observed. From around 0.20 mol_{CO2}/mol_{amine} to 0.36 mol_{CO2}/mol_{amine} only traces (lower than 0.01 mol/L) are formed, and its amount increases until

1 around 0.75-0.94 mol_{CO2}/mol_{amine}. Around this region, the concentration is quite stable, and
2 from 0.94 mol_{CO2}/mol_{amine} onward, a decrease (hydrolysis reaction, as shown in reaction 23) is
3 observed.

4 In a previous NMR study on 0.5 M and 1M MAPA, the species MAPA(COO⁻)₂ was also found
5 in negligible amount.¹¹ The reason is that the nitrogen not bound to CO₂ in MAPACOO⁻_{(p)/(s)}
6 may not be available for reaction with CO₂ because, as given in reaction 20, it could be
7 protonated and act thus as counterion to the negative charged groups.
8 The concentration of MAPA/MAPA(H⁺)₂ constantly decreases until around 1.12
9 mol_{CO2}/mol_{amine} and, from here onward, it starts increasing due to the hydrolysis of
10 MAPA(H⁺)COO⁻_(p) and MAPA(H⁺)COO⁻_(s). Not all the MAPA in solution reacts with CO₂ as
11 it also goes through the reaction of protonation (Reaction 18 and 19) and, in the protonated
12 form, it can act as counter ion, like e.g. in reaction 21 and 22 the “Amine” may be MAPA.

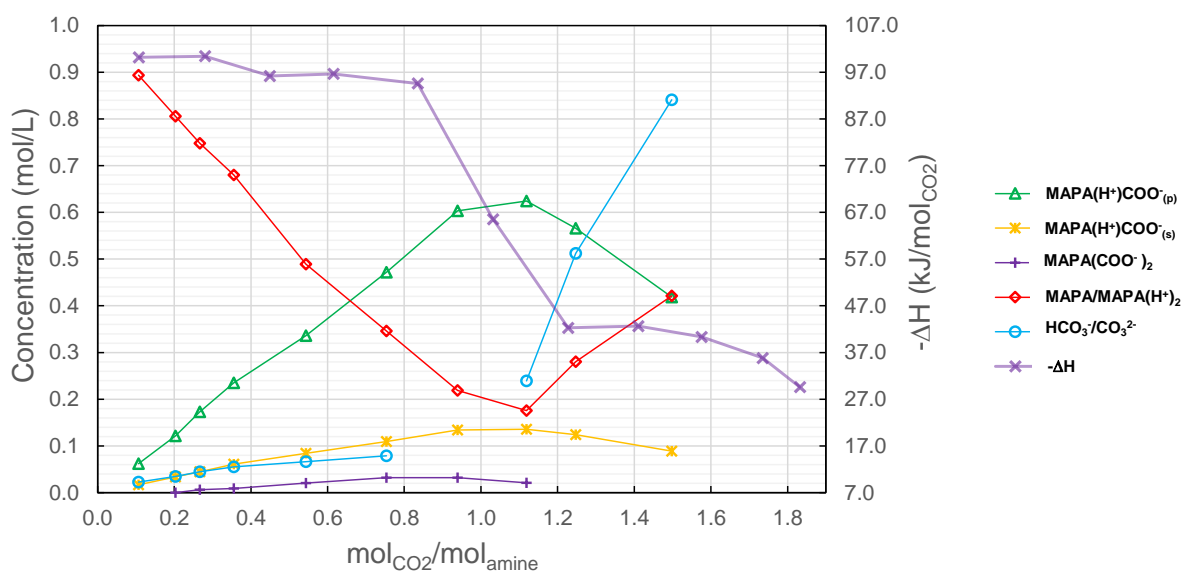
13 (Bi)carbonate is formed throughout the CO₂ loading range. It is mainly in concentration higher
14 than MAPA(COO⁻)₂ and lower than MAPACOO⁻_(p) but, around the region of the maximum
15 mol_{CO2}/mol_{amine}, it reaches concentrations higher than all the species in solution. Specifically,
16 its presence is more evident from the region of hydrolysis onward (at loading 0.94 the HCO₃⁻
17 /CO₃²⁻ signal was not detected because it was probably overlapped with the carbamate signals).

18 Concerning the pH, it decreases at increasing CO₂ loading. As soon as CO₂ is added into the
19 solution, a ΔpH drop of 0.74 is observed and, during the CO₂ addition, the pH values span from
20 11.66 at about 0.1 mol_{CO2}/mol_{amine} to 7.77 at about 1.5 mol_{CO2}/mol_{amine}. A change in the slope
21 of the curve is observed at about 0.76-1.17 mol_{CO2}/mol_{amine} and, specifically, a unit pH drop
22 occurs between 0.95 and 1.17 mol_{CO2}/mol_{amine}, which is the region where the MAPA
23 carbamates hydrolyses start.

