

Direct Air Capture with Amino Acid Solvent: Operational Optimization using a Crossflow Air-Liquid Contactor¹

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Abstract

Direct air capture (DAC) is a negative emission technology for removing CO₂ from the atmosphere to maintain the CO₂ level within a reasonable range so as to address greenhouse effects. In this study, the operational optimization of lab-scale DAC has been investigated using a crossflow air-liquid contactor loaded with a three dimensionally printed Gyroid packing structure and a potassium sarcosinate solvent. The effects of various parameters, including feed air flow rate, liquid solvent flow rate, contactor geometry, and ambient temperature, are examined. The results demonstrate that the Gyroid packing design achieves comparable CO₂ capture performance to conventional packed beds but with a significantly lower pressure drop of up to 77.8%, suggesting its potential as an efficient and cost-effective solution for gas-liquid contactors in DAC. Additionally, the study explores the climate impact on CO₂ capture performance and finds that as the air temperature increases from 35°F to 95°F at fixed relative humidity at 80%, the CO₂ capture rate increased from 23.2 to 46.8% with better stability. The research highlights the importance of optimizing contactor design and operational conditions to improve the CO₂ capture rate and feasibility of DAC systems as a negative emission technology for addressing greenhouse effects.

Keywords: Direct Air Capture, CO₂ Capture Capacity, Contactor Design, Potassium Sarcosinate, Weather Condition

1. Introduction

As more nations pledge to achieve net zero emissions, policymakers are increasingly focusing on translating these commitments into tangible policy actions and strategies. Currently, only a limited number of countries and companies have developed detailed plans or pathways to reach their net zero goals[1]. Carbon capture, storage, and utilization (CCSU) are widely acknowledged as significant components for addressing global warming. To achieve significant reductions in atmospheric CO₂, a wide range of technologies are required, including capturing emissions from specific sources and removing CO₂ directly from the atmosphere[2-4]. Negative emissions technologies (NETs) play a crucial role in reducing CO₂ levels by extracting CO₂ from the atmosphere. Direct air capture (DAC), bioenergy with carbon capture and storage, ocean fertilization, and afforestation are four prominent approaches to achieving negative emissions[3, 4]. NETs, particularly

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DAC, are essential elements of a comprehensive response to climate change, offering diverse solutions to tackle the challenge. By promoting transparency and integrating carbon dioxide removal (CDR) into net zero strategies, policymakers can identify the technological, policy, and market needs within their jurisdictions while enhancing public understanding of these approaches.

DAC employs chemical processes to directly extract CO₂ from the atmospheric air. Although DAC shows promise in terms of scalability, modularity, and the ability to utilize arid land, it currently remains costly, considering the dollars required per tonne of CO₂ removed[5, 6]. Nevertheless, DAC offers notable advantages as a CDR option, including its ability to achieve high storage permanence when coupled with geological storage and its minimal impact on land and water resources[7]. Additionally, the captured CO₂ can be used, such as in food processing or in combination with hydrogen to generate synthetic fuels. As the transition to net zero emissions continues, the CO₂ used for synthetic fuel production would increasingly need to be sourced from sustainable bioenergy sources or directly extracted from the atmosphere. This approach is crucial in order to avoid delayed emissions from CO₂ derived from fossil fuels when the synthesized fuel is burned[8, 9]. Hence, DAC emerges as a viable option to achieve this objective.

Among the various DAC technologies, solvent-based absorption is considered one of the most advanced and well-established methods. In conventional solvent-based absorption systems, a solution of CO₂-absorbing solvent is introduced into a packed column, where it comes into contact with the dilute CO₂ present in the air. The gaseous CO₂ is absorbed into the liquid solvent, resulting in an outlet stream with reduced CO₂ concentration and a liquid stream that is rich in CO₂. To enhance the contact between the gas and liquid phases, structured packing is often employed in solvent-based systems, providing a large interfacial area for efficient interaction between the dilute CO₂ and the liquid solvent. The performance of CO₂ capture in solvent-based approaches largely relies on the characteristics of the packing material, which facilitates the contact and reaction between the dilute CO₂ and the solvent[10]. In previous studies, conventional structured packing designs such as stainless steel[11, 12] and PVC[13, 14] have been utilized for liquid solvent-based DAC systems although there have been limited advancements and changes. The capital cost associated with packing and the column itself typically constitutes around 75% of the total costs, which is primarily determined by the column size and the amount of packing[15]. However, advancements in structured packings have the potential to significantly reduce capital costs and energy requirements by minimizing the column size and packing volume[6, 16, 17]. Recent research has focused on exploring the use of advanced packing columns specifically tailored for DAC applications[10, 18].

