

Dissociation Constants of Some Alkanolamines at 293, 303, 318, and 333 K

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The pK_a values were determined potentiometrically for the conjugate acids of 2-(2-aminoethoxy)ethanol (DGA), 2-(methylamino)ethanol (MMEA), 2-(*tert*-butylamino)ethanol (TBAE), 2-amino-2-methyl-1-propanol (AMP), *N*-methyldiethanolamine (MDEA), 2-(dimethylamino)ethanol (DMMEA), 2-(diethylamino)ethanol (DEMEA), and 2-(diisopropylamino)ethanol (DIPMEA) at 293, 303, 318, and 333 K.

1. Introduction

Alkanolamine solutions are used frequently to remove acidic components (e.g., H_2S , COS, and CO_2) from natural and refinery gases. Industrially important alkanolamines for this operation are the secondary amines diethanolamine (DEA) and diisopropanolamine (DIPA) and the tertiary amine *N*-methyldiethanolamine (MDEA). Usually aqueous solutions of these alkanolamines are applied; however, combined solvents like water and sulfolane in the Shell Sulfinol process are also used (1). Often there is no reason to remove CO_2 ; therefore, selective absorption of sulfur compounds from acid gas streams may lower the costs of the treating process considerably (2).

Selective absorption of particularly H_2S is among other achieved by reducing the reaction rate of CO_2 . Generally this reaction rate is for tertiary amines lower than for primary and secondary amines, which explains the increasing popularity of MDEA-based processes for selective absorption. Also processes based on sterically hindered (secondary) amines, like the Exxon Flexsorb process, seem to be suitable for this purpose. In order to be able to improve the selectivity, extensive knowledge of the reaction of CO_2 with alkanolamines is needed. In the past decade, a large number of articles have been published on this subject (e.g., Versteeg and Van Swaaij (3, 4)). Apart from the reaction rate of CO_2 with alkanolamines, also the capacity of alkanolamines for acidic components is an important factor in gas treating processes.

For both CO_2 reaction rate and capacity, the dissociation constant of the alkanolamine applied is an important variable. Versteeg and Van Swaaij (3) reported for primary and secondary amines a Brønsted relationship between the second-order rate constant for the formation of the zwitterion and the dissociation constant of the alkanolamine. Also for tertiary amines, a similar relationship at 293 K between the second-order rate constant and the dissociation constant of the alkanolamine was observed (4).

Dissociation constants of various alkanolamines at various temperatures have been accumulated by Perrin (5). Schwabe et al. (6) published data for *N*-methyldiethanolamine at 298, 308, 318, and 333 K. For diethanolamine and diisopropanolamine in an aqueous 1.00 M potassium chloride solution at temperatures of 293, 303, 313, and 323 K, information was reported by Blauwhoff and Bos (7). Antelo et al., (8) studied the influence of the ionic composition of the medium on the dissociation constants of monoethanolamine, diethanolamine, and triethanolamine at 298 K. They found a linear relationship between pK_a and $I^{1/2}$.

Table I. pH Values of the Calibration Buffers

temperature, K	pH			
	buffer 1	buffer 2	buffer 3	buffer 4
293	2.00	4.00	7.00	10.00
303	2.00	4.01	6.98	9.89
318	2.00	4.67	6.84	9.04
333	2.00	4.69	6.84	8.96

Although the dissociation constants of well-known and industrially important alkanolamines have been reported at several temperatures, dissociation constants of less common alkanolamines are still lacking. Therefore, the aim of the present investigation was to provide additional and reliable dissociation constants for various alkanolamines at several temperatures.

2. Experimental Section

2.1. Chemicals. For the experiments, the chemicals used were obtained from Janssen Chimica [2-(methylamino)ethanol (MMEA), 2-(*tert*-butylamino)ethanol (TBAE), 2-amino-2-methyl-1-propanol (AMP), *N*-methyldiethanolamine (MDEA), and 2-(diisopropylamino)ethanol (DIPMEA)] and Merck [2-(dimethylamino)ethanol (DMMEA) and 2-(diethylamino)ethanol (DEMEA)]. All these compounds were p.a. quality. Texaco Chemical Co. provided 2-(2-aminoethoxy)ethanol (DGA). All compounds were used as received.

Aqueous hydrochloric acid (0.1000 M) was used as titrant and prepared from Merck Titrisol ampules solved in aqueous 0.1 M tetraethylammonium chloride (TEACl, Merck p.a.; recrystallized twice from chloroform/ethyl acetate, washed with diethyl ether, and dried under vacuum conditions at 120 °C).

