

CO₂ Capture from Flue Gases by Absorption Using Ionic Liquid

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ABSTRACT: In this study, to evaluate the performance of 1-butyl-3-methylimidazolium acetate ([BMIM][AcO]) as a solvent for CO₂ absorption process in bubble column reactor, the absorption capacity and CO₂ removal efficiency were investigated under different operating conditions, including solvent concentration of 0.05-0.1 M, CO₂ inlet concentration of 10000-30000 ppm, and gas flow rate of 0.5-2 L/min. The results show that when the CO₂ concentration was increased from 10000 to 30000 ppm, the removal efficiency and absorption capacity decreased from 87% to 55% and from 9.8 to 4.8 gCO₂/kg solvent, respectively. The CO₂ removal efficiency increased from 55 to 98% when the gas flow rate was reduced from 2 to 0.5 L/min. Similarly, CO₂ removal was improved by increasing the solvent concentration from 0.05 to 0.1 M. The results of this study show that 1-butyl-3-methylimidazolium acetate ([BMIM][AcO]) can be used as a solvent to remove CO₂ from flue gases with high removal efficiency and absorption capacity.

KEYWORDS: Absorption, Ionic Liquid, CO₂ capture, Global warming

I. INTRODUCTION

Human activity has recently resulted in an increase in the average air temperature, creating an important environmental concern. In order to reduce greenhouse gas emissions and associated adverse impacts, such as acidic rain and global warming, many researchers have carried out experiments. Among the greenhouse gases, carbon dioxide is the primarily responsible for the global warming. Therefore, providing effective CO₂ capture techniques has been recognized as a successful climate change strategy. There are three different approaches applied to capture CO₂: pre-combustion, oxy-fuel combustion, and post-combustion. Post-combustion CO₂ capture, one of these approaches, attracted the most interest due to the advantages it

provides, including outstanding efficiency for the separation of CO₂ and low partial pressure of CO₂ in gas stream. Up to date, many post-combustion carbon dioxide capture methods such as absorption [1], adsorption [2], cryogenics [3], and membrane technology [4] have been suggested.

Absorption of CO₂ by physical and chemical absorbents was a common industrial method among the methods applied in post-combustion. Between them, amine solutions were attracted considerable attention by numerous researchers. The amine solutions widely used for CO₂ absorption are monoethanolamine (MEA) [5, 6], diethanolamine (DEA) [7], methyldiethanolamine (MDEA) [8], 2-amino-2-methyl-1-propanol (AMP) [9], and piperazine (PZ) [10]. The first-generation solvent, such as Monoethanolamine (MEA), has a number of challenges, including a limited capacity to capture CO₂, high volatility, equipment corrosion, and negative environmental effects. Although these alkanolamine-based systems are very effective at capturing acid gas, they have a number of disadvantages, including the uptake of water into the recovered acid gas stream, amine loss during reclamation due to their volatility, and amine degradation by small amounts of sulfur-containing compounds. The usage of the alkanolamines at high concentrations is limited by these issues. A new kind of solvent such as ionic liquids has been used as an acid gas capture reagent in recent research studies. It has been shown that 1-butyl-3-methylimidazolium acetate ([BMIM][AcO]), presents very high ability for the absorption of acid gases [11].

In this work, we focus on the CO₂ absorption performance of the 1-Butyl-3-methylimidazolium Acetate ([BMIM][AcO]). Experiments were carried out using a Bubble column reactor, where the gas is continuously flowing and the liquid is a batch system. The effects

of initial CO₂ concentration, gas flow rate and ionic liquid concentration on the CO₂ capture performance were determined. The absorption capacity (g CO₂/kg ([BMIM][AcO])) and CO₂ removal efficiency were also calculated.

II. EXPERIMENTAL DETAILS

2.1 Calculation of Absorption Capacity and Removal Efficiency

The total amount of CO₂ absorbed by the solution is calculated by the area above the CO₂-time profile graph. The overall gas flow rate and the inlet concentration of CO₂ allowed us to calculate the CO₂ input flow rate. The CO₂ flow rate at the exit can also be calculated by using the gas flow rate and the measured CO₂ concentration at the effluent gas. The detail of the calculation can be found elsewhere [5].