24 The hydrolyses of MAPA carbamate species occurred at about pH 9.5 and the reactions 23-25
25 show how they may be driven by the acidity of the solution. The reaction of the nitrogen atom

1 of the amine carbamates with a proton makes the compound unstable, leading to the formation
 2 of protonated MAPA species and HCO_3^- . Alike, in a study by Lv et al.³⁴ it was reported that
 3 the hydrolysis of MEA carbamate was occurring at about pH 9. Similarly, Zhang et al.¹⁰ in the
 4 study of CO_2 loaded aqueous solution of MAPA 2M at 40 °C, showed that the reactions of
 5 carbamate hydrolysis started at about pH 9.5. The speciation trends in MAPA 2M are
 6 comparable to those observed in our study, but due to the different reaction conditions and
 7 concentrations, some differences are observed. In the present study, a lower end pH value at
 8 the same loading ($1.5 \text{ mol}_{\text{CO}_2}/\text{mol}_{\text{amine}}$) was reached since the work is dealing with MAPA 1M
 9 aqueous solution as compared to MAPA 2M. Hence, with respect to $\text{MAPACOO}^-_{(p)}$, more
 10 (bi)carbonate than reported by Zhang et al. is observed. Moreover, in our study, at the pH value
 11 of the carbamate hydrolysis, not all the MAPA in solution has reacted with CO_2 , as it was
 12 instead for MAPA 2M in Zhang et al.¹⁰

13 In Figure 8, the concentration of the species in aqueous MAPA 1M at increasing CO_2 loading
 14 are reported together with the heat of absorption.



15
 16 **Figure 8.** NMR speciation and heat of absorption data in aqueous MAPA 1M at increasing
 17 CO_2 loading (25 °C).

18

1 The heat of absorption is quite constant and stays approximately the same until 0.84
2 mol_{CO2}/mol_{amine}, where a sudden dropping off is observed (0.84-1.23 mol_{CO2}/mol_{amine}).
3 Carbamate is known to be an exothermic reaction releasing high heat of reaction, and the heat
4 released decreases as soon as the hydrolysis occurs and/or the carbamate is not formed
5 anymore.

6 Hence, in the range of the CO₂ loading (from the beginning to the maximum loading), two
7 main regions can be identified:

8 1) In the first region, going from about 0.11 to about 0.94 mol_{CO2}/mol_{amine}, the following
9 reactions are mainly occurring:

- 10 • Protonation of MAPA (reaction 18-19);
- 11 • Formation of primary and secondary MAPA carbamate (reaction 20-21), and in
12 negligible amount, formation of MAPA(COO⁻)₂ (reaction 22).
- 13 • Formation of (bi)carbonate in small amount through the reactions 5-7. Additionally,
14 MAPA may act by providing basicity (reaction 18-19), leading to a kind of CO₂
15 hydration catalysis (reaction 4) (as DEEA singly in reaction 13). However, in MAPA
16 solution, the carbamate formation is favored.

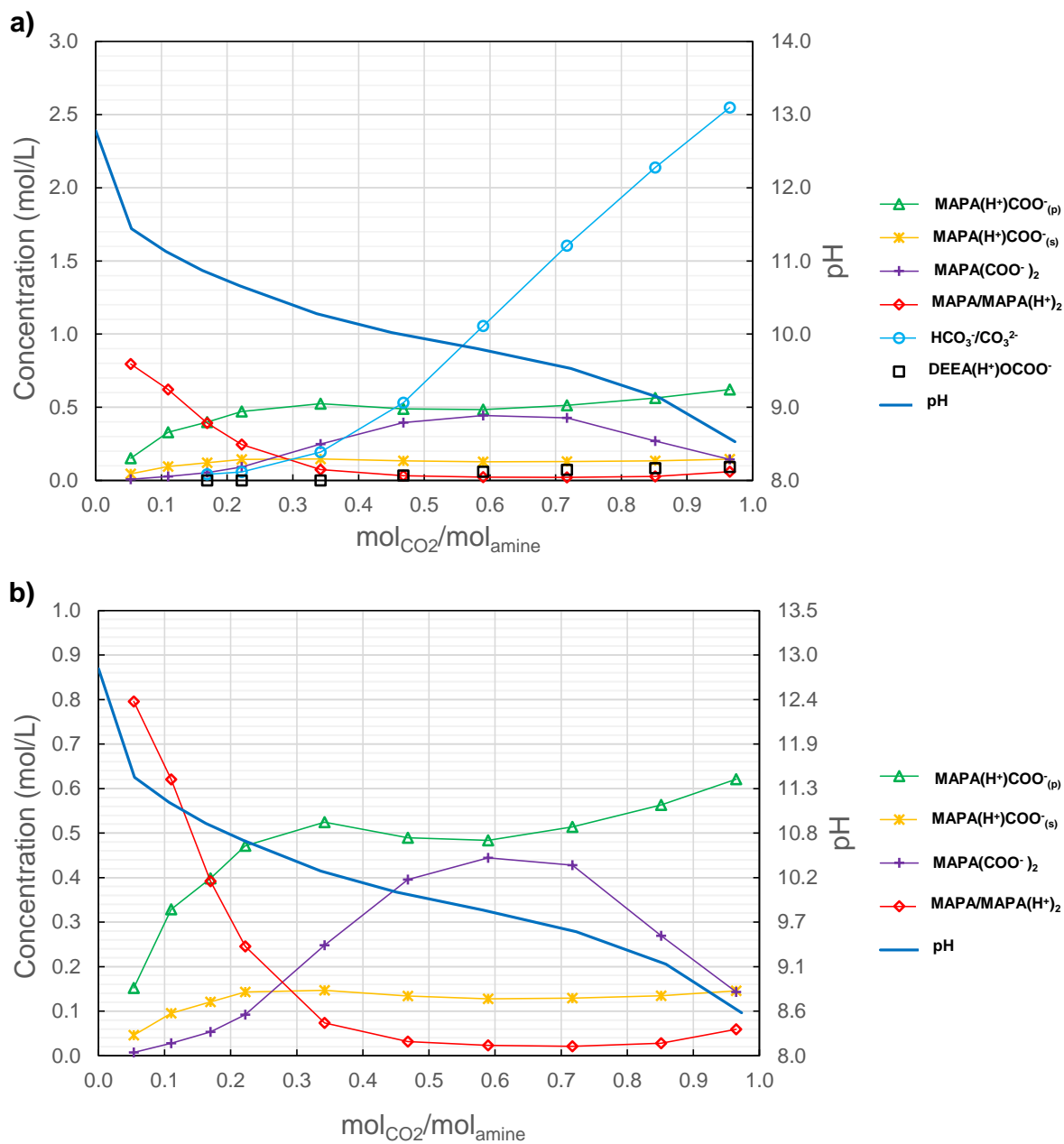
17 2) In the second region, from about 0.94 mol_{CO2}/mol_{amine} onward, the hydrolysis reactions
18 mainly take place:

