

# **Heat of reaction for CO<sub>2</sub> absorption using aqueous K<sub>2</sub>CO<sub>3</sub> solution with homopiperazine**

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## **Abstract**

Carbon dioxide (CO<sub>2</sub>) is emitted from the consumption of fossil fuels and CO<sub>2</sub> emission is directly caused by the phenomenon of global warming. Coal-fired power plant is considered as a major source of CO<sub>2</sub> emission and it takes about 32~40 % of the total emission. The separation and capture of CO<sub>2</sub> from power plant is gaining interest as a method to reduce greenhouse gas emissions. Various methods for CO<sub>2</sub> separation and capture are having been developed, and some commercialized technologies for post-combustion having been applied to natural gas industry for 60 years. However, the technical improvements, such as high CO<sub>2</sub> loading and low generation energy should be achieved. Reaction heat can be explained by the summation of sensible heat, evaporation heat and heat of CO<sub>2</sub> absorption reaction. Since the heat of absorption reaction is considered as the energy needed to desorb CO<sub>2</sub> from rich absorbent. The CO<sub>2</sub> desorption energy and binding energy of each solvent have almost the same value. As for the heat of reaction, the enthalpy of the standard state( $\Delta H^\circ$ , heat of reaction) was obtained by calculating the heat of reaction generated per one mole of CO<sub>2</sub>.

In this work, the heat of reaction between absorbent and CO<sub>2</sub> was measured using differential reaction calorimeter (DRC, SETARAM Co. Ltd.) for comparing with generation efficiency of absorbent at higher than atmospheric pressure and refers to the entire quantity of heat generated from reactions between solvents and CO<sub>2</sub>. In order to

measure the total amount of CO<sub>2</sub> dissolved in solution, CO<sub>2</sub> consumption were analyzed by gas chromatography (GC). Aqueous absorbents were tested with more commonly used material, such as Monoethanolamine (MEA) and Potassium carbonate (K<sub>2</sub>CO<sub>3</sub>). Homopiperazine is used as a promoter of K<sub>2</sub>CO<sub>3</sub> solution to prevent crystalline formation and increase absorption capacity of K<sub>2</sub>CO<sub>3</sub> absorbent. The experimental results are compared with heat of reaction and values derived from solubility measurements. In solubility of CO<sub>2</sub>, the CO<sub>2</sub> loading capacity (mol-CO<sub>2</sub>/mol-solvent) of MEA and K<sub>2</sub>CO<sub>3</sub> + homopiperazine are shown approximately 0.5 and 0.73 at 313K. Also, in our experiment the reaction heat of K<sub>2</sub>CO<sub>3</sub> + homopiperazine was shown to be lower than MEA solution. On the other hand, homopiperazine was shown to have the highest loading capacity and heat of reaction. Therefore, K<sub>2</sub>CO<sub>3</sub> + homopiperazine is to be more favorable in the case of MEA. The results will be presented to comparison with MEA and an interpretation of the behavior of the K<sub>2</sub>CO<sub>3</sub> + homopiperazine. The experimental results are compared with respect to heat of reaction and values derived from CO<sub>2</sub> loading capacity.

Table 1. Overall heat of reaction of various absorbents at 313K.

Absorbent	Loading (mol CO <sub>2</sub> /mol solvent)	-Δh (kJ / mol CO <sub>2</sub> )	-Δh (kJ / g CO <sub>2</sub> )
MEA 30 wt%	0.54	82.38	1.87
MDEA 30 wt%	0.59	44.65	1.01
K <sub>2</sub> CO <sub>3</sub> 15 wt%	0.79	27.81	0.63
HomoPZ 10 wt%	1.04	94.58	2.15
K <sub>2</sub> CO <sub>3</sub> 15 wt% + HomoPZ 10 wt%	0.73	70.77	1.61