

# **Compact, Low Energy CO<sub>2</sub> Management using Amine Solution in a Packed Bubble Column**

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## **Abstract**

A novel method for removing CO<sub>2</sub> from gas streams is described. The Carbon Dioxide Continuous Scrubber, CDOCS, takes advantage of the intimate liquid-gas contact afforded in a packed bubble column to both absorb the CO<sub>2</sub> from a gas stream, and to regenerate the solution. The design relations and performance of a prototype CDOCS system using amine solution are presented. Over a 30 day trial, 20 m<sup>3</sup>/hr of air was continuously scrubbed to 60-80 ppm. The CDOCS footprint is less than 0.18 m<sup>3</sup> with power consumption around 300 W. Applications for air scrubbing include alkaline fuel cells, small scale processing, and industrial safety. A concept is proposed to sweeten bio-gas from dairy farm effluent for on-farm electricity generation. While industrial processes involving packed or plate trickle columns are well known for CO<sub>2</sub> production and sweetening, these conventional methods are complex and expensive, and do not scale down to air scrubbing or to small scale biogas production.

Key Words: CO<sub>2</sub> Scrubbing, CO<sub>2</sub> Management, Amine, Packed Bubble Column

## 1. Introduction

The objective of this project was to develop a feasible technology solution for the problem of scrubbing CO<sub>2</sub> from the air supply of Alkaline Fuel Cells. Although AFC's have been successfully deployed in the space program for many years, terrestrial application has always been problematic because the acid gas, CO<sub>2</sub>, in atmospheric air reacts with the base liquid electrolyte to form precipitates which can degrade the electrode performance and reduce the activity of the electrolyte [1]. At the present time, researchers and system developers employ a solid absorbent, soda lime, to remove the CO<sub>2</sub> from the oxidizer stream. However, soda lime cannot be regenerated, must be replaced frequently, and adds considerable operating cost. Soda lime has never been considered as an option for commercial AFC systems. AFC stack materials and architecture have continued to develop but a method to continuously scrub CO<sub>2</sub> from air that meets the system requirements for AFC plants has not previously been found [2].

Several applications currently exist for managing CO<sub>2</sub> levels in air including submarines [3,4], space craft [5], diving re-breathers [6], and the operating theater [7]. We suggest that if a low-cost air scrubber were technically feasible, then new applications may include high CO<sub>2</sub> work environments like breweries and coal mines. CO<sub>2</sub> removal could also possibly reduce HVAC energy requirements in crowded environments like meeting rooms. Some people may even prefer to have a low CO<sub>2</sub> work or home environment with air pollution at pre-industrial levels.

CO<sub>2</sub> is separated from a wide range of gas flows in industrial processes. Combustion gas is stripped with aqueous amine solution in order to produce bottled gas and dry ice using the common industrial method of wet scrubbing. Packed bed or plate columns are used to contact a liquid adsorbent with the gas stream [8]. Typically, industrial wet scrubbers utilize a temperature or pressure swing process to separate CO<sub>2</sub> from petrochemicals, methane (natural gas sweetening), and other gas streams [9]. Industrial separation technology is complex and expensive, and thus economy of scale means that most wet scrubbing equipment is large scale. Using the height equivalent of theoretic plate (HETP) design process, a packed column to scrub CO<sub>2</sub> from air to below 10 ppm would approach an infinite number of theoretical stages.

Monoethanolamine (MEA) is the most widely employed solvent for CO<sub>2</sub> absorption, used for over seventy years in the CO<sub>2</sub> production and gas sweetening industries. Though new amines and amine mixes have been developed, MEA is still the preferred absorbent for low pressure and low concentration CO<sub>2</sub> absorption. The properties and behavior of amine solutions, while not entirely understood, have been empirically documented over time to provide engineers with a useful database for design [10]. Absorption occurs at temperatures up to approximately 60°C. The MEA – CO<sub>2</sub> reaction is exothermic and reversible by supplying heat to the system. The temperature swing absorption/evolution process reverses at approximately 70°C [11]. The basic MEA temperature swing chemistry is given in Equation 1.



The packed bubble column has not been reported in literature or industry as an apparatus for CO<sub>2</sub> absorption from air or other gas streams. It is well known that a bubble column provides excellent liquid/gas contact. A high surface area, low volume packing in the bubble column increases the gas hold-up, reduces geysering, and maintains a small bubble size [12].

The carbon dioxide continuous scrubber (CDOCS) system is a rather simple concept utilizing an MEA solution in a packed bubble column (PBC) as the CO<sub>2</sub> absorber, and a smaller PBC containing a heating element as the regenerator [13]. Air is scrubbed during bubbling through the absorber, and a flush gas is bubbled through the regenerator to carry away the regenerated CO<sub>2</sub> from the heated MEA solution. A low head flow, induced by the liquid lift from the air hold-up in the absorber PBC, is used to move solution from the top of the absorber, to the top of the regenerator, then return it to the bottom of the absorber.

Section 2 gives brief descriptions of the CDOCS system concept employed as an AFC air scrubber and for biogas sweetening. Section 3 outlines the operating theory and design relations for the CDOCS absorber and regenerator and the system integration dynamics. Section 4 describes the prototype system and the experiments performed. Section 5 gives the experimental results, with the conclusions following in Section 6.

