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CO₂ Capture by Dry Alkanolamines and an Efficient Microwave Regeneration Process

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Removal of acidic gases such as H₂S and CO₂ is performed during the purification of raw natural gas, most commonly using amine gas treatment. However, this industrially entrenched method is limited by significant shortcomings including low operational capture efficiency, amine pipeline corrosion and a large energy penalty due to the sorbent regeneration process. To address these shortcomings, we have studied the use of perfluorinated silica-stabilized dry alkanolamines (DAₙ) for CO₂ capture. Due to their micronized liquid domains, DAₙ display high operational CO₂ capture efficiency. Further, to minimize energy requirements for sorbent regeneration, microwave-assisted regeneration of the spent DAₙ sorbent was also studied and shown to decrease the energy requirements by about ten times. In contrast to very recent work, our results show that the use of DAₙ exhibits extraordinary recyclability, with negligible decrease in absorption capacity over at least ten absorption-regeneration cycles, indicating the potential of this material for gas treatment applications.

1 Introduction
Carbon dioxide capture is crucial not only for mitigating greenhouse gas emissions but also for natural gas processing. The recent United States shale gas boom has precipitated a rush in the production of natural gas, making the need for effective CO₂ capture technologies more pressing than ever. Examples of solid sorbents for CO₂ capture include metal-organic frameworks, porous polymers and zeolites. Such sorbents have the advantage of a high surface area to accelerate CO₂ capture. However, liquid sorbents, which often require agitation to achieve high gas-liquid interfacial areas, are still predominant in industrial processes. In fact, amongst various CO₂ capture technologies, the most established is amine gas treatment, which relies on the use of alkanolamines to separate acidic gases such as H₂S and CO₂ from raw natural gas. Despite amine-based gas sweetening being an already well-established industrial process, it faces multiple shortcomings including low operational CO₂ capture efficiency, amine corrosivity and a significant energy penalty due to the sorbent regeneration process. Although primary and secondary alkanolamines react with CO₂ in a 2:1 mole ratio to form their corresponding carbamates with a theoretical CO₂ absorption capacity of 50 mol% (Scheme 1), industrial systems utilizing alkanolamines typically only operate at 15 to 30 mol% capture efficiency. In addition, the high viscosity and corrosivity of alkanolamines necessitate their dilution with water to approximately 30 wt.% of amines. This leads to associated energy penalty during sorbent regeneration, where CO₂-loaded alkanolamine sorbents are heated to regenerate amines, since the additional water content increases the overall heat capacity of the sorbent. To address this problem, researchers have turned to utilizing ionic liquid-alkanolamine blends or to amine-functionalized porous solids for CO₂ capture. Dawson et al. recently reported an alternative approach whereby they used hydrophobic silica-stabilized dry alkanolamines (DAₙ) for CO₂ capture. However, the DAₙ exhibited low recyclability and substantial sorbent loss during regeneration, as well as significantly decreased CO₂ uptake after the first cycle. Nevertheless, this approach circumvents the need for dilution of the sorbent with water, thereby decreasing the energy penalty incurred during sorbent regeneration.

Amine-related corrosion problems are also potentially reduced because the amine is isolated by the silica particles from pipelines.

Scheme 1. Reaction of (a) monoethanolamine (MEA) and (b) diethanolamine (DEA) with CO₂ to form the corresponding carbamate salts.
For dry liquid formation to occur, the stabilizing particles must be hydrophobic towards the encapsulated liquid, otherwise a foam or paste may ensue. Generally, for a flat surface to be hydrophobic towards a liquid (contact angle > 90°), the solid surface tension γS must be γS ≪ γL/4, where γL is the surface tension of the liquid. It has recently been demonstrated that the formation of dry oils with liquids of decreasing surface tension requires the use of nanoparticles with increasing degrees of fluorination on their surfaces. Based on the surface tensions of MEA and DEA (Table S1), it can be deduced that oleophobic perfluoroalkyl-functionalized nanoparticles are preferable over oleophilic hydrocarbon-functionalyzed nanoparticles for stabilization of DA (Figure S1).