- 19 • Around 0.94-1.25 mol_{CO2}/mol_{amine}, the hydrolysis of MAPA(COO⁻)₂ (reaction 23)
20 occurs;
- 21 • Around 1.12-1.5 mol_{CO2}/mol_{amine}, the hydrolyses of primary and secondary MAPA
22 carbamate (reactions 24 and 25) are predominant;
- 23 • (Bi)carbonate found in solution is mainly coming from the reactions of amine
24 carbamate hydrolysis (23-25), but the reactions 5, 7 and 11 may also take place
25 (reaction 5 is however negligible at pH lower than 8²⁵). Reaction 11 is supported by

1 the chemical shift value of the peaks corresponding to the $\text{HCO}_3^-/\text{CO}_3^{2-}$ species in the
2 ^{13}C NMR spectra. The ppm values are consistent with the presence of both CO_3^{2-} and
3 HCO_3^- until about 1.12 mol $_{\text{CO}_2}$ /mol $_{\text{amine}}$. From here onward, the ppm values are in line
4 with the only presence of HCO_3^- (the chemical shifts and the estimated concentrations
5 are available in Table S11 and in Figure S1 of the Supporting Information,
6 respectively).

1 3.3.3 MAPA 1M-DEEA 3M-CO₂-H₂O system

2 In Figure 9, the amount of each species in MAPA 1M-DEEA 3M-CO₂-H₂O is reported for each
3 CO₂ loading and interrelated with pH data.



4
5 **Figure 9.** Concentration of a) all the species and b) MAPA derivative species only, and pH
6 data in aqueous MAPA 1M-DEEA 3M at increasing CO₂ loading (25 °C).

7

1 The same species observed in MAPA 1M and DEEA 3M are found, but at different
2 concentrations and ratios.

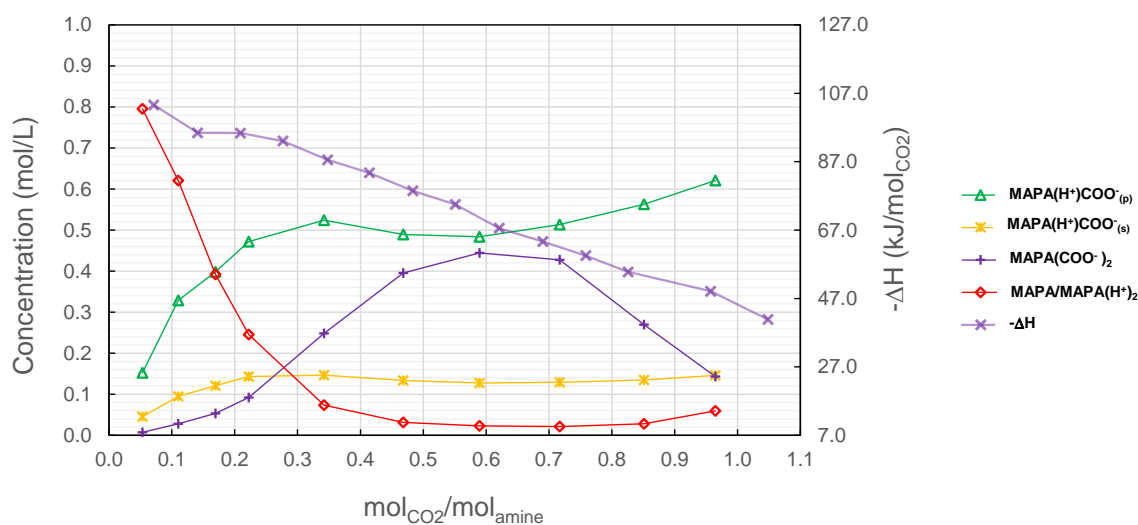
3 DEEAOCOO⁻ (given by reaction 14) has a trend like that observed in DEEA 3M, but it is not
4 detected since the early loadings.

5 MAPACOO⁻_(p) and MAPACOO⁻_(s), given by reactions 20-21, increase until around loading
6 0.34 mol_{CO2}/mol_{amine}. In the region around 0.34-0.59 mol_{CO2}/mol_{amine} a slight decrease and/or
7 stabilization of their concentrations is observed, whereas from 0.59 mol_{CO2}/mol_{amine} onward an
8 increase is again observed. From about 0.22 mol_{CO2}/mol_{amine} until around 0.59 mol_{CO2}/mol_{amine},
9 the concentration of MAPA(COO⁻)₂ (formed as in reaction 22) deeply increases to the
10 detriment of MAPACOO⁻_{(p)/(s)}. From 0.59 mol_{CO2}/mol_{amine} until maximum loading, it goes
11 through the hydrolysis process (reaction 23) and, therefore, decreases in favor of formation of
12 further MAPACOO⁻_{(p)/(s)}.