## 2. CO<sub>2</sub> Management Applications

Current applications for CO<sub>2</sub> management by scrubbing with MEA exist in manned space flight, submarines and methane gas sweetening. CO<sub>2</sub> production also uses MEA solution in packed or plate columns to separate CO<sub>2</sub> from a combustion gas stream. The two future applications featured in this paper require small-scale, low-cost scrubbing. The relative simplicity of the dual packed bubble column DCOCS system is well suited to these emerging energy applications.

### 2.1 *Alkaline Fuel Cells*

The CDOCS was developed to specifically meet the requirements for a 5 kW AFC, but can be scaled for either smaller and larger systems. The air scrubber is positioned between the air blower and the humidification/pre-heating system and can be integrated with the air supply conditioning system to optimize space and power utilization [14]. A system schematic showing the operation of the CDOCS scrubber in the context of a typical AFC system is shown in Fig. 1. In this configuration, the previously scrubbed AFC exhaust gas is used as a source for the regenerator flush gas. Thermal integration includes use of AFC exhaust heat to pre-condition the incoming air. The incoming and outgoing regenerator flows pass through a counter flow heat exchanger to reduce heat energy input into the regenerator.

The CDOCS system prototype described in this paper was developed specifically to meet the system size, maintenance, and parasitic power requirements for a demonstration project. The demonstration system coupled a wind turbine, electrolyzer and AFC system to provide continuous power for a remote telecommunications facility. The system specifications limited parasitic power to 300 W and set a maintenance target of 30 days. The materials required for the CDOCS are also required to be compatible with the AFC.

### 2.2 *Small scale biogas sweetening*

Intensive animal husbandry and industrial agriculture improve economic performance in the agricultural industry, but generate large animal and plant processing

waste streams. In particular, swine and dairy farming have high localized effluent accumulation near housing sheds and milking barns. This kind of waste poses issues of odors, water pollution, and greenhouse gas emissions, particularly methane,  $\text{CH}_4$ , during decomposition. These agricultural waste streams must be treated to mitigate environmental impact, and the biogas energy potential has been investigated [15]. One of the negative energy aspects is transporting the waste to a large processing plant. On-farm biogas generation has been developing in recent years [16], but a major issue with biogas is the variable and possibly large portions of  $\text{CO}_2$  and moisture, and the presence of  $\text{H}_2\text{S}$ . Unless the biogas is used directly for heating, storage and use in an internal combustion engine for power generation are problematic due to corrosion and variability of the gas heating value and combustion characteristics. A low-cost method for separating the acid gases and moisture from the biogas, termed “sweetening”, could facilitate higher value uses and storage.

The schematic diagram in Fig. 2 is a concept of an on-farm biogas system with integrated sweetening. An MEA and glycol solution readily absorbs water vapor from the gas stream when the water in the solution is less than 30% wt. In an experiment with air and a solution with 10 wt% water, the solution absorbed 3 liters of water from  $1000\text{m}^3$  of atmospheric air. The absorbed water can be removed through the regenerator and either condensed and collected or exhausted as vapor. The absorption of  $\text{CO}_2$  is per kg solution is higher for higher MEA concentration, but water is required during regeneration or degradation of the MEA can occur. The concept biogas CDOCS system is integrated with the biogas energy system by using waste heat from boiler combustion or an engine to supply heat for the regenerator.

An important property of amine solutions is the absorption of acid gases, including the  $\text{H}_2\text{S}$  which will form  $\text{SO}_2$  upon combustion. The  $\text{H}_2\text{S}$  is regenerable from the solution with heat in the same process as the  $\text{CO}_2$  regeneration [17]. The anaerobic digester can be operated at higher pressure (2.5-3 bar) which suppresses ammonia production, but also increases the  $\text{CO}_2$  fraction [18]. This pressure is used in the concept system to drive the gas flow through the CDOCS system. Once the biogas is scrubbed, a compressor can be used to raise the pressure to a level suitable for storage and injection into an internal combustion engine or gas turbine generator.

### 3. Continuous Scrubber Design

This paper presents the results of research to determine design relationships and develop the system architecture for this novel application of a packed bubble column. The CDOCS system requires an injection gas pressure in the range of 300 mm H<sub>2</sub>O. The gas stream is injected into the base of the packed bubble column with a depth of absorbent solution set by the level of CO<sub>2</sub> removal required. The packing breaks up the air stream into small bubbles and creates a well mixed frothy liquid. Liquid lift caused by the gas hold-up is used to drive flow of the solution through a heated regenerator. The regenerator is also a packed bubble column with a small stream of flushing gas provided by a diaphragm pump. Corrosion and oxidation were found to be manageable due to the low temperature regeneration and stainless steel construction. The following sections describe the two key system components, the absorber and regenerator, and the system integration with an overview of the governing design and operating relations.