Experimental

Materials

All materials were purchased from commercial sources and used as received. 1H,1H,2H,2H-perfluorooctyltriethoxysilane (purity 97%) was purchased from Fluorochem. Ethanolamine (purity 98%), diethanolamine (purity 98%) and fumed silica powder with an average particle size (aggregate) of 0.2-0.3 µm (product number S5505) were purchased from Sigma Aldrich. Ethanol (purity 99%) was purchased from International Scientific. Aqueous ammonia (28%) was purchased from Malayan Acid Works. Compressed CO2 (purity 99.8%) was purchased from Singapore Oxygen Air Liquide.

Methods

Synthesis of perfluoroalkyl-functionalized oleophobic SiO2 nanoparticles

10 g of fumed SiO2 was added to a well-mixed solution of 220 mL ethanol, 11 mL 28% aqueous NH3 and 0.5 mL of 1H,1H,2H,2H-perfluorooctyltriethoxysilane, which was then homogenized (using an IKA T18 basic Ultra Turrax digital homogenizer) for 30 min until a homogeneous suspension was obtained. The resulting dispersion was stirred at 400 ± 50 rpm at room temperature overnight, after which the dispersion was heated on a hot plate to evaporate the solvents and subsequently dried in an oven at 80 °C for 1-2 days to collect the dry perfluorinated silica powder. Thermogravimetric analysis (TGA) was performed using a TGA Q500 v6.7 Build 203 instrument, from room temperature to 800 °C at a rate of 20 °C/min with 40 mL/min of nitrogen gas as balance gas and 60 mL/min of air as sample gas. TGA of perfluorinated silica versus unmodified fumed silica particles showed that the perfluorocarbon groups comprised approximately 35 wt.% of the functionalized silica (Figure S2). Dynamic Light Scattering (DLS) studies were performed with a Brookhaven ZetaPlus Zeta Potential Analyzer, to determine the size distribution of
unmodified fumed silica particles versus perfluorinated fumed silica particles after dispersing them in ethanol. DLS studies showed that the average aggregate size of perfluorinated silica particles was smaller than that of unmodified fumed silica particles (282 nm vs 367 nm, see Table S2).

Preparation of dry alkanolamines

5 g of perfluorinated SiO$_2$ nanoparticles was placed in a plastic beaker while 5 mL of the selected alkanolamine (pre-heated 1 hr. at 100 °C to minimize the water content) was slowly added into the beaker and blended with the particles using a simple coffee whisk (Figure S3a) at a rate of 500–1800 rpm for 1 min or until formation of a free-flowing powder.

Absorption of CO$_2$ into bulk or dry alkanolamines

The reaction was performed at room temperature and pressure. 5 mL of alkanolamine or approximately 10 g of DA$_f$ powder (containing 5 mL of alkanolamine) was put into a round bottomed flask placed in a water bath to modulate the heating effects arising from the exothermic nature of the reaction. CO$_2$ gas was bubbled at a flow rate of 500 ± 50 mL/min through the flask with the outlet connected to a gas bubbler until the absorption profile plateaued. At short time intervals of 30 sec, 1 min, the flask was taken out of the water bath, dried weighed on a weighing balance to record the increase in mass. The maximum CO$_2$ uptake by alkanolamines (in the absence of water) was calculated by assuming a reaction mole ratio of 2:1 alkanolamine to CO$_2$.

Regeneration of sorbents by conventional hotplate heating

The CO$_2$-loaded alkanolamine or DA$_f$ powder was regenerated by bubbling argon gas at a flow rate of 500 ± 50 mL/min through the sorbent while the flask was heated in an oil bath at 120 °C to release the absorbed CO$_2$. The flask was equipped with a reflux condenser maintained at 15 °C to minimize the evaporative loss of the sorbents and the outlet was connected to a gas bubbler.

Results and discussion

Brunauer-Emmett-Teller (BET) analysis performed on perfluorinated SiO$_2$ particles showed that CO$_2$ adsorption on these particles at room temperature was negligible (Figure 2), because 1 g of perfluorinated SiO$_2$ nanoparticles absorbed approximately 6 mg of CO$_2$ at atmospheric pressure while 1 g of the DA$_f$ (containing 0.48 g of perfluorinated silica) absorbed approximately 100 mg of CO$_2$. Of this 100 mg, about 3% is attributed to adsorption by silica. Hence, the mass increase is largely due to CO$_2$ uptake by the alkanolamine.