The CO₂ removal efficiency was determined by the difference between gas

concentrations, read by CO₂ analyzer, at the input and the output of bubble column reactor (Eq.1).

$$\text{CO}_2 \text{ Removal Efficiency} = \frac{C_i - C_e}{C_i} \times 100 \quad (1)$$

where C_i is the CO₂ concentration (ppm) of the inlet gas stream, C_e is the CO₂ concentration (ppm) of the effluent gas stream.

2.2 Materials

In this work, 1-butyl-3-methylimidazolium acetate ([BMIM][AcO]) (Sigma-Aldrich) was used as a solvent. Carbon dioxide CO₂ (> 99.95 %, 150 bar, Oksan Gas, Turkey) and Nitrogen N₂ (> 99.99 %, 200 bar, Oksan Gas, Turkey) were mixed and used to simulate the flue gases. The molecular structure of the [BMIM][AcO] which has molecular weight of 198.26 g/mol is shown in Fig.1. In the experimental studies, the solvent concentration was adjusted to be 0.05, 0.08 and 0.1 Molar with distilled water which obtained by ultrapure water device (Thermoscientific, Germany).

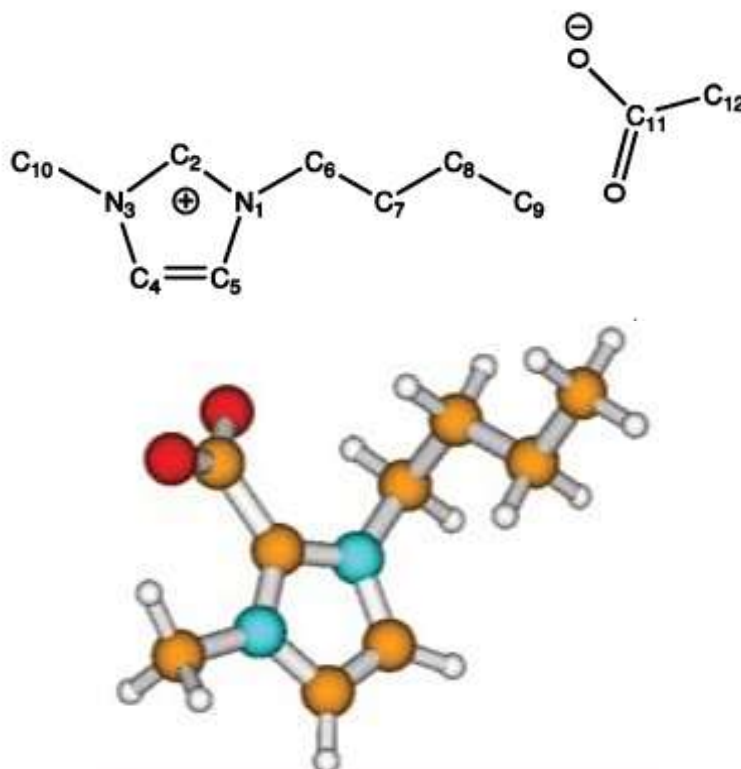


Fig. 1: The molecular structure of the [BMIM][AcO] [12]

2.3 Experimental Set-Up

Bubble column reactor used in this study was a cylindrical vessel with a gas diffuser at the bottom. The gas was dispersed into the liquid phase in the form of bubbles, thus providing high contact areas. Bubble column reactor was preferred over other reactors due to its simple structure, absence of mechanical moving parts, easy use, cost efficiency,

high heat and mass transfer rates, and low maintenance requirements. The reactor can be seen from Fig. 2.

The bubble column reactor used in this study was designed and manufactured from plexiglass by a local company. The column is 40 cm high and 5 cm in diameter. To simulate the flue gas, a mixture of CO₂ and N₂ gases was prepared using a

mass flow controller (MKS Multi Gas Controller 647B, USA). The gas mixture was fed into the solvent through the diffuser placed at the bottom of the reactor. 0.5 L of solvent was pumped into the reactor at the beginning of the experiment and was not changed during the experiment. The gas mixture was continuously fed into the reactor, and the gas

flow rate and CO₂ concentration were measured at the inlet and outlet of the reactor via mass flow meters and CO₂ analyzer (Vernier, Labquest 2, USA). The gas mixture leaving the reactor passed through the dehumidifier before entering the CO₂ analyzer. The experimental set up used in the experiments is shown in Figures 3 and 4.