#### 3.1 Absorber

The CDOCS system makes novel use of a packed bubble column (PBC) for both CO<sub>2</sub> absorption and regeneration. While the PBC has been well known in Chemical and Process Engineering, it is not as widely used in industry as packed trickle columns. A PBC is a liquid reservoir filled to depth,  $h$ , with the absorbing liquid with chemical solubility for CO<sub>2</sub> component,  $\varepsilon_o$ . Air with CO<sub>2</sub> concentration,  $C_{amb}$ , is injected at the bottom of the PBC, is broken up into small bubbles by the packing material, and rises due to buoyancy. The residence time of the air flow,  $Q_a$ , in the liquid absorbent in a PBC of cross-section area,  $A$ , is given as  $t_r = hA/Q_a$ . The PBC has proved an effective means to remove CO<sub>2</sub> from air because of the high gas-liquid contact area and vigorous mixing [19].

The PBC absorber for a CDOCS system can be designed using the following relation for scrubbed CO<sub>2</sub> concentration:

$$C(t_r) = BC_{amb} \exp(-h_m A_p t_r) + \varepsilon_o C_{amb} \quad (2)$$

Where  $B$  is an experimentally determined entrance constant for a given solution,  $h_m$  is the bubble flow mass transfer coefficient, and  $A_p$  is a packing factor expressing a representative bubble cross section area. In previously reported investigations, the liquid depth for high efficiency scrubbing (98+% CO<sub>2</sub> removal) was determined to be 250-300 mm.

### 3.2 Regenerator

In our previous research we have shown that a high concentration aqueous solution (50% wt MEA) buffered with glycol (35% wt) can be regenerated at atmospheric pressure and temperatures as low as 70°C without boiling when a packed bubble column is used. The PBC regenerator performance was investigated with the major finding being that regeneration at 100°C in a PBC is faster than in a stirred beaker at 140°C or higher [20]. Nitrogen was used as the flush gas in the previous regeneration studies of fully CO<sub>2</sub> saturated solution. CO<sub>2</sub> was measured in the exhaust gas stream, and the regeneration rate was shown to increase with flush gas flow and with temperature. The solution was re-saturated with CO<sub>2</sub> by dropping in dry ice, and regenerated as many as ten times without loss of regeneration capacity or change in regeneration rate. Experimentally determined optimal regeneration conditions were used for the prototype system design, a regenerator heater temperature of 120°C, residence time of at least 10 minutes, and flush gas flow capable of producing a fully frothy condition.

### 3.3 System dynamics

The operating dynamics of the CDOCS system used to model the system are shown in Fig. 3. For continuous scrubbing, the rate of CO<sub>2</sub> removed from the regenerator must equal the rate of CO<sub>2</sub> scrubbed from the air stream. The rate of solution flow through the regenerator,  $Q_s$ , is the primary system control parameter for achieving continuous scrubbing performance. Performing a mole balance on CO<sub>2</sub>, the necessary flow can be related to the absorber efficiency,  $\eta_A$ , the regenerator efficiency,  $\eta_R$ , the mole rate of CO<sub>2</sub> influx to the scrubber with the air,  $\dot{c}$ , the mole fraction of MEA in the solution,  $f_{MEA}$ , and the CO<sub>2</sub> loading of the absorber solution,  $\alpha$ :

$$Q_s = 2 \frac{\eta_A}{\eta_B} \frac{\dot{c}}{\alpha_{MEA}} \frac{M_s}{\rho_s} \quad (3)$$

Where  $M_s$  and  $\rho_s$  are the solution molecular weight and density respectively.

#### 4. Experimental set-up

A prototype CDOCS system was constructed to evaluate long-term performance [21]. Tests have not yet been conducted on methane or biogas, but the chemistry is well known for this process, and the design relations and system performance for air should be applicable to other small-scale scrubbing applications. The regeneration temperature and the regenerator flow rate are the primary variables which affect the steady state performance of the system, as shown in Eqn. 3. The regenerator liquid flow was controlled with a manual valve and the regenerator element temperature was thermostatically controlled.

The experimental set-up is shown schematically in Fig. 4. The overall proportion of the prototype system is 0.49 m x 0.35 m x 0.6 m. The fill level of MEA solution in the absorber was 300 mm and determined via a sight glass. 17 liters of fresh solution (50% wt MEA, 35% wt glycol, 15% wt water) were placed in the absorber at the beginning of each experiment in the long-term testing. The dynamic liquid level of approximately 500mm is caused by the gas hold-up in the PBC during operation. This dynamic head is used to collect fluid at the top of the PBC and generate the gravitationally fed flow through the regenerator. The air-induced solution flow from the absorber through the regenerator was determined to be in the range of 4 liters/hour for steady operation.

During operation, some solution is absorbed into the coalescer material and escapes in the exhaust as vapor. The liquid level was maintained over the course of the investigation via a peristaltic pump from a solution sump. While the testing program was carried out using 20 m<sup>3</sup>/hr air flow, the prototype absorber, with dimensions 500 mm height and 260 mm diameter was capable of scrubbing well over 40 m<sup>3</sup>/hr air flow [21]. The long term tests used a lower flow rate because the low-cost blower would overheat if run at high power for long continuous periods.