Carbon dioxide absorption results for both DA$_f$ and bulk liquid are shown in Figure 3. The theoretical maximum alkanolamine CO$_2$ uptake (in the absence of water) was calculated by assuming a reaction mole ratio of 2:1 alkanolamine to CO$_2$. Hence, 1 g MEA theoretically absorbs about 0.36 g CO$_2$ at maximum, and 1 g DEA theoretically absorbs 0.21 g CO$_2$. Based on Figure 3, CO$_2$ absorption by dry and bulk DEA in the first 10 min of CO$_2$ exposure reached 98 and 60 wt% of the theoretical maximum absorption respectively. Similarly, dry and bulk DEA achieved about 90 and 35 wt% of the theoretical maximum within 20 min. Mass changes due to replacement of air in the reaction vessel headspace with CO$_2$ were taken into account in the calculations.

Figure 2. Isothermal CO$_2$ adsorption at 22 °C by perfluorinated SiO$_2$ nanoparticles showing negligible CO$_2$ uptake (0.135 mmol g$^{-1}$ at 100 kPa).

For both amines, the DA$_f$ showed significantly higher CO$_2$ absorption than their corresponding bulk liquid counterparts (Figure 3). The lower absolute absorption of the bulk amines is due to mass transport effects arising from the large increase in the liquid viscosity caused by carbamate salt formation, which was verified by measuring the viscosities of the alkanolamine liquids before and after CO$_2$ absorption (Table S3). The effect of viscosity of a liquid on the mass transfer kinetics is described by the Stokes-Einstein equation:

\[ D = \frac{kT}{6\pi\eta r} \]
\[ D_{ij} = \frac{kT}{h_{SE} \eta_j R_i} \left( \frac{R_j}{R_i} \right) \]

(1)

**Figure 3.** Results of the absorption profile of CO\(_2\) into bulk alkanolamine or DA for (a) MEA (b) DEA at room temperature and pressure.

**Figure 4.** (a) and (b) TG-MS graphs of CO\(_2\)-loaded bulk DEA, (c) and (d) TG-MS graphs of bulk DEA without CO\(_2\) loading. (a) The black line shows a decrease of 18.92% of the initial sample mass for CO\(_2\)-loaded bulk DEA with a concomitant increase in sample temperature from 30 to 120 °C as depicted by the blue line, (b) graph corresponding to (a), depicting ion current detection of both CO\(_2\) and H\(_2\)O. (c) The black line shows a decrease of 16.03% of the initial sample mass for bulk DEA, (d) graph corresponding to (c) depicting detection of only H\(_2\)O but not CO\(_2\).

In this equation, \(D_{ij}\) is the diffusion coefficient of solute \(i\) in pure solvent \(j\), \(k\) is the Boltzmann constant, \(T\) is the temperature, \(h_{SE}\) is the Stokes-Einstein number, \(\eta_j\) is the solvent viscosity, \(R_i\) is the radius of the solute molecule and \(R_j\) is the radius of the solvent molecule. According to Equation (1), the diffusion coefficient of CO\(_2\) gas in the amine sorbent decreases with increasing solvent viscosity. Further, given that the system studied is stationary and unagitated, increased
viscosity also significantly affects the replenishing of unreacted amine near the gas-liquid interface, which is dependent upon movement of alkanolamine molecules towards the interface via convection currents or diffusion.

However, due to the small domain sizes within DA\textsubscript{f} (approximate diameter of a microscopic liquid marble is 80±30 µm, as measured by cryo-SEM), the required distance of CO\textsubscript{2} diffusion within the droplets to contact unreacted alkanolamine is significantly less than for the bulk liquid. This helps to explain why the DA\textsubscript{f} achieve a much larger gas absorption relative to the theoretical capacity within the timeframe of the experiment than do the bulk amines. In addition, the reason for the slightly higher absorption capacity for dry MEA compared to dry DEA is due to the inherently lower viscosity of the former compared to the latter.