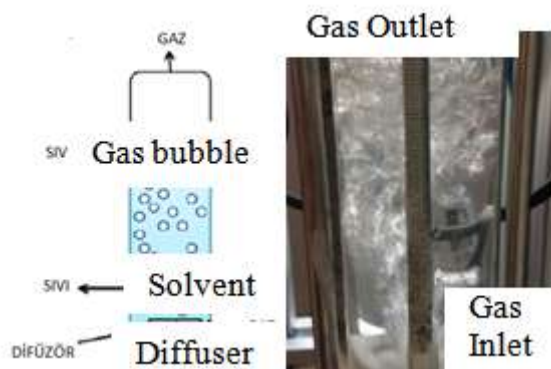


Fig. 2: Bubble column reactor



Fig. 3: Experimental set-up

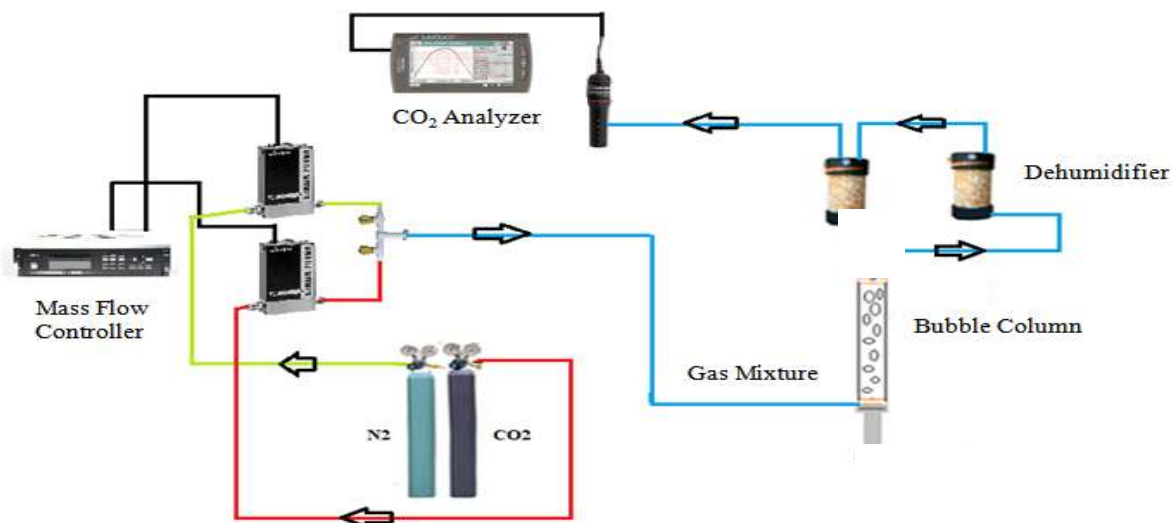


Fig. 4: Flow chart of experimental set-up.

III. RESULTS AND DISCUSSION

3.1 Effect of CO₂ Concentration

The initial CO₂ concentration changes the amount of pollutant loaded into the solvent per unit volume of gas. The change in CO₂ concentration directly affects the CO₂ holding capacity and CO₂ removal efficiency. The effect of CO₂ concentration (C_{CO_2}) was search using CO₂ concentrations of 10000 ppm, 20000 ppm and 30000ppm at gas flow rate (G) of 2.0L/min and solvent concentration ($C_{[bmim][Ac]}$) of 0.05M. The effects of initial CO₂

concentration on CO₂ effluent concentration, removal efficiency and absorption capacity (g CO₂/g solvent), are given in Figures 5 and 6.

As seen in Figures 5 and 6, CO₂ removal decreased when the initial CO₂ concentration increased from 10000 ppm to 30000 ppm. As the CO₂ concentration increased, the solution reached the saturation in a shorter time and the CO₂ holding capacity was full. The results show consistency when compared with the literature [13, 14].