The regenerator was constructed from stainless steel tube and fittings with dimensions 300 mm height and 22.5 mm diameter. The flush gas was atmospheric air



supplied by a diaphragm pump at a rate of 1 to 4 liters/hour and pre-heated by passing through a coil in an oil bath. The flush gas was cooled in a stainless steel coil condenser before passing through the sampling chamber of a Drager infrared CO<sub>2</sub> detector. The continuous flow through the regenerator was cooled by a fan and stainless steel radiator prior to being returned to the absorber. In this experiment, the regenerator was not insulated and 310 W was continuously supplied to the heating element.

Atmospheric air was supplied to the absorber by a blower at a rate determined from a float flow meter. The air injection at the base of the rolled stainless steel absorber was accomplished by a simple 8-port sparger from a swimming pool filter. The absorber and the regenerator were packed with commercially available 15 mm diameter stainless Pall rings. Wire mesh stainless mist eliminators were layered on top of the packing to collect droplets. A commercial Pall coalescer containing pleated paper-like filter eliminated mist from the scrubbed air outlet, and collected fluid drained back into the absorber.

A Shimadzu GC-9A Gas Chromatograph with Thermal Conductivity Detector was used to measure the CO<sub>2</sub> concentration in the processed air stream. The inlet air CO<sub>2</sub> concentration was measured with a Draeger Polytron Infra Red CO<sub>2</sub> detector, and atmospheric levels in our laboratory were found to be 380±5 ppm. The MEA concentration in the solution was monitored at regular intervals by titration of the solution with acid (0.1 molar HCl) and methyl red indicator. The acid reacts only with free MEA in the solution, not with MEA which is bound to CO<sub>2</sub> but still regenerable (carbamate). The solution pH was also monitored using an Orion Research 701A digital Ionalyzer.

## **5. Results**

The experimental prototype system was used to investigate the scrubbing performance relationships with several operational variations, including solution strength, solution depth, scrubbing solution addition and regenerator temperature over several extended periods of 20-30 days. Table 1 gives a summary of the results of the experiments with the CDOCS system. As a comparison, air was scrubbed with the regenerator heater switched off, and a linear increase in CO<sub>2</sub> in the scrubbed air was measured from 0 to 380 ppm over 12 days.

The regeneration temperature of 120°C provided good long term performance without risking solution degradation. Higher regenerator flush gas flow enhanced performance, but with the current prototype, 4 liters/min is the upper limit without causing operating issues with the fluid level at the top of the regenerator.

Using water to maintain the operating liquid level resulted in diluted solution over time and poor performance. Several MEA solutions were investigated for long term behavior. A slight improvement in performance may be gained by using the 60/25/15 solution. The hygroscopic nature of the 80/20/0 solution became apparent when 3 liters of water were absorbed from the first 1000m<sup>3</sup> of scrubbed air. The MEG-free solution had markedly reduced performance and a much higher rate of solution loss due to the higher solution vapor pressure.

Using the optimal operating conditions determined by the experimental test program, the CDOCS system was operated over a period of 32 days. The CO<sub>2</sub> concentration was reduced to less than 60ppm for the first 11,000 m<sup>3</sup> of scrubbed air, and to around 80 ppm for a further 3000 m<sup>3</sup> scrubbed air. A sudden darkening of the solution at 11,000 m<sup>3</sup> coincided with an increase in the rate of solution addition and a change in the rate of MEA wt% remaining in the solution as determined by the titration analysis. Over the whole test, the system required 10 liters of solution over the one month period, or 0.7 liters/1000 m<sup>3</sup> scrubbed air.

## **6. Conclusion**

A prototype carbon dioxide continuous scrubber (CDOCS) system has been developed which utilizes the packed bubble column (PBC) gas-liquid contactor for absorbing CO<sub>2</sub> from a high volume flow of air in a compact space, and for regenerating the absorbent solution at low temperature. The novel concept involves wet scrubbing with an aqueous solution of monoethanolamine (MEA) and glycol. The gas hold-up in the absorber is used to drive the flow through the regenerator, a smaller, heated PBC which is flushed with air. The PBC allows for a small footprint (0.1 m<sup>3</sup>) for scrubbing capability of at least 40 m<sup>3</sup>/hr.

The performance of the overall system is determined by the balance between absorption and regeneration. The prototype system was tested over one month of

continuous operation for scrubbing 20 m<sup>3</sup>/hr air. The air was scrubbed to less than 40ppm for the first four days of operation, to 60±20 ppm for the following 11 days, then to 80±10 ppm for the remainder of the 30 day test period. Over thirty days of operation, 15,000 m<sup>3</sup> of air was scrubbed to acceptable CO<sub>2</sub> concentration for alkaline fuel cells while consuming 12 kg of MEA and 310 W electric power for the regenerator heater, which is within the specifications for a 5 kW (output) AFC demonstration system.

## **7. Acknowledgements**

This work was supported by the Foundation for Research Science and Technology New Zealand by a Technology for Industry Fellowship, and by ASCO CO<sub>2</sub> Ltd. The authors wish to acknowledge the collaboration with Dr. Owen Curnow, Department of Chemistry, University of Canterbury, and Mr. Warren Holland of ASCO CO<sub>2</sub> Ltd. The contributions of other students in the Advanced Energy and Material Systems Laboratory have helped in the overall product development, Lee Liaw, Elizabeth Jenkins, Christopher Carl. Thanks to the AEMS lab technician, Ron Tinker and GC-TD technician Trevor Berry.