Based on the work of Wang et al.,\textsuperscript{31} who postulated that the higher methane gas hydrate formation rate in dry versus bulk water was due to the higher surface area to volume ratio in dry water, and taking the view that the rate of gas uptake is dependent upon the gas-liquid interfacial area, we expect that the CO\textsubscript{2} absorption rate by DA\textsubscript{f} would follow a similar trend (estimated surface area to volume ratio in DA\textsubscript{f} is 75,000 m\textsuperscript{-1} based on volume of alkanolamine present). Nevertheless, the gas-liquid interfacial area for the microscopic liquid droplets is reduced by the presence of the silica shell around each droplet; CO\textsubscript{2} must diffuse across this porous shell to react with the encapsulated alkanolamines.

Our CO\textsubscript{2} uptake studies showed that for MEA, the absorption rate by the bulk liquid in the first minute was faster than that by dry MEA (Figure S4). This is presumably because the lower liquid viscosity of MEA compared with DEA means that CO\textsubscript{2} mass transfer effects do not outweigh the effect of the silica barrier initially. However, as carbamate formation proceeds, the viscosity of the liquid mixture increases (Table S3), and mass transfer effects become more important. The rate of CO\textsubscript{2} uptake therefore becomes faster for dry over bulk MEA. For DEA, dry DEA exhibits faster CO\textsubscript{2} absorption than bulk DEA throughout the entire absorption cycle (Figure S4), because the mass transfer rate in the more viscous DEA liquid was much slower and hence the effect of higher surface area to volume ratio of dry DEA was more prominent.

Since the carbamate salts decompose in the temperature range of 100-150 °C,\textsuperscript{35,36} the CO\textsubscript{2}-loaded bulk alkanolamines were heated to 120 °C to release the absorbed CO\textsubscript{2} gas and regenerate the alkanolamines. The decrease in mass was attributed to both CO\textsubscript{2} and H\textsubscript{2}O release, which was verified by thermogravimetric mass spectroscopy (TG-MS) analysis (see Figure 4).

For each sample of bulk alkanolamine and DA\textsubscript{f}, three cycles of absorption and desorption of CO\textsubscript{2} were performed (Figure 5). Based on Figure 5, DEA appears to be more recyclable than MEA, with negligible change in CO\textsubscript{2} uptake capacity over three cycles. This is because secondary amines like DEA form weaker bonds with CO\textsubscript{2} than primary amines like MEA, allowing for easier regeneration of the free amines.\textsuperscript{37} This is supported in the findings of McCann et al., where the enthalpy of carbamate formation for MEA and DEA was found to be −29.7 kJ/mol and −23.7 kJ/mol respectively.\textsuperscript{38} Moreover, MEA has a lower boiling point than DEA (170 °C vs 271 °C) and a much higher vapour pressure, leading to a greater degree of sorbent loss per cycle.

As heating is essential for the sorbent regeneration process, it is imperative to maximize the heat transfer efficiency within the amine sorbents. This is dependent on the three main mechanisms of heat transfer, namely conduction, convection, and radiation. In the case of DA\textsubscript{f}, these three mechanisms happen to be hindered by the silica shell encasing the micronized alkanolamine droplets and the air pockets throughout the dry liquids. For heat transfer via conduction, fumed silica particles exhibit low thermal conductivity due to the nanoscale pores among them in their fractal arrangement,\textsuperscript{39,40} causing the silica shell to block effective heat conduction from the heat source to the droplets. Importantly, the many air pockets between the microscopic particle-coated liquid marbles in DA\textsubscript{f} also limit heat conductivity. With regards to convection, the liquid in DA\textsubscript{f} exists as dispersed microscopic droplets, thereby limiting convective heat transfer. As for radiative heat transfer, Taylan and Berberoğlu have shown that the silica shell in dry water greatly attenuates infrared radiative heat transfer due to its high single-scattering albedo, i.e. a significantly larger proportion of heat radiation is scattered compared to that absorbed by the silica shell,\textsuperscript{41} thus preventing infrared radiative heat transfer from the heat source to the droplets.

![Graph](Image)
Figure 5. Comparison of solvent recyclability for (a) bulk MEA versus dry MEA, whereby heating lasts for 2 hr. for each cycle of regeneration and (b) bulk DEA versus dry DEA, whereby heating lasts for 1 hr. for each cycle of regeneration.