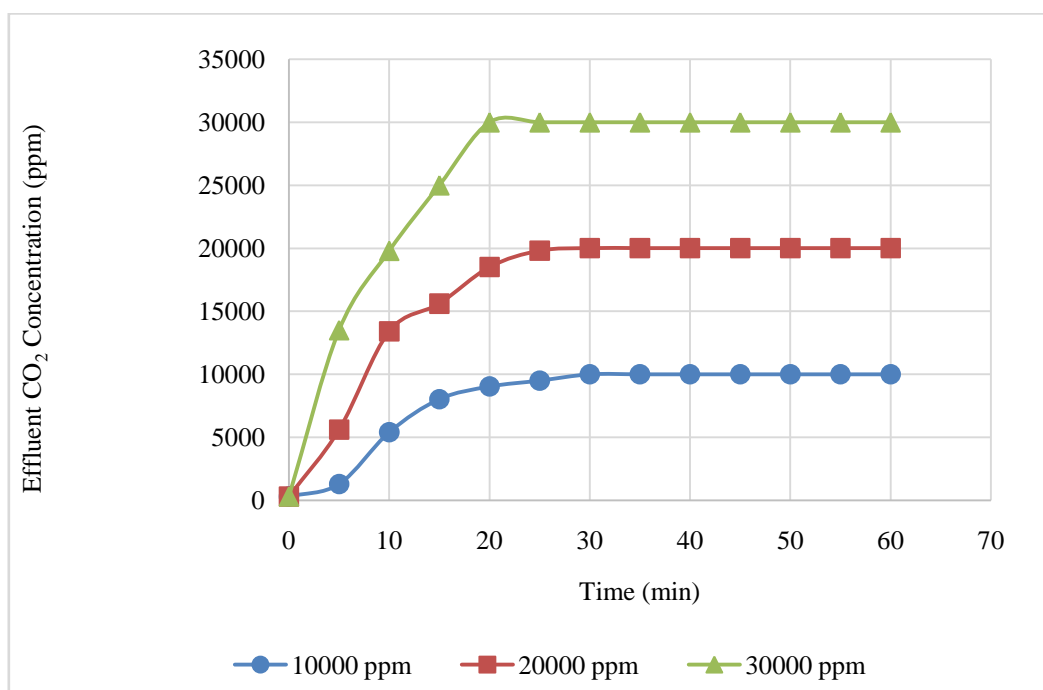


Fig. 5: The changes of effluent CO₂ concentrations with different initial CO₂ concentrations ($C_{[bmim][Ac]}$:0.05 M, G: 2.0 L/min)

As seen in Fig.6, the CO₂ removal efficiency and absorption capacity decreased with the increase of the CO₂ concentration from 10000 ppm to 30000ppm. The removal efficiency increased from the 55 to 87% when the initial CO₂ concentration decreased from 30000 ppm to 10000

ppm. It is well known in the absorption operation that a lower CO₂ concentration led to a higher CO₂ removal efficiency. The absorption capacity of 9,79 decreased to 4,81 gCO₂/kg solvent with the increase in the CO₂ concentration.

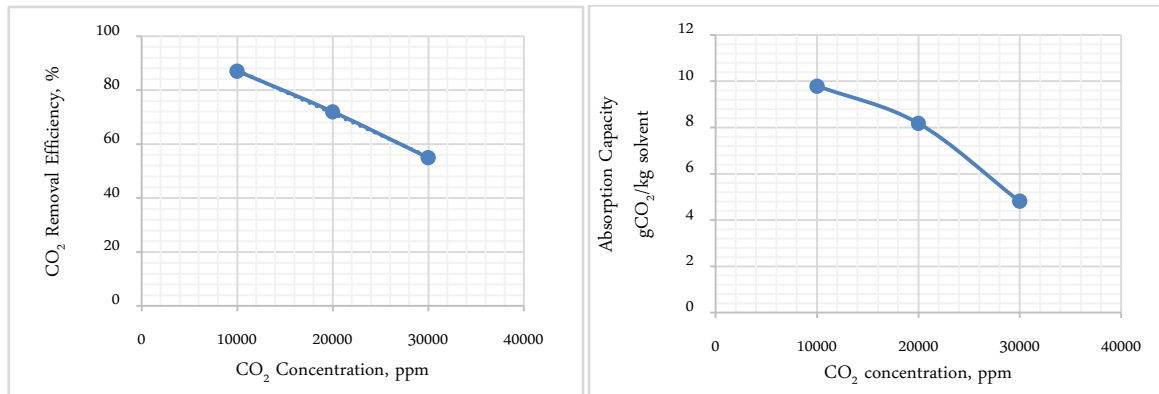


Fig. 6: The effect of initial CO₂ concentration on the absorption capacity and CO₂ removal efficiency

3.2 Effect of Gas Flow Rate

Gas flow rate which conducts with an L/G ratio is an important parameter that affects the CO₂ holding capacity. As the gas flow rate increases, the amount of CO₂ fed to the unit solvent volume per unit time increases. The effect of gas flow rate was searched using three different gas flow rates, 0.5, 1

and 2 L/min at 0.05M solvent concentration and 30000ppm CO₂ concentration. The solvent volume was kept constant as 500mL throughout the experiment, but the feed gas flow rate was increased. The results can be seen from Figures 7 and 8.