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## Figure Captions

**Figure 1.** Schematic diagram of an alkaline fuel cell power generation system with the CDOCS integrated into the air supply pre-conditioning, and utilizing AFC exhaust air as the regenerator flushing gas.

**Figure 2.** Conceptual schematic diagram of an on-farm biogas processing system with the CDOCS system providing fuel gas sweetening and de-watering prior to storage or injection into an engine.

**Figure 3.** Flow balance on the absorber and regenerator showing the definitions of the efficiencies for CO<sub>2</sub> stripping and for regeneration, respectively.

**Figure 4.** Schematic diagram of the CDOCS prototype system used in the experimental investigations.

**Figure 5.** Long term scrubbing performance of DCOCS system with measurements of the MEA solution strength and tracking of consumption of solution from the scrubber sump.

Figure 1

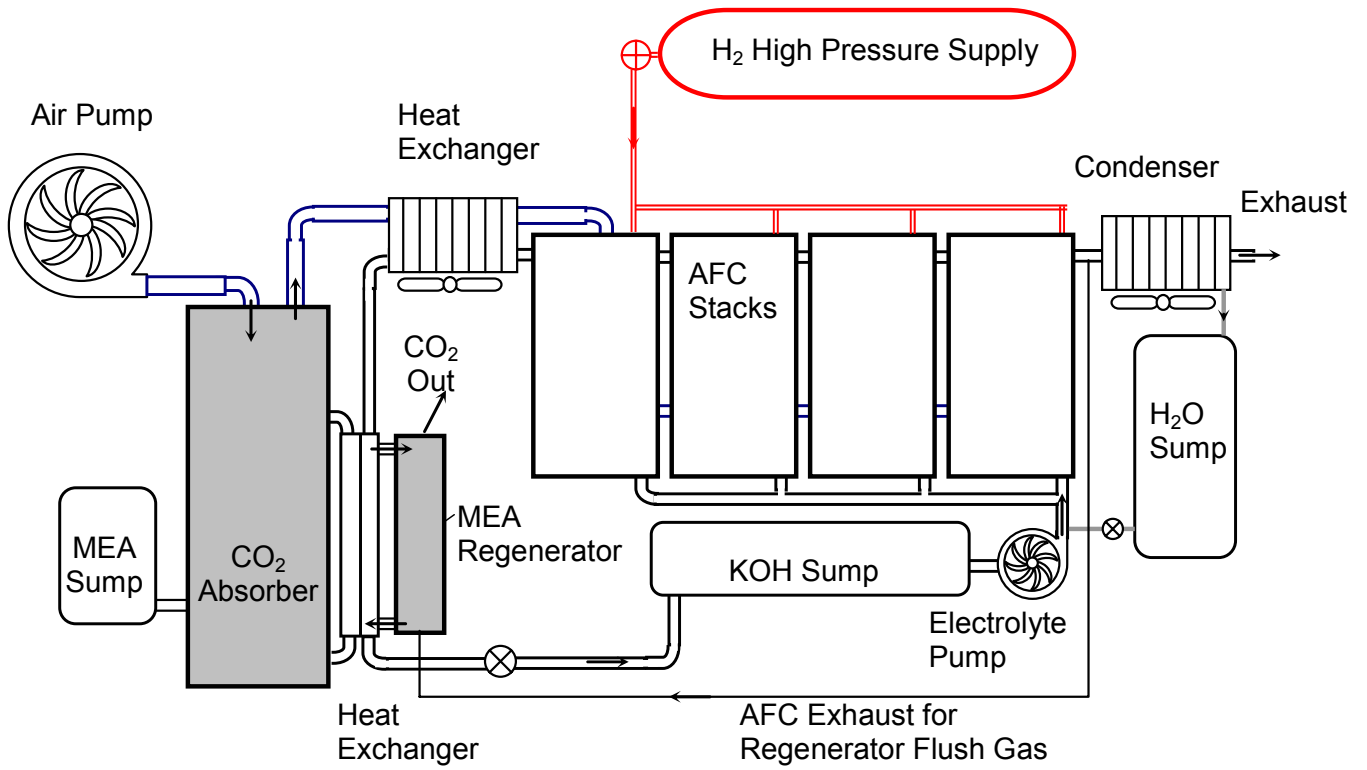


Figure 2

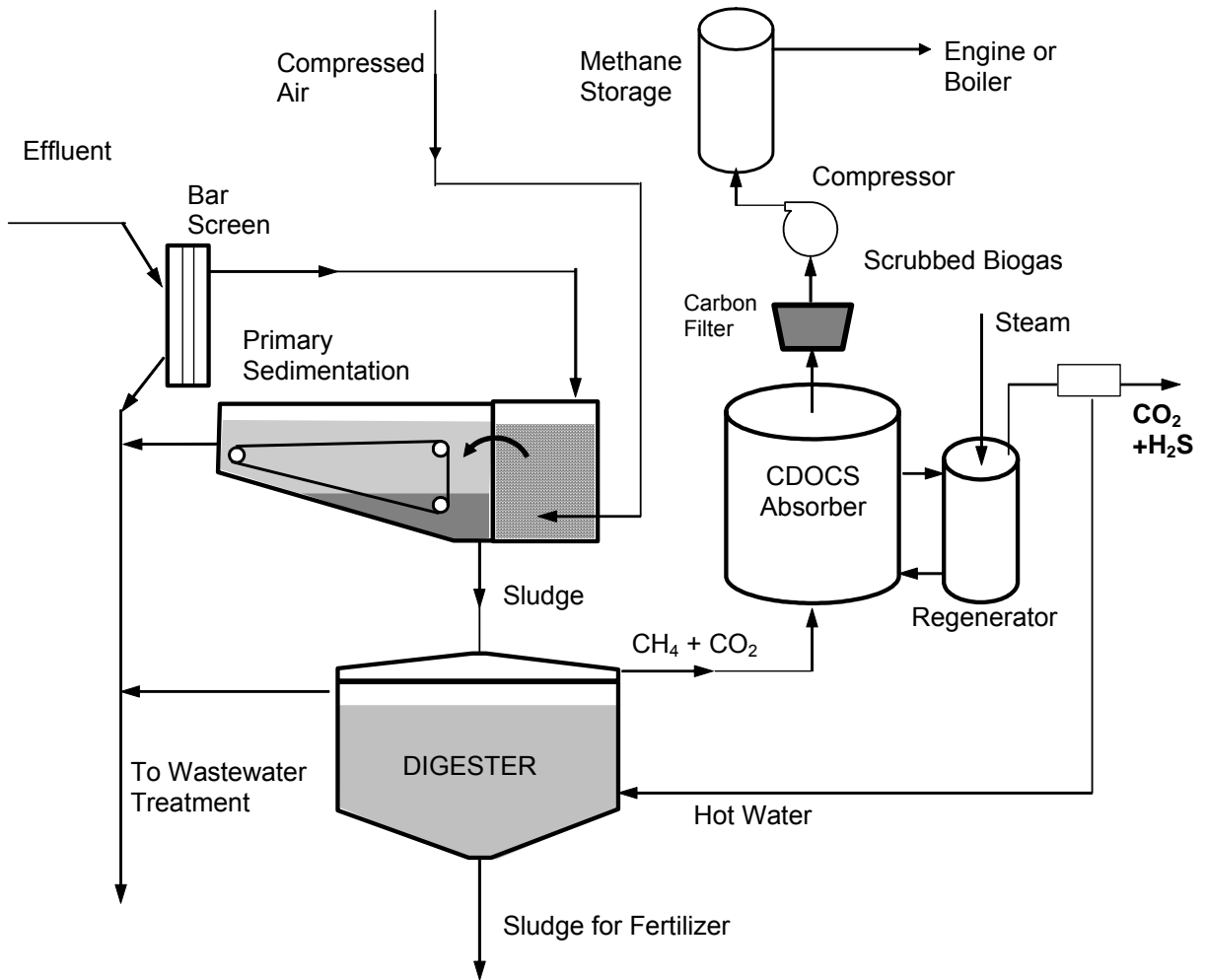




Figure 3

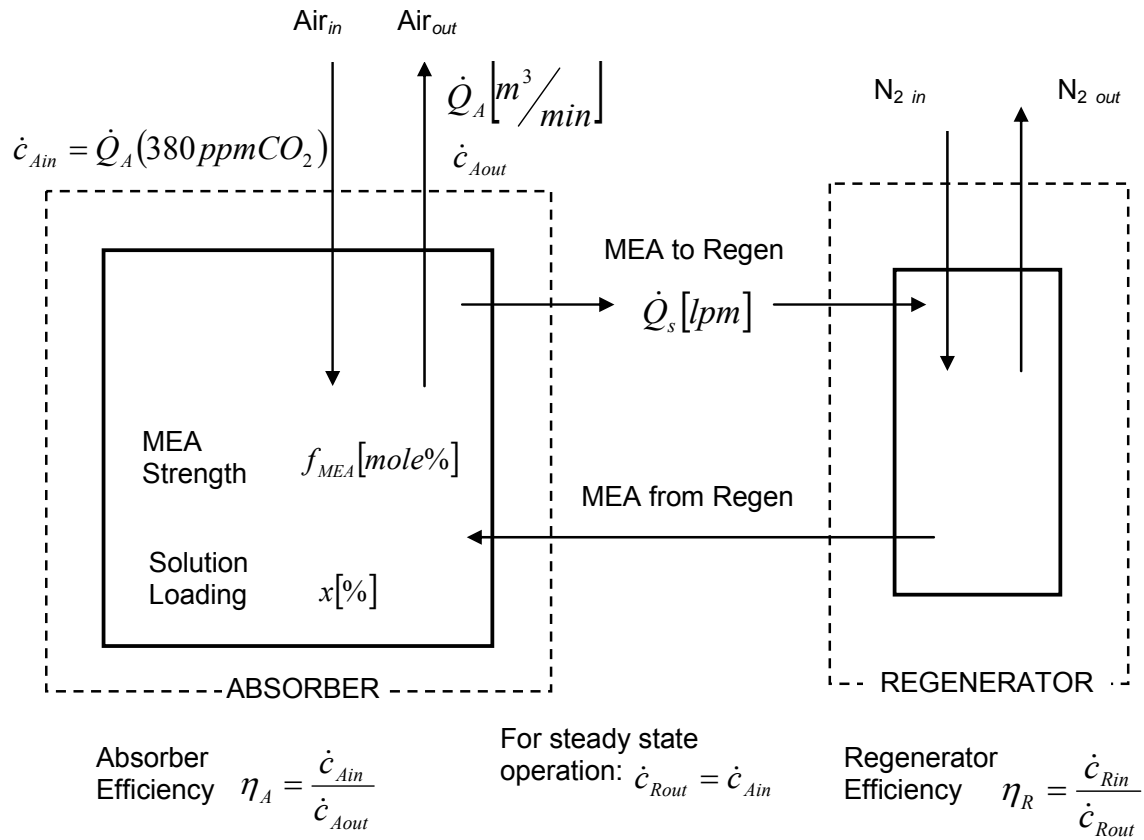


Figure 4

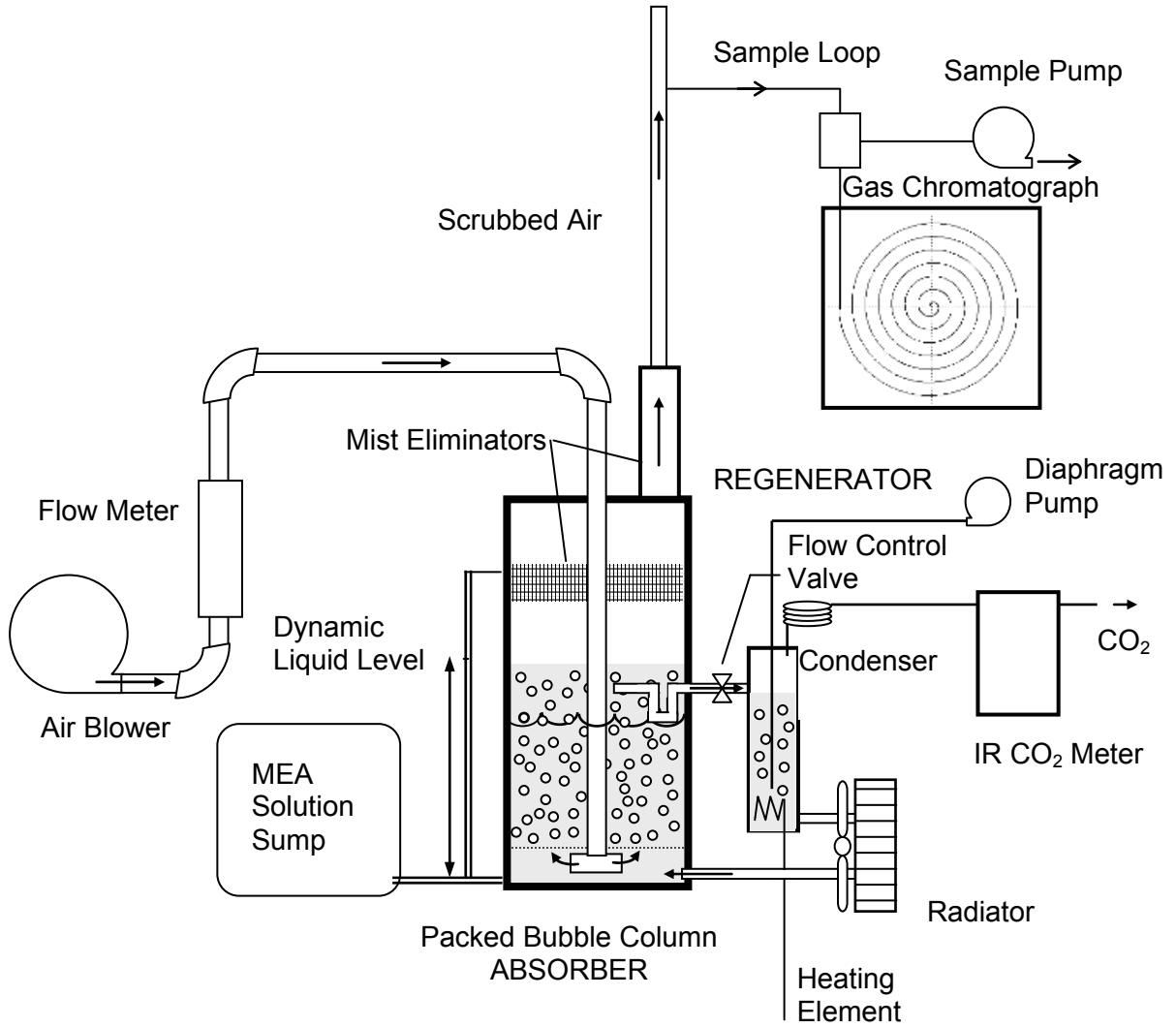
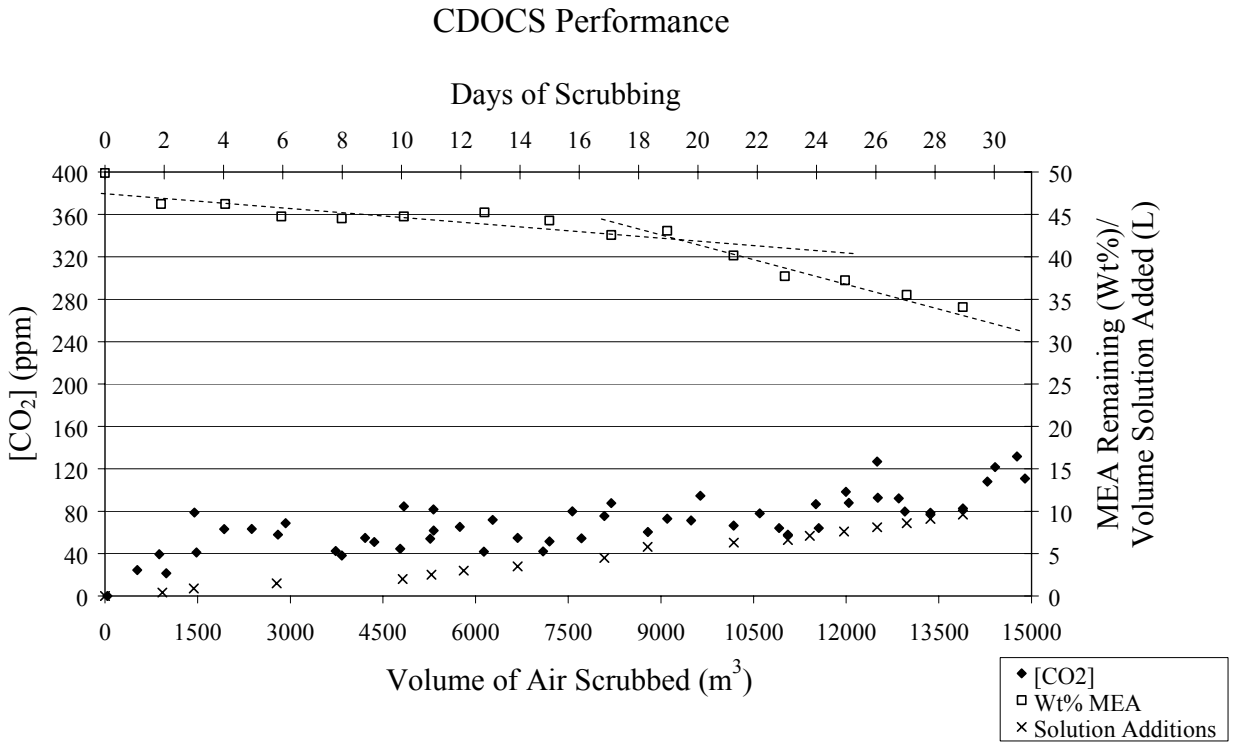