This study investigates the design of a gas-liquid contactor with a large surface area for the chemical absorption of CO₂ from air using an aqueous potassium sarcosinate (K-SAR) solutions. Previous research has shown that aqueous basic salts of amino acids hold promise for DAC applications[18-21]. In recent years, aqueous amino acid salt solutions, particularly K-SAR, have garnered attention as superior alternatives to conventional amine solvents. These solutions offer several advantages such as higher loading capacities, faster CO₂ absorption kinetics, lower volatility and corrosivity, and greater chemical stability. This research paper focuses on the development of a high surface area gas-liquid contactor for the chemical absorption of CO₂ from ambient air using aqueous amino acid solutions containing K-SAR.

In this work, we propose a framework to create, manufacture, and experimentally evaluate advanced packing geometries based on a Gyroid design with a triply periodic minimal surface (TPMS) structure. TPMS structures consist of two interpenetrating fluid domains separated by a thin wall, exhibiting periodic properties in all three dimensions[22]. These nature-inspired surfaces minimize area, possess zero mean curvature, and offer smooth paths for fluid flow. Prior investigations have demonstrated that TPMS geometries can improve mass transfer performance by 49-61% and increase effective gas-liquid interfacial area by 91-140% compared to conventional packing materials like Mellapak 250.Y[10]. In this study, we

evaluated the CO₂ capture performance of a Gyroid geometry with an advanced structured packing and compared it to a conventional packed bed packing using polypropylene balls of different sizes. Three dimensionally (3D) printed Gyroid packing structure tested with varying air flowrates and solvent conditions to determine the optimal operational parameters. Then, the Gyroid packing test was performed in an environmental chamber under different temperature and humidity conditions to check its climate impact in terms of CO₂ capture performance.

2. Experimental Setup

2.1. Packing structure for air contactor

In the current study, a modular absorber device on a liter-scale was employed to assess the efficiency of packing materials in capturing CO₂ from ambient air using an aqueous 1M K-Sar solution (a mixture of potassium hydroxide and sarcosine). We investigated novel packing geometries for CO₂ capture from ambient air by employing 3D-printed structures known as TPMS with a Gyroid pattern. A comparative analysis was conducted between the Gyroid-structured packing and conventional packings composed of polypropylene balls of different sizes. The Gyroid packings demonstrated comparable or enhanced maximum fluid capacity and pressure drops compared to traditional packings, indicating their potential suitability for industrial applications.[10].

The Gyroid structure's geometric design was generated using the open-source software MSLattice, allowing for precise control over the desired thickness and unit length of each layer. Subsequently, STL (Stereo Lithography) files were generated using MSLattice to represent the Gyroid structure. The specimens were designed as Gyroid unit cells, with a total of seven unit cells per side length, resulting in a configuration measuring $3 \times 2 \times 2$ in³. To fill the $4 \times 4 \times 6$ in³ air contactor, eight 3D-printed specimens were produced. The Stratasys Fortus 450 printer was utilized for the 3D printing process, with each layer of the Gyroid structure having a thickness of 0.01 in. Stratasys water was used to dissolve the support structure within the printed part. ABS-30m was chosen as the material for the Gyroid structure due to its remarkable resistance to heat, cold, and chemicals, making it a crucial contributor to the advancement and proliferation of three-dimensional printing technology[23].

Figure 1 (a) provides visual representations of the packed beds comprising polypropylene particles, showcasing side and top views for three different particle diameters: 1/4 in, 5/16 in, and 3/8 in. In Figure 1 (b), the 3D-printed Gyroid packing, created using ABS-30m material, is presented from both the side and top perspectives. For a comprehensive overview of the various packing types, Table 1 offers a comparative analysis between the polypropylene balls and the 3D-printed Gyroid structure. Notably, the specific surface area (SSA) of the 3D-printed Gyroid structure, intentionally set at 579 m²/m³, is specifically compared to the smallest polypropylene packing (3/8 in) with a SSA of 578 m²/m³. A notable distinction between Gyroid packing and the packing with the smallest polypropylene (3/8 in) is the significantly higher voidage observed in the Gyroid structure. This disparity arises from the Gyroid's flexible design, which allows for adjustments of various functional parameters such as unit cell size, wall thickness, and curvature. The higher voidage offers increased open space within the packing material, potentially enhancing gas-liquid contact and improving mass transfer efficiency during CO₂ capture processes[24]. This information highlights the subtle differences in SSAs between the two packing types, emphasizing the potential advantages offered by the Gyroid structure in terms of surface area-to-volume ratio and its potential impact on CO₂ capture efficiency.