13 MAPA is almost totally consumed and the curve shows a steep decrease until loading 0.34
14 mol_{CO2}/mol_{amine}, (region where MAPA(H⁺)COO⁻_{(p)/(s)} do not show any more a steep increase).
15 From here to 0.72 mol_{CO2}/mol_{amine}, the decrease becomes slightly less evident, and it starts
16 slightly increasing from 0.72 mol_{CO2}/mol_{amine} to the maximum loading. The hydrolysis of
17 MAPACOO⁻_{(p)/(s)} may go along that of MAPA(COO⁻)₂, but its evidence is only given around
18 the last-loadings region where a slight increase of MAPA is observed.

19 Concerning the pH trend, it has the tendency of decreasing at increasing addition of CO₂.
20 Specifically, throughout the CO₂ loading range, the pH spans from 11.44 at 0.05
21 mol_{CO2}/mol_{amine} to 8.53 at maximum loading. A clear drop is observed between 0.00 and 0.05
22 mol_{CO2}/mol_{amine} (ΔpH=1.34), and the curve gets slightly steeper close to the maximum loading.
23 However, similarly to MAPA 1M, the amine carbamate hydrolysis starts at pH less than 10,
24 i.e. about pH 9.8.

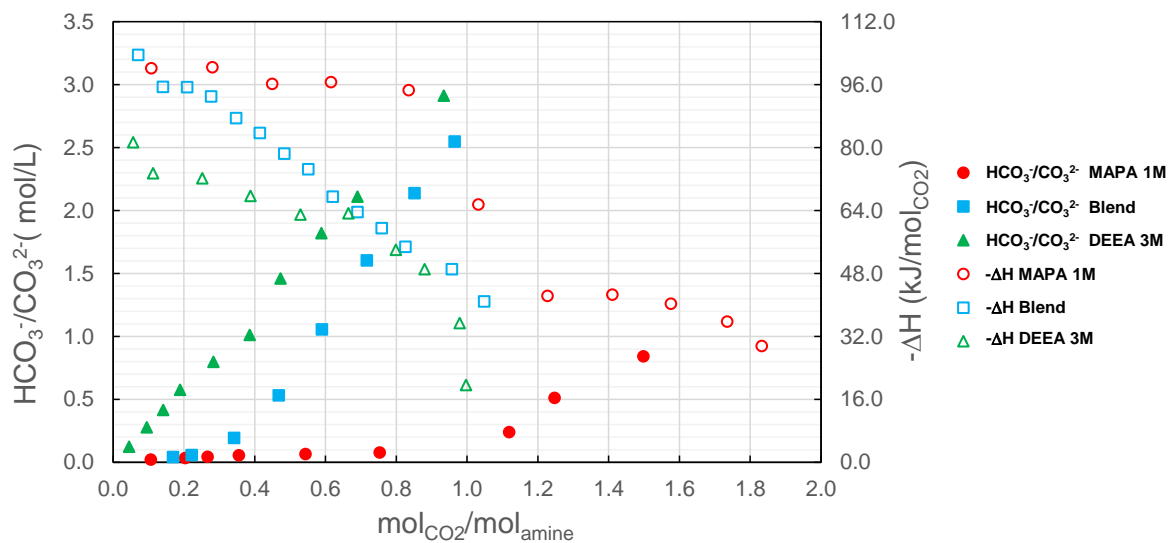
1 In Figure 10, the heat of CO₂ absorption in blend has been interrelated with the concentrations
 2 of the MAPA-derivative species.



3
 4 **Figure 10.** Concentration of MAPA-derivative species at different CO₂ loadings (25 °C) and
 5 heat of absorption trends in aqueous MAPA1M-DEEA3M.

6
 7 Throughout the CO₂ loading range, the heat continues to decrease steadily without any evident
 8 drop. Rather, the heat of absorption appears not to be influenced by the carbamate hydrolysis
 9 as it was instead observed in MAPA 1M.

10 The heat released by the DEEA-activity may somehow compensate the heat released by the
 11 carbamate formation, and it may lead to an absence of a rapid drop off in the region of
 12 carbamate hydrolysis. For clarifying the behavior, in the Figure 11, the (bi)carbonate
 13 concentrations and the heat of absorption at the different CO₂ loadings in the three systems are
 14 compared.

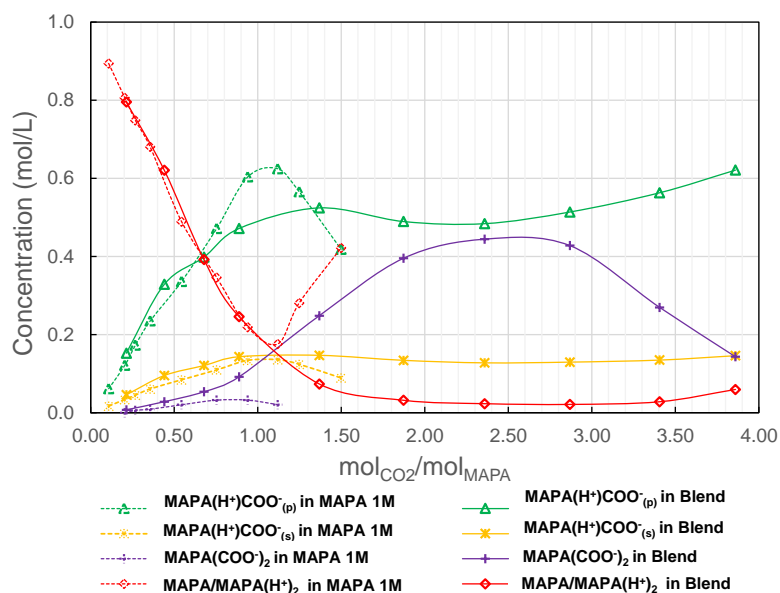


1
 2 **Figure 11.** Comparison of (bi)carbonate and heat of absorption in MAPA 1M, DEEA 3M and
 3 their blend.

