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Sengeløv, Louise With; Thomsen, Kaj

Publication date: 2011

Document Version Early version, also known as pre-print

Link back to DTU Orbit

Citation (APA): Sengeløv, L. W., & Thomsen, K. (2011). Phase Equilibrium in Amino Acid Salt Systems for CO2 Capture [Sound/Visual production (digital)]. WORKSHOP - Amino Acid Salts for CO2 Capture, Adminiet, Porsgrunn, Norway, 01/01/2011

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Phase Equilibrium in Amino Acid Salt Systems for CO₂ Capture

Louise With Sengeløv and Kaj Thomsen (kth@kt.dtu.dk)

Porsgrunn, February 3rd 2011



DTU Chemical Engineering Department of Chemical and Biochemical Engineering

Outline



Introduction

Advantages in using amino acid salts instead of alkanolamines:

- Might be less prone to degradation in oxygen rich environments
- Less volatile than alkanolamines due to their ionic nature
- Same affinity towards CO₂ as alkanolamines
- Form carbamates

Amino acids participate in the transportation of CO_2 in the blood by the formation of carbamate

Problems and drawbacks:

- More expensive
- Limited solubility of amino acids in water
- Limited amount of experimental data available in literature







Experimental data for amino acid salt solutions

• CO₂ Equilibrium data

- Potassium taurate Kumar *et al.* (2003) **38 data points**, T=25 and 40°C
- Potassium glycinate (Portugal et al. 2009) 103 data points, T= 20-50°C
- Potassium methionate (Kumelan et al. 2010) 65 data points, T= 80-120°C
- Potassium sarcosinate (Aronu *et al.* 2010) graph with 55 points, T=40-120°C
- Potassium prolinate (unpublished, DTU, UTwente)
- Absorption kinetics
 - Kumar et al. (2003) (potassium salt of taurine and glycine)
 - Van Holst *et al.* 2009 (potassium salts of 7 different amino acids)
 - Prakash et al. 2010 (potassium salts of taurine and glycine)
- Equilibrium constants
 - Sharma *et al.* 2003
 - Hamborg et al. 2007
- Heat capacity and other thermal properties none
- Freezing point depression data
 - Sengeløv (2010) (loaded and unloaded potassium salt of methionine) 20 data points -14 to 0°C

Methionine, C₅H₁₁NO₂S



- Molar mass 149.21 g/mol
- Decomposition temperature 281°C
- Solubility in water at 25°C: 0.38 molal
- pKa 2.28 and 9.21
- Found in cereal grains and in nuts

Iso-electric point of methionine



рΗ

Solubility of methionine in water **O** and in NaCl solution **X**



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Hill and Robson, Biochemical Journal, 28(1934)1008-1013

Potassium salt of methionine – Equilibrium constants

- Sharma VK; Zinger A; Millero FJ; De Stefano C; *Dissociation constants of protonated methionine species in NaCl media*, Biophysical Chemistry, 105(2003)79-87
 - Determined in temperature range 5 45°C
- Hamborg ES; Niederer JPM; Versteeg GF; Dissociation constants and thermodynamic properties of amino acids used in CO₂ absorption from (293 to 353) K, J. Chem. Eng. Data, 52(2007)2491-2502
 - Determined in temperature range 20 80°C
- Equilibrium constant for the carbamate formation of methionine not available
- Equilibrium constant for the protonated form of methionine not relevant at pH above 6

Potassium salt of methionine – experimental data

- Kumelan J, Pérez-Salado Kamps A, Maurer G, Solubility of CO₂ in Aqueous Solutions of Methionine and in Aqueous Solutions of (K₂CO₃ + Methionine), Ind. & Eng. Chem. Res. 49(2010)3910-3918
 - VLE-data for the system CO_2 -H₂O-K₂CO₃-Methionine at 80-120°C
 - It corresponds to loaded solutions of potassium salt of methionine
 - Some precipitation in experiments at 80°C with 2.5 molal $\rm K_2CO_3$ and 0.83 molal methionine
- Sengeløv L., Bachelor Thesis, Technical University of Denmark, 2010
 - Freezing point depression data for potassium salt of methionine and for loaded solutions of this salt



Results for the CO_2 -H₂O-K₂CO₃-Methionine system (Equilibrium constant from Sharma et al.)



 $m(K_2CO_3) \approx 1.27 \text{ mol/(kg H}_2O) \& m(Met) \approx 0.42 \text{ mol/(kg H}_2O)$

Measurements of freezing point depressions for mixtures of 1:1 (mole ratio) aqueous Met-KOH and 1:3 (mole ratio) aqueous Met-K₂CO₃



Freezing point depressions: Results and model correlation (Equilibrium constant from Sharma *et al.*)



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The potassium salt of methionine as an absorbent in postcombustion contra MEA, MDEA and K₂CO₃

Desorber 10000 1 molal absorbent, 115 °C 2 molal absorbent, 115 °C conditions 1400 9000 Potassium salt of Partial pressure of CO₂ in kPa 8000 1200 Partial pressure of CO₂ in kPa Potassium salt of methionine methionine 7000 MDEA 1000 Equilibrium MDEA 6000 - MEA constant from 800 -MEA 5000 Potassiumcarbonate Sharma *et al.* 600 4000 Potassiumcarbonate 3000 400 2000 200 1000 0 n 0.1 0.2 0.3 0 0.2 0.3 0.4 0.5 0 0.1 0.4 0.5 CO₂ loading CO₂ loading 1 molal absorbent, 115 °C 3330 2 molal absorbent, 115 °C 4330 Heat of reaction in kJ/(kg CO₂ captured) Heat of reaction in kJ/(kg CO₂ captured) ---- Potassium salt of — Potassium salt of 3830 methionine 2830 methionine ---- MEA ----MEA 3330 2330 2830 1830 Potassiumcarbonate 2330 Potassiumcarbonate 1830 1330 1330 830 830 330 330 0 0.2 0.4 0.6 0.8 1 0 0.2 0.4 0.6 0.8 CO₂ loading CO₂ loading

The potassium salt of methionine as an absorbent in postcombustion contra MEA, MDEA and K₂CO₃



Conclusion I

- The extreme difference between the properties of the potassium salt of methionine at absorber conditions and at desorber conditions indicates:
 - A strong temperature dependence of the equilibrium constant of methionine
 - Acidic properties of methionine strongest at desorber temperature
- Portugal et al. found that the carbon dioxide solubility in potassium glycinate did not change much in the temperature range investigated (20-50°C)

Conclusion II

- Due to the limited amount of experimental data available, it is possible to model CO₂ solubility in amino acid salt solutions without taking carbamate formation into account
- The potassium salt of methionine seems to have very interesting properties for post-combustion usage. However VLE-data at absorber column conditions are needed to validate the results
- The limited solubility and risk of precipitation provide challenges in the use of the potassium salt of methionine and must be investigated further
 - The 2 molal solutions for which calculations were shown here might be precipitating!
- More research is required to clarify the role of methionine in the absorption process: Especially regarding carbamate formation and kinetics
- Much more research is required on amino acid salt solutions. They have very interesting properties and could be real alternatives to alkanolamines.