The following pH buffers were used in the calibration of the glass electrode: pH = 2.00, 4.00, 7.00, 9.00, 10.00, 4.66, 6.88, and 9.22 (all from Merck; pH values given at 293 K).

All solutions were prepared with carbon dioxide free double-distilled water.

2.2. Procedure. The pK_a values of the conjugate acids were determined by potentiometric titration of the basic compounds ($0.30\text{--}0.35 \times 10^{-3}$ mol in 25×10^{-3} L of solution) at constant ionic strength (0.1 M TEACl) with 0.1000 M HCl (in 0.1 M TEACl). The titration curves were recorded by computerized equipment consisting of a Mettler Model DV11 motorburet, a Knick industrial pH meter, type DIN, and a thermostated titration vessel. The pH meter was read by a digital voltmeter (Tekelec Airtronic, 4 digits), which was connected to the computer (PDP-11/84, Digital Equipment Corp.) via a DR11 digital I/O interface. The motorburet was driven by the computer also via this interface. Calibration of the combined glass/reference electrode (Metrohm Model AG9100) was done by linear regression of the millivolt response of the electrode on four buffer solutions. The pH values used are presented in Table I.

The sample and buffer solutions were thermostated at the required temperature (± 0.1 K) before use. Carbon dioxide was excluded from the titrations by flushing the titration vessel with nitrogen.

2.3. Calculation. The pK_a values of the samples were calculated from titration data (per titration curve, 60 or more pH- 10^{-3} L data pairs) with the use of the SUPERQUAD computer program (Gans et al. (9)). As this program does not take into account the values of the activity coefficients in its equations,

Table II. pK_a Values of Conjugate Acids of Alkanolamines^a

compound	293 K	303 K	318 K	333 K
<i>N</i> -methyldiethanolamine	8.76	8.49	8.28	7.99
2-(dimethylamino)ethanol	9.23	8.99	8.69	8.36 ^b
2-(2-aminoethoxy)ethanol	9.62	9.31	8.98	8.60
2-(diethylamino)ethanol	9.76	9.47 ^b	9.16 ^b	8.71 ^c
2-amino-2-methyl-1-propanol	9.88	9.52	9.20	8.78
2-(methylamino)ethanol	9.95	9.64	9.35	8.94
2-(diisopropylamino)ethanol	10.14	9.82	9.53	9.13
2-(<i>tert</i> -butylamino)ethanol	10.29	9.94	9.64	9.28

^a Accuracy, ± 0.02 pK_a unit. K_a values in mol/L. ^b Accuracy, ± 0.03 pK_a unit. ^c Accuracy, ± 0.04 pK_a unit.

the following steps were used to obtain the thermodynamically correct pK_a values for infinite dilution: (1) conversion of the thermodynamic K_w value to the "formal" value to be used as input by the SUPERQUAD program and (2) conversion of the pK_a value given by the computer program to its thermodynamic value. The activity coefficients needed for steps 1 and 2 were calculated by the Debye-Hückel equation:

$$-\log(\gamma_i) = \frac{Az_i^2 I^{1/2}}{1 + Bk_i I^{1/2}} \quad (1)$$

The values of A and B in this equation at the various temperatures were taken from Manov et al. (10). The ionic radius parameters k used in these calculations were obtained from Kielland (11): $k = 9$ (H), $k = 3$ (OH), $k = 4$ (MMEA, AMP, DMMEA, DGA), $k = 5$ (TBAE, DIPMEA, MDEA, DEMEA). The pK_w values used were taken from literature (12). The Debye-Hückel equation (1) represents activity coefficients with very good accuracy up to $K = 0.1$ M (13, 14).

3. Results and Discussion

In the procedure outlined in the preceding text, it should be noted that the response of the glass electrode must be calibrated in terms of the concentration (not the activity) of the hydrogen ion. For titrations in a medium of constant ionic strength, this difference tends to be canceled by the difference of the liquid junction potentials between standard buffers and the titration medium (15). For the correctness of the complete procedure to be ascertained, the pK_a values of benzoic acid and imidazole were determined at 298 K. The value for benzoic acid was found to be 4.22 ± 0.02 and equals the literature value of 4.20 (16) within experimental error. For imidazole a value of 7.08 ± 0.02 was found. The literature values for imidazole (17-20) were extrapolated to zero ionic strength with the help of eq 1 and coincide with the present result within experimental error. Furthermore, the inertness of the ionic strength buffer toward the alkanolamines was checked by carrying out some titrations at various buffer concentrations.

All determinations were carried out in triplicate, and the results were reproducible within ± 0.02 pK_a unit. The data given in Table II are the mean values of these triplicate measurements. The SUPERQUAD program calculates the accuracy of its results based on the fit of the experimental data to the theoretical equations and the experimental error in the volume of titrant added ($\pm 0.02 \times 10^{-3}$ L). In almost all cases, this error was ± 0.02 pK_a unit.