4
 5 In the blend, until about 0.34 mol_{CO2}/mol_{amine}, the concentrations of (bi)carbonate is more like
 6 MAPA 1M than DEEA 3M, indicating that the activity of DEEA as Brønsted base has not yet
 7 an influence on the bicarbonate formation (as it is in DEEA 3M). Such activity is more evident
 8 from 0.34 mol_{CO2}/mol_{amine} onward, where the (bi)carbonate species has the same trend as
 9 DEEA 3M, but still lower concentrations. Similarly, the heats of absorption in MAPA 1M and
 10 MAPA 1M-DEEA 3M were comparable and almost constant until about 0.3 mol_{CO2}/mol_{amine}.
 11 From here, in MAPA 1M-DEEA3M it constantly decreased, while in MAPA 1M it stayed
 12 approximately the same until a loading of about 0.84 mol_{CO2}/mol_{amine}, the region where the
 13 amine carbamate hydrolyses take place.

14 The speciation data shown in Figures 9-11 represents the thermodynamic composition of the
 15 solution at the specific CO₂ loading reported. In the case of the blend, until around 0.34
 16 mol_{CO2}/mol_{amine}, we observe (Figure 9) that primary MAPA carbamate is the main product,
 17 followed by secondary MAPA carbamate and finally MAPA(COO⁻)₂. From about 0.34
 18 mol_{CO2}/mol_{amine}, along with formation of amine carbamates, further addition of CO₂ contributes

1 to $\text{HCO}_3^-/\text{CO}_3^{2-}$ formation, which is probably due to activity of DEEA as a Brønsted base.
 2 Moreover, from $0.59 \text{ mol}_{\text{CO}_2}/\text{mol}_{\text{amine}}$ onwards, MAPA carbamate hydrolysis reactions also
 3 contribute to bicarbonate formation. A similar situation is observed for 1M MAPA at
 4 approximately $0.8 \text{ mol}_{\text{CO}_2}/\text{mol}_{\text{amine}}$ (Figure 7 and 8). For MAPA 1 M this point corresponds to
 5 a significant drop in ΔH and pH indicating a change in formation of the predominant reaction
 6 product, i.e. towards bicarbonate. In the blend, the presence of 3M DEEA seems to blanket this
 7 clear change, and as soon as the trend for (bi)carbonate formation goes as in DEEA 3M, the
 8 released heat steadily decreases. However, due to the formation of the MAPA carbamate
 9 species, it is still higher than in the single DEEA 3M, as well as the amount of (bi)carbonate is
 10 lower.
 11 Moreover, to have a visual picture of DEEA's activity and influence on MAPA, the MAPA
 12 derivative species both in blend and in MAPA 1M are plotted as a function of $\text{mol}_{\text{CO}_2}/\text{mol}_{\text{MAPA}}$
 13 (Figure 12).



14
 15 **Figure 12.** Comparison of MAPA derivatives species in single and blended aqueous MAPA
 16 1M as a function of $\text{mol}_{\text{CO}_2}/\text{mol}_{\text{MAPA}}$
 17

1 It is possible to observe that the speciation curves for both single and blended MAPA are
2 similar until around 0.7 mol_{CO2}/mol_{MAPA}. Around this loading, in MAPA 1M the species
3 MAPA(H⁺)COO⁻_(p) continues increasing, while in the blend its concentration goes towards a
4 stabilization in favor of MAPA(COO⁻)₂ formation. Around 1.1 mol_{CO2}/mol_{MAPA}, in MAPA1M
5 the carbamate hydrolysis of MAPA(H⁺)COO⁻_{(p)/(s)} starts, while MAPA species in the blend
6 continue to react with CO₂ (e.g. MAPA(COO⁻)₂ starts swiftly increasing).

7 At increasing CO₂ loading, the amino-sites for absorbing CO₂ are saturated (not available), the
8 pH decreases, and the carbamate hydrolysis starts. In the presence of DEEA, MAPA is almost
9 totally consumed, meaning that the amino functional groups are more available for the
10 reactions of carbamate formation, and MAPA is then less protonated. The same is true for the
11 amino functional groups not bound to CO₂ in MAPACOO⁻_{(p)/(s)}.

12 Then, in the present study, the role played by DEEA in the presence of MAPA and CO₂
13 confirms the findings reported by Perinu et al.¹¹ for MAPA in blend with several tertiary
14 amines. Specifically, DEEA (pK_a=9.84), having the pK_a value larger than pK_{a2} of MAPA
15 (pK_{a1}= 10.6 and pK_{a2}=8.6), may withdraw protons from protonated MAPA species (reactions
16 15-16) and/or lead the protonation process (reaction 12 and 17), and in the protonated form it
17 acts as counter ion of the carbamate species (the term “Amine” in reactions 21-22). Jang et al.
18 2018 also concluded that DEEA, and generally the tertiary amines, can induce the reaction
19 from protonated MEA to active molecule MEA.⁴⁸

20 To summarize, in the blend with MAPA, the Brønsted base DEEA plays two main roles in the
21 presence of CO₂. It acts both by increasing the availability of MAPA’s nitrogens to form
22 carbamates (e.g., MAPA(COO⁻)₂ is formed in significant amount as compared to single
23 MAPA, where its concentration is negligible), and by catalyzing the hydration of CO₂ for
24 (bi)carbonate formation (as it is in the absence of MAPA).

