

CHARACTERISTICS STUDY OF CO₂ INTO AQUEOUS POTASSIUM AMINO ACID SALT SOLUTION BY NMR SPECTROSCOPY

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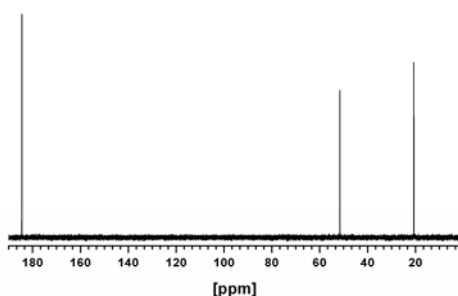
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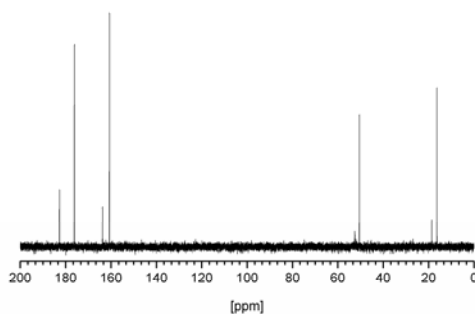
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Rapid industrialization and industrial activities since the industrial revolution have resulted in a sharp increase of greenhouse gas concentration in the atmosphere and are causing global warming. CO₂ generated by combustion of fossil fuels consists of around 80% of total greenhouse gas emissions and is considered the most important of the 6 major greenhouse gases. To reduce greenhouse gases and cope with the climate change problem at its root, the development of CO₂ capture technology is critical [1]. The wet absorption process, which is a post-combustion CO₂ capture technology using liquid absorbent, has been studied extensively in many countries for its ability to process high concentrations of CO₂ using retrofits of the existing process. The most important factor in the CO₂ absorption process is the selection of absorbent as this will determine the performance of the process. Generally, alkanolamine absorbents are effectively used to remove acid gas. In this study, the aqueous amino acid salt solutions were investigated as one of the carbon dioxide absorbent. The advantages of amino acid salts are their high resistance to degradation, low vapor pressure, and small amounts of oxidative degradation products [2]. The proton in carboxyl group of amino acid is substituted with metal ion by neutralization. The CO₂ loadings of potassium salt of amino acid were higher than those of sodium and lithium salt of amino acid [3]. The absorption capacity of the aqueous potassium salts of the L-alanine was measured and compared with MEA (monoethanolamine) and DEA (diethanolamine). The equilibrium partial pressure of CO₂ ($P_{CO_2}^*$) and absorption rate were measured by using VLE (Vapor-Liquid Equilibrium) equipment. The experiment conditions were temperature at 298, 313 and 353K and concentration of 1 mol·L⁻¹. The result of CO₂ loading high absorption capacity and absorption rate were faster than that of amine absorbent. Though good absorbent was considered by

many variables, absorption rate and capacity was more important factor. The CO₂ loaded solutions of MEA, DEA and aqueous potassium salts of the L-alanine solution were prepared by VLE apparatus at 333K. ¹H NMR and ¹³C NMR spectra were obtained for these systems at 293 K. This study shows that it is possible to use this method to confirm the reaction mechanism of aqueous potassium salts of the L-alanine solution with CO₂.



(a)



(b)

Figure 1. ¹³C spectra of potassium salt of L-alanine compared carbon dioxide before (a) and after (b) the reaction

References

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