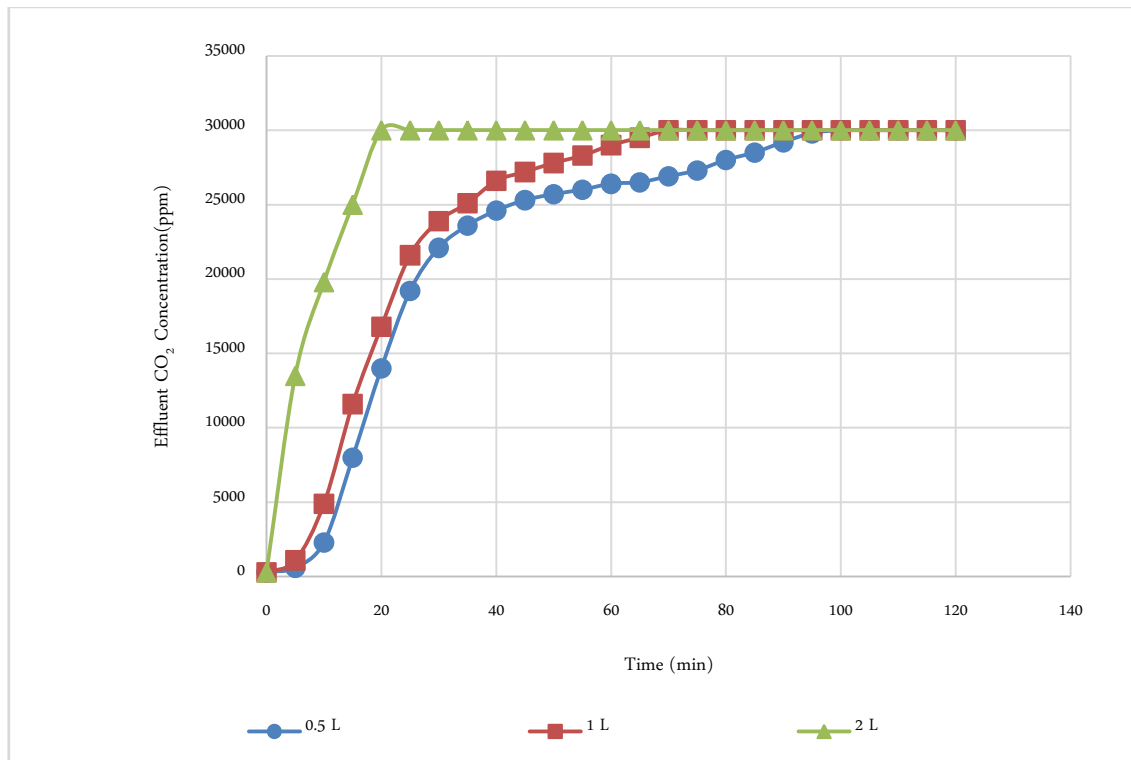


Fig. 7: Effect of gas flow rate on the effluent CO₂ concentration ($C_{[bmim][Ac]}:0.05\text{ M}$, $C_{CO_2}: 30000\text{ppm}$)

As can be seen from Fig. 7, the CO₂ holding capacity of the solvent was rapidly consumed with increasing gas flow rate. When the gas with a concentration of 30000 ppm enters the reactor at a rate of 0.5L/min, it took 100 minutes for

the run of the CO₂ holding capacity of the solvent out, while the solvent holding capacity was exhausted within 20 minutes when it was fed at a rate of 2 L/min.