1 Based on the above results and discussion, during the addition of CO₂ in aqueous MAPA 1M-
2 DEEA 3M, three main regions can be identified:

3 1) In the first region, spanning from 0.05 to 0.34 mol_{CO2}/mol_{amine}, the following reactions take
4 place:

- 5 • Amines protonation, i.e. MAPA protonation reactions (18-19), together with the
6 exchange-proton reactions with DEEA (reactions 15-17).
- 7 • Reactions 20-22 which lead to formation of MAPA carbamates.
- 8 • (Bi)carbonate is formed in small amount, and the reactions 5 and 6 may mainly occur (pH
9 is higher than 10 and DEEA is not yet involved in the CO₂ hydration catalysis).

10 2) The second region can be identified around 0.34-0.59 mol_{CO2}/mol_{amine}:

- 11 • Formation of MAPA(COO⁻)₂ (reaction 22)
- 12 • The DEEA has similar activity as in DEEA 3M.

13 3) In the third region (0.59-0.96 mol_{CO2}/mol_{amine}), carbamate hydrolysis reactions are mainly
14 occurring:

- 15 • MAPA(COO⁻)₂ hydrolysis is predominant (Reaction 23),
- 16 • Hydrolysis of MAPA(H⁺)COO⁻_{(p)/(s)} (reactions 24-25) may be considered a minor one
- 17 • Along with the reactions as in DEEA 3 M singly, (bi)carbonate found in solution is coming
18 from the carbamate hydrolysis (reactions 23-25). The reaction 11 may also occur and it is
19 supported by the chemical shift values of HCO₃⁻/CO₃²⁻ and corresponding concentrations
20 (see the chemical shift values in Table S11 and the estimated concentrations in Figure S2
21 of the Supporting Information).

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1 4. CONCLUSIONS

2 In this study, aqueous MAPA 1M, DEEA 3M and their blend were investigated in the presence
3 of CO₂. Based on speciation, heat of absorption and pH data analyses, the main reactions
4 occurring at various CO₂ addition were identified and discussed.

5 At increasing CO₂ loading, the basicity of the solution decreases, as well as the capability of
6 each amine to absorb CO₂. In the aqueous MAPA systems, this leads to the hydrolysis of the
7 carbamate species, which also corresponds to a drop in the released heat. The reactions of
8 carbamate hydrolysis in both single and blended MAPA occur at similar pH values (i.e. about
9 pH 9.8-9.5), but in the blended MAPA, due to the activity of DEEA, the drop in both pH and
10 heat of absorption is not so noticeable as in MAPA singly. The Brønsted basicity of DEEA
11 indeed acts both by catalyzing the CO₂ hydration reaction and by allowing the diamine to be
12 more available for the direct reaction to CO₂, i.e. carbamate formation.

13 In the investigated amine systems, a saturation region can be identified in the region close to
14 the maximum loading, and it is observed when the amine capability, in terms of conjugate
15 Brønsted base and/or Lewis base, is reduced and/or saturated.

16 With the present study, crucial insights for MAPA and DEEA solvents, singly and in blend,
17 are given and can be considered valuable for the selection of new amine solvents with improved
18 properties.

19

20 ASSOCIATED CONTENT

21 **Supporting Information.** The following file is available free of charge. Density, pH, heat of
22 absorption, NMR speciation, chemical shifts and estimated concentration of bicarbonate and
23 carbonate species (PDF).

24

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4

5 **Notes**

6 The authors declare no competing financial interest.

7

8 ACKNOWLEDGMENT

9 This work was supported by the financial assistance and scholarships (C.P. and I.M.B.)
10 provided by the Research Council of Norway (CLIMIT, grant nr. 243620/E20) which is
11 gratefully acknowledged. The NMR spectra were recorded by C.P. in the NMR Laboratory at
12 the Faculty of Natural Science of the Norwegian University of Science and Technology
13 (NTNU, Trondheim) and the assistance of the staff is acknowledged.

14

15

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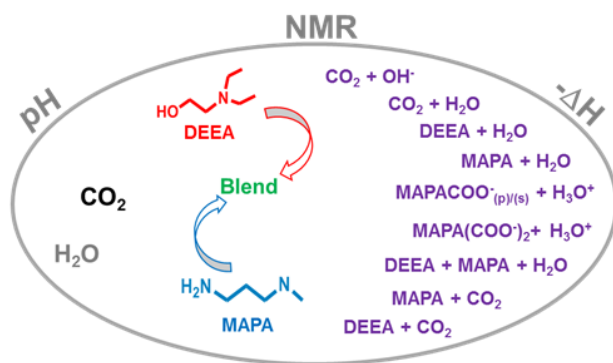
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Supporting Information

Aqueous MAPA, DEEA and their blend as CO₂ absorbents: Interrelationship between NMR speciation, pH and heat of absorption

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Table S1: Density, as measured by Anton Paar DMA 4500 M densitometer, of aqueous MAPA 1M, DEEA 3M and MAPA 1M-DEEA 3M before addition of CO₂.

	density (g/ml)
MAPA 1M - DEEA 3M	0.97572
MAPA 1M	0.99094
DEEA 3M	0.98334

Table S2. Weight of 1 mL of each CO₂ loaded sample.