For *N*-methyldiethanolamine, some pK_a values were determined by Schwabe et al. (6). Their values deviate systematically from our values, the difference being -0.12 pK_a units. This discrepancy cannot be explained from the approximations they make (i.e., equal activity coefficients for the H^+ and the Cl^- ion and the use of the limiting law of Debye-Hückel up to concentrations of 0.04 M), which account only for a 0.01-0.04 pK_a units difference over the concentration range studied. It may be possible that the discrepancy between the present results and those of Schwabe et al. originates from the use of two different calibration methods for the pH scale. Schwabe et al. use the standard potential of the calomel electrode as given by Hills and Ives (21) (i.e., 0.26796 B) to fix their pH scale. In this work the pH scale was established with the use of the newer NBS standard buffer solutions (22).

Glossary

A, B	constants, see eq 1
I	ionic strength
k	ionic radius parameter, see eq 1
M	molarity [mol/L]
K_a	acid dissociation constant at $I = 0$ [mol/L]
z	charge number
γ	activity coefficient

Registry No. DGA, 929-06-6; MMEA, 109-83-1; TBAE, 4620-70-6; AMP, 124-68-5; MDEA, 105-59-9; DMMEA, 108-01-0; DEMEA, 100-37-8; DIPMEA, 96-80-0.

Literature Cited

- (1) Kohl, A. L.; Riesenfeld, F. C. *Gas Purification*; Gulf Publishing Co.: Houston, TX, 1979.
- (2) Blauwhoff, P. M. M.; Kamphuis, B.; van Swaaij, W. P. M.; Westerterp, K. R. *Chem. Eng. Proc.* **1985**, *19*, 1.
- (3) Versteeg, G. F.; van Swaaij, W. P. M. *Chem. Eng. Sci.* **1988**, *43*, 573.
- (4) Versteeg, G. F.; van Swaaij, W. P. M. *Chem. Eng. Sci.* **1988**, *43*, 587.
- (5) Perrin, D. D. *Dissociation Constants of Organic Bases in Aqueous Solution*; Butterworth: London, 1965.
- (6) Schwabe, K.; Graichen, W.; Spiethoff, D. *Z. Phys. Chem. (Munich)* **1959**, *20*, 68.
- (7) Blauwhoff, P. M.; Bos, M. J. *Chem. Eng. Data* **1981**, *26*, 7.
- (8) Antelo, J. M.; Arce, F.; Casado, J.; Sastre, M.; Varela, A. *J. Chem. Eng. Data* **1984**, *29*, 10.
- (9) Gans, P.; Sabatini, A.; Vacca, A. *J. Chem. Soc., Dalton Trans.* **1983**, 1195.
- (10) Manov, G. G.; Bates, R. G.; Hamer, W. J.; Acree, S. F. *J. Am. Chem. Soc.* **1943**, *65*, 1765.
- (11) Kielland, J. *J. Chem. Am. Soc.* **1937**, *59*, 1675.
- (12) Landolt-Bornstein. *Zahlenwerte und Funktionen*; Springer-Verlag: Berlin, 1960; Vol. II, Band 7, p 841.
- (13) Kortüm, G. *Textbook of Electrochemistry*; Elsevier: Amsterdam, 1951; Vol. I.
- (14) Robinson, R. A.; Stokes, R. M. *Electrolyte Solutions*; Butterworth: London, 1959.
- (15) Irving, H. M.; Miles, M. G.; Pettit, L. D. *Anal. Chim. Acta* **1967**, *38*, 475.
- (16) Landolt-Bornstein. *Zahlenwerte und Funktionen*; Springer-Verlag: Berlin, 1960; Vol. II, Band 7, p 867.
- (17) Tanford, C.; Wagner, M. L. *J. Am. Chem. Soc.* **1953**, *75*, 434.
- (18) Edsall, J. T.; Felsenfeld, G.; Goodman, D. S.; Gurd, F. R. N. *J. Am. Chem. Soc.* **1954**, *76*, 3054.
- (19) Mickel, B. L.; Andrews, A. C. *J. Am. Chem. Soc.* **1955**, *77*, 5291.
- (20) Brooks, P.; Davidson, N. *J. Am. Chem. Soc.* **1960**, *82*, 2118.
- (21) Hills, G. J.; Ives, D. J. G. *J. Chem. Soc.* **1951**, 318.
- (22) Bates, R. G.; Guggenheim, E. A. *Pure Appl. Chem.* **1960**, *1*, 163.

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