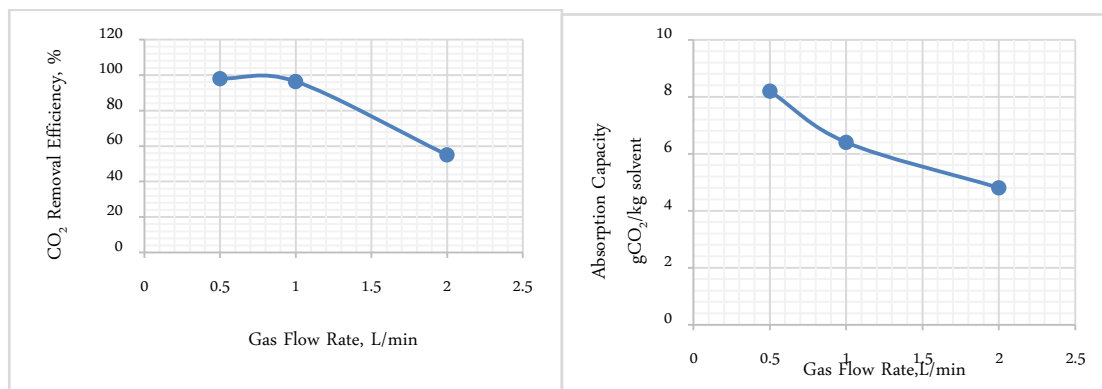


Fig. 8: Effect of gas flow rate on the absorption capacity and effluent CO₂ removal efficiency ($C_{[bmim][Ac]}$: 0.05 M, C_{CO_2} : 30000ppm)

As seen from Fig. 8, with the increasing gas flow rate, the CO₂ removal efficiency decreased. The removal efficiency increased from 55% to 98% when the gas flow rate decreased from 2 L/min to 0.5 L/min. In general, as the gas flow increases the amount of CO₂ molecules available for the absorption increases which leads to a higher mass transfer flux. Additionally, the gas flow rate determines the detention time of the gas component and effects the mixing regime of the solution. By increasing gas flow rate, the gas-film coefficient increases. This can be explained by the fact that increased gas velocity results in a reduction in film thickness as well as an increase in interfacial turbulence, both of which can improve solute transport from gas to liquid [15]. On the other hand, the amount of CO₂ passed through the system was increased and the contact time between gas and liquid was reduced, these two facts led to negative effects on CO₂ capture efficiency [16]. Therefore, increasing the gas flow rate has a limited influence on the overall mass transfer coefficient [17]. Based on the data obtained in this study, it can be concluded that the negative effects on CO₂ capture efficiency outweighed the positive effects, resulting in a decrease in absorption capacity. A similar trend has been report by Kuntz and Aroonwilas [18] and

by Qi et al. [14] for the chemical absorption of CO₂ into MEA solutions.

3.3 Effect of Solvent Concentration

The effect of solvent concentration was searched using three different [BMIM][AcO] concentration at 30000ppm CO₂ concentration and 2 L/min gas flow rate. Results can be seen from Figures 9 and 10.

Concentration of [BMIM][AcO] increases absorption capacity because mass transfer coefficient typically increases as absorbent concentration does. By increasing solvent concentration, a greater amount of active absorbent is available to diffuse towards the gas-liquid interface and react with CO₂. Therefore, increasing the [BMIM][AcO] amount in the solvent increases the chemical reaction. With increasing solvent concentration, CO₂ removal efficiency increases as well. Similar results are also found by the various researchers [19]. The removal efficiency does not double when the solvent concentration double because the relationship between the mass transfer coefficient and the rise in solvent concentration is not totally linear. On the other hand, the viscosity of the solvent is increased by the high solvent concentrations. Therefore, optimization is necessary when determining the solvent concentration.

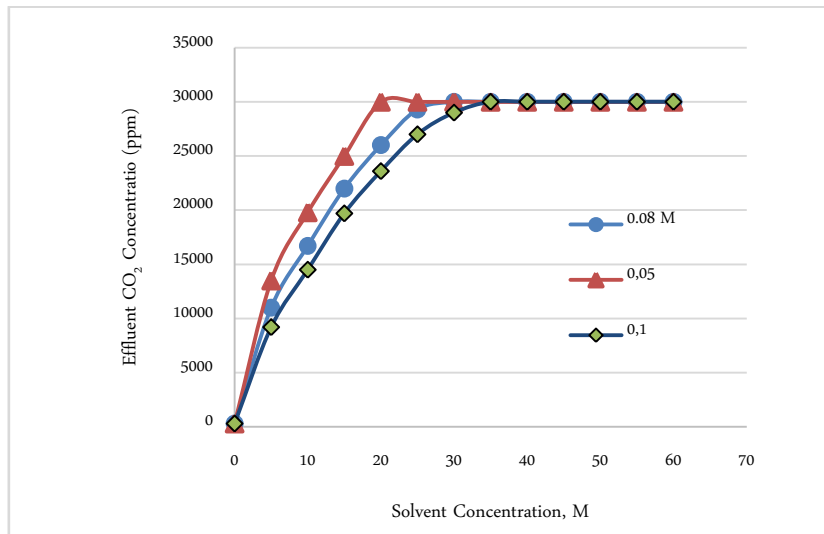


Fig.9: Effect of solvent concentration on effluent CO₂ concentration (G:2L/min, C_{CO₂}: 30000ppm)