MAPA 1M - DEEA 3M										
mol_{CO2}/mol_{amine}	0.05	0.11	0.17	0.22	0.34	0.47	0.59	0.72	0.85	0.96
density (g in 1ml)	0.9797	0.9851	0.999	1.0132	1.026	1.049	1.0685	1.084	1.102	1.1121
MAPA 1M										
mol_{CO2}/mol_{amine}	0.11	0.20	0.27	0.36	0.54	0.75	0.94	1.12	1.25	1.50
density (g in 1ml)	0.9866	0.9908	1.0056	1.0061	1.005	1.0201	1.0268	1.0247	1.0359	1.0444
DEEA 3M										
mol_{CO2}/mol_{amine}	0.04	0.10	0.14	0.19	0.28	0.39	0.47	0.59	0.69	0.93
density (g in 1ml)	0.9881	0.9879	1.0026	1.0016	1.0209	1.0312	1.0462	1.055	1.0633	1.0846

Table S3. pH measured in each sample.

MAPA1M- DEEA 3M											
mol_{CO2}/mol_{amine}	0.00	0.05	0.11	0.16	0.22	0.34	0.45	0.58	0.72	0.86	0.97
pH	12.78	11.44	11.13	10.87	10.66	10.28	10.02	9.8	9.53	9.13	8.53
MAPA 1M											
mol_{CO2}/mol_{amine}	0.00	0.10	0.18	0.27	0.38	0.57	0.76	0.95	1.17	1.36	1.47
pH	12.4	11.66	11.3	11.12	10.97	10.58	10.23	9.53	8.45	7.9	7.77
DEEA 3M											
mol_{CO2}/mol_{amine}	0.00	0.05	0.09	0.14	0.18	0.27	0.37	0.47	0.56	0.67	0.98
pH	12.07	10.79	10.53	10.38	10.27	10.11	10.01	9.82	9.67	9.57	8.41

Table S4. Differential heat of absorption of CO₂ in aqueous MAPA 1M at 25°C.

mol_{CO2}/mol_{amine}	-ΔH (kJ/mol_{CO2})
0.11	100.205
0.28	100.458
0.45	96.224
0.62	96.670
0.84	94.611
1.03	65.527
1.23	42.289
1.41	42.655
1.58	40.320
1.74	35.798
1.83	29.562

Table S5. Differential heat of absorption of CO₂ in aqueous MAPA 1M at 40°C.

mol_{CO2}/mol_{amine}	-ΔH (kJ/mol_{CO2})
0.21	98.2799
0.43	94.2602
0.65	93.6375
0.87	88.0429
1.08	63.7486
1.27	43.9388
1.46	36.7198
1.70	35.1281

Table S6. Differential heat of absorption of CO₂ in aqueous DEEA 3M at 25°C.

mol_{CO2}/mol_{amine}	-ΔH (kJ/mol_{CO2})
0.06	81.33
0.11	73.51
0.25	72.23
0.39	67.74
0.53	62.99
0.67	63.32
0.80	54.02
0.88	49.09
0.98	35.38
1.00	19.61

Table S7. Differential heat of absorption of CO₂ in aqueous DEEA 3M at 40°C.

mol_{CO2}/mol_{amine}	-ΔH (kJ/mol_{CO2})
0.06	78.99
0.12	73.38
0.23	70.02
0.34	67.54
0.44	65.37
0.56	60.14
0.72	57.31
0.88	51.35
0.98	37.01

Table S8. Differential heat of absorption of CO₂ in aqueous MAPA 1M-DEEA 3M at 25°C.

mol_{CO2}/mol_{amine}	-ΔH (kJ/mol_{CO2})
0.07	103.60
0.14	95.41
0.21	95.35
0.28	93.04
0.35	87.53
0.41	83.73
0.48	78.48
0.55	74.49
0.62	67.55
0.69	63.65
0.76	59.52
0.83	54.78
0.96	49.11
1.05	40.88

Table S9. Concentration (mol/L) of each species in aqueous MAPA 1M-CO₂ systems.

mol_{CO2}/mol_{amine}	0.11	0.20	0.27	0.36	0.54	0.75	0.94	1.12	1.25	1.50
MAPACOO⁻_(p)	0.062	0.121	0.173	0.235	0.336	0.472	0.604	0.625	0.566	0.419
MAPA(COO⁻)₂		0.000	0.006	0.009	0.020	0.032	0.032	0.021		
MAPACOO⁻_(s)	0.016	0.034	0.045	0.061	0.084	0.109	0.134	0.135	0.124	0.089
MAPA	0.894	0.806	0.748	0.680	0.489	0.346	0.219	0.176	0.281	0.421
HCO₃⁻/CO₃²⁻	0.022	0.035	0.044	0.055	0.066	0.078		0.239	0.513	0.841

Table S10. Concentration (mol/L) of each species in aqueous DEEA 3M-CO₂ systems.

mol _{CO2} /mol _{amine}	0.04	0.10	0.14	0.19	0.28	0.39	0.47	0.59	0.69	0.93
DEEA	2.806	2.999	2.979	3.261	2.893	3.023	3.163	2.963	3.018	2.991
DEEAOCOO ⁻	0.01	0.02	0.023	0.021	0.043	0.057	0.072	0.078	0.087	0.108
HCO ₃ ⁻ /CO ₃ ²⁻	0.123	0.279	0.417	0.575	0.797	1.012	1.459	1.820	2.110	2.912

Table S11. Concentration (mol/L) of each species in aqueous MAPA1M-DEEA3M-CO₂ systems

mol _{CO2} /mol _{amine}	0.05	0.11	0.17	0.22	0.34	0.47	0.59	0.72	0.85	0.96
MAPACOO ⁻ _(p)	0.152	0.328	0.398	0.472	0.524	0.489	0.484	0.514	0.563	0.621
MAPA(COO ⁻) ₂	0.007	0.028	0.054	0.092	0.248	0.395	0.444	0.428	0.269	0.143
MAPACOO ⁻ _(s)	0.046	0.095	0.121	0.143	0.147	0.134	0.128	0.129	0.135	0.146
MAPA	0.795	0.621	0.392	0.246	0.074	0.032	0.023	0.021	0.028	0.059
HCO ₃ ⁻ /CO ₃ ²⁻			0.041	0.057	0.193	0.531	1.056	1.604	2.138	2.549
DEEA	3.063	3.255	2.956	2.888	3.023	3.163	3.230	3.173	2.897	2.769
DEEAOCOO ⁻			0.000	0.000	0.000	0.034	0.059	0.073	0.084	0.092

Table S12. Chemical shift values of the signal corresponding to HCO₃⁻/CO₃²⁻ in the ¹³C NMR spectra of aqueous MAPA 1M, DEEA 3M and MAPA 1M – DEEA 3M at increasing addition of CO₂ into the solution.