Figure 5



**Table 1.** Results of long term air scrubbing investigations, time to reach steady state operation, and the steady scrubbing performance, with effects of processing variables; Regeneration Temperature, Flush Gas Flow, Water and MEA solution additions.

| <b>Regen Temp.</b> | <b>MEA/MEG/H<sub>2</sub>O wt%</b> | <b>Flush Gas [l/min]</b> | <b>Water* Add Rate</b> | <b>Soln.* Add Rate</b> | <b>Time to Reach SS</b>       | <b>Steady State CO<sub>2</sub> ppm</b> |
|--------------------|-----------------------------------|--------------------------|------------------------|------------------------|-------------------------------|--|
| 100±10°C           | 50/35/15                          | 1.8                      | 2                      | 0                      | 9 days<br>4500m <sup>3</sup>  | 160 ±20                                |
| 120 °C             | 50/35/15                          | 4                        | 2.2                    | 0                      | 6 days<br>3000m <sup>3</sup>  | 120 ±20                                |
| 120 °C             | 50/35/15                          | 4                        | 0                      | 0.38                   | 4 days<br>2000m <sup>3</sup>  | 80 ±20                                 |
| 120 °C             | 60/25/15                          | 4                        | 0                      | 0.6                    | 4 days<br>2000m <sup>3</sup>  | 60 ±20                                 |
| 120 °C             | 80/20/0                           | 4                        | 0                      | 0                      | 12 days<br>5000m <sup>3</sup> | 120 ±20                                |
| 120 °C             | 50/0/50                           | 4                        | 0                      | 1.5                    | 4 days<br>2000m <sup>3</sup>  | 80 ±20                                 |
| → 120 °C           | 50/35/15                          | 4                        | 0                      | 0.7                    | 4 days<br>2000m <sup>3</sup>  | 60 ±20<br>→ 80 ±20                     |

\* Liquid additions measured in liters/1000<sup>3</sup> scrubbed air

→ Optimal system 30 day test: 60ppm scrubbing for 23 days, 80ppm thereafter.