Since the three major forms of heat transfer are significantly inhibited for DA$_f$, we turned to microwave heating as an alternative method to improve the efficiency of heat transfer during the regeneration process. This method is dependent on the ability of the irradiated material, e.g. a solvent, to absorb microwave energy and convert it to heat, which is quantified by a dielectric parameter called the loss factor $\tan \delta$. This loss factor is expressed as:

$$\tan \delta = \frac{\varepsilon''}{\varepsilon'}$$  \hspace{1cm} (2)

where $\varepsilon''$ is the dielectric loss, which is indicative of the efficiency with which electromagnetic radiation is converted into heat, and $\varepsilon'$ is the dielectric constant describing the ability of molecules to be polarized by the electric field. A solvent with a high $\tan \delta$ value of more than 0.5 is required for efficient microwave absorption. DEA, with a high $\tan \delta$ value of ~ 0.6 at 2.45 GHz at 278 K can be rapidly heated to temperatures >100 °C within a minute when irradiated under microwave conditions.$^{42,43}$

Owing to the higher recyclability of DEA over MEA, we chose to study DEA for microwave heating-based regeneration. Samples of approximately 10 g of dry DEA containing 5 mL of liquid DEA were loaded with CO$_2$ until the absorption plateaued, and the CO$_2$-loaded samples were subsequently heated in a programmable microwave synthesizer at 120 °C for approximately 1 hr. until no further mass loss occurred. To test the recyclability of the sorbent and confirm that the observed mass loss was correctly attributed to CO$_2$ loss, 10 repeated cycles of CO$_2$ absorption and removal were carried out, as shown in Figure 6. The level of CO$_2$ uptake by dry DEA was found to be stable over at least 10 cycles, showing excellent recyclability of dry DEA. Similar regeneration results were obtained using a simple household microwave instrument, although the heating periods were limited to 10-20 sec in order to avoid overheating.

Regeneration of DA$_f$ via microwave heating was found to be more efficient than conventional hotplate heating under the same conditions (whereby no purge gas was used). Conventional hotplate heating took more than 3 hr and consumed about 0.38 kWh per cycle on average. Microwave heating on the other hand took about 1 hr and consumed about 0.034 kWh per cycle on average, which is less than 10% of the energy consumption by conventional hotplate heating.

Finally, the recyclability of the perfluorinated silica particles was investigated. Alkanolamines typically undergo degradation over multiple absorption and regeneration cycles after which they must be replaced.$^{44}$ Nevertheless, the perfluorinated silica particles could be separated from the amines by dispersion of DA$_f$ in ethanol and then centrifuged, and re-used in at least three subsequent batches of DA$_f$. This leads to significant cost-savings since only the alkanolamines need to be replaced.

Conclusions
In conclusion, the dry alkanolamines described herein demonstrate significantly enhanced performance with regards to CO₂ absorption capacity and sorbent recyclability. Furthermore, the novel dielectric heating technique for sorbent regeneration proposed here has proven to possess the benefits of reduced energy consumption and minimal use of purge gas. The perfluorinated silica particles can also be recycled and reused to prepare DA₆ with negligible change in their wettability and overall quality for at least three cycles. Moreover, as suggested by the recent work of Dawson et al., the amine component of the DA₆ is also isolated by the silica particles from pipelines and other corrosion sensitive materials, leading to further reduction in maintenance costs associated with adding corrosion inhibitors and repairing damage to pipelines. Last but not least, this novel method of using DA₆ is not restricted to MEA and DEA; in principle any dry liquid can be formed provided that nanoparticles of the appropriate surface energy are selected. The implications are that this idea can be extended to other liquid CO₂ sorbents, possibly leading to a similar improvement in absorption rate and capacity. However, there still exist several challenges for the application of DA₆ industrially. Firstly, the powdery nature of DA₆ will likely require significant changes in the current process technology and facilities specialized in handling liquid amine sorbents. Secondly, due to the nanoscale primary particle size of fumed silica, careful confinement of the powder needs to be put in place to prevent contamination in the outlet gas stream. Thirdly, stripper columns need to be modified to incorporate dielectric heating capabilities. Nevertheless, given the multiple advantages of utilizing DA₆ for CO₂ absorption and desorption, we believe that they should still be considered promising candidates for practical applications in CO₂ capture.

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Notes and references
Dry alkanolamines are shown here to be highly efficient CO$_2$ sorbents, recyclable via microwave regeneration.