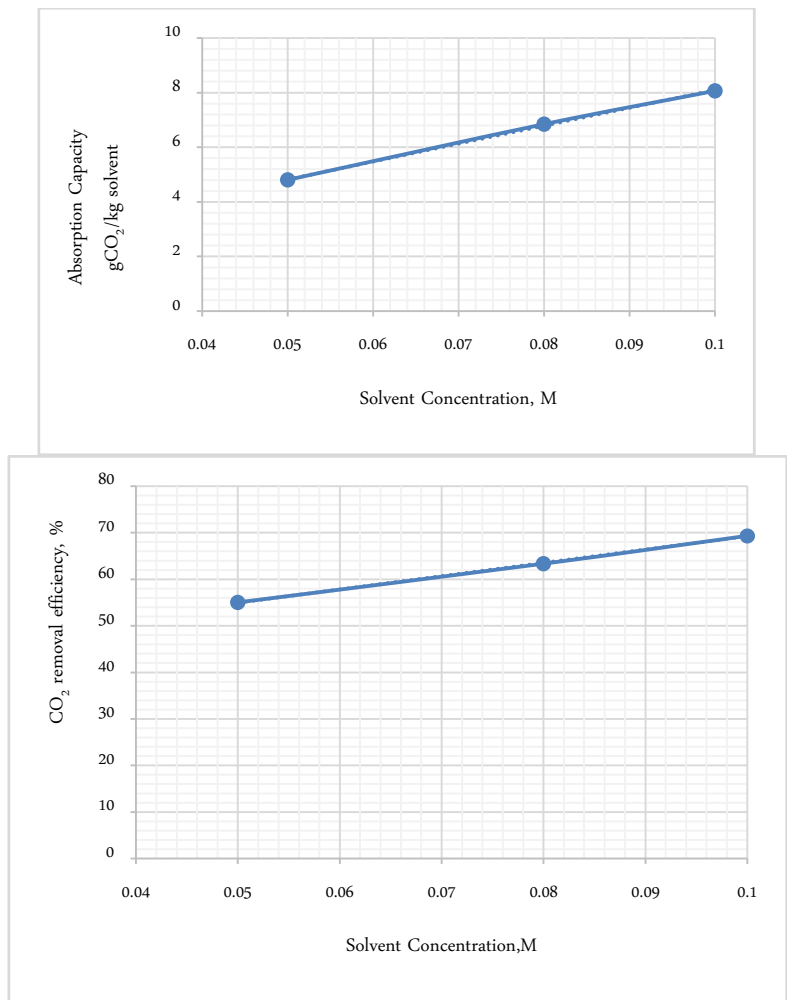


Fig. 10. Effect of solvent concentration on the absorption capacity and the CO₂ removal efficiency

IV. CONCLUSION

In this work, the absorption of CO₂ into aqueous 1-butyl-3-methylimidazolium acetate ([BMIM][AcO]) solution has been used to CO₂ capture from flue gas. The bubble column reactor was used to determine the effects of several operational parameters such as inlet CO₂ concentration (10000-30000 ppm), gas flow rate (0.5-2 L/min) and solvent concentration (0.05-0.1 M) on the absorption capacity and removal efficiency. The following results were obtained.

* CO₂ removal efficiency and absorption capacity increased from 55% to 87% and from 4.8 to 9.8 gCO₂/kg solvent, respectively, when CO₂ concentration was increased from 10000 to 30000 ppm.

*CO₂ removal efficiency increased from 55 to 98% when the gas flow rate was reduced from 2 to 0.5 L/min. Similarly, CO₂ removal was improved by increasing the solvent concentration from 0.05 to 0.1 M.

The results of this study show that 1-butyl-3-methylimidazolium acetate ([BMIM][AcO]) can be used as a solvent to remove CO₂ from flue gases with high removal efficiency and absorption capacity.

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