MAPA 1M	mol _{CO2} /mol _{amine}	0.11	0.2	0.27	0.36	0.54	0.75	0.94	1.12	1.25	1.5
	δ (ppm) HCO ₃ ⁻ /CO ₃ ²⁻	168.2	168.1	167.8	167.8	167.3	166.4		161.9	161.1	161
DEEA 3M	mol _{CO2} /mol _{amine}	0.04	0.1	0.14	0.19	0.28	0.39	0.47	0.59	0.69	0.93
	δ (ppm) HCO ₃ ⁻ /CO ₃ ²⁻	165.7	165.2	164.8	164.5	164.1	163.7	163.3	162.8	162.3	161.3
MAPA1M- DEEA 3M	mol _{CO2} /mol _{amine}	0.05	0.11	0.17	0.22	0.34	0.47	0.59	0.72	0.85	0.96
	δ (ppm) HCO ₃ ⁻ /CO ₃ ²⁻			165.7	165.1	163.9	163	162.4	161.9	161.4	161.2

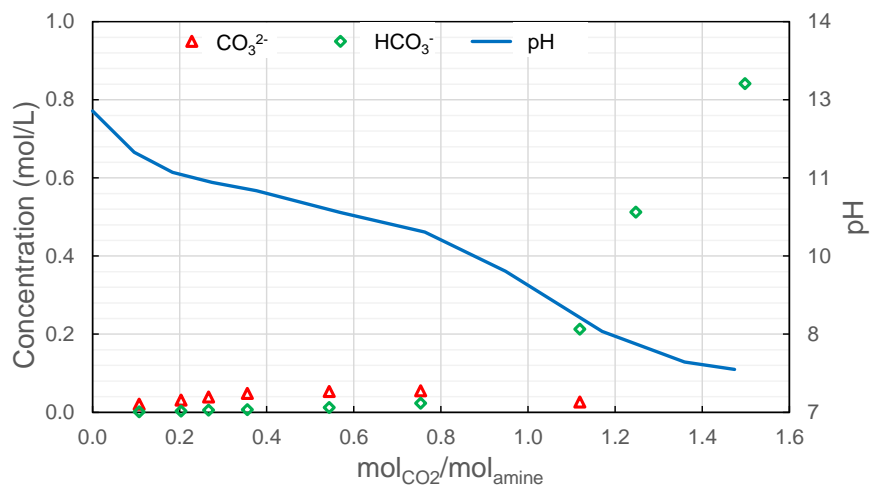


Figure S1. Estimation of the HCO_3^- and CO_3^{2-} content in MAPA 1M- CO_2 - H_2O systems. Calculation based on the calibration curve developed in Perinu et al. 2014.¹

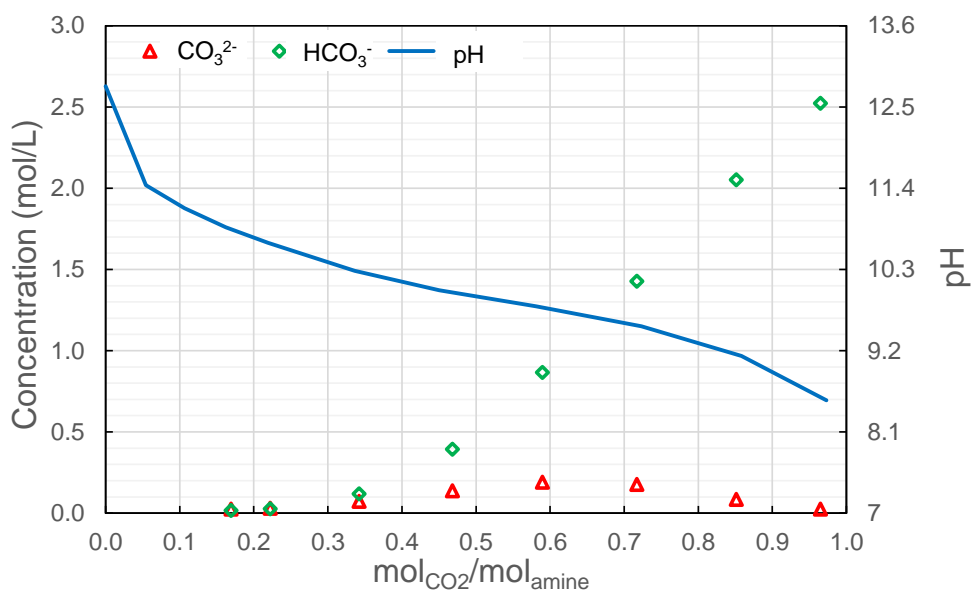


Figure S2. Estimation of the HCO_3^- and CO_3^{2-} content in MAPA 1M-DEEA 3M- CO_2 - H_2O systems. Calculation based on the calibration curve developed in Perinu et al. 2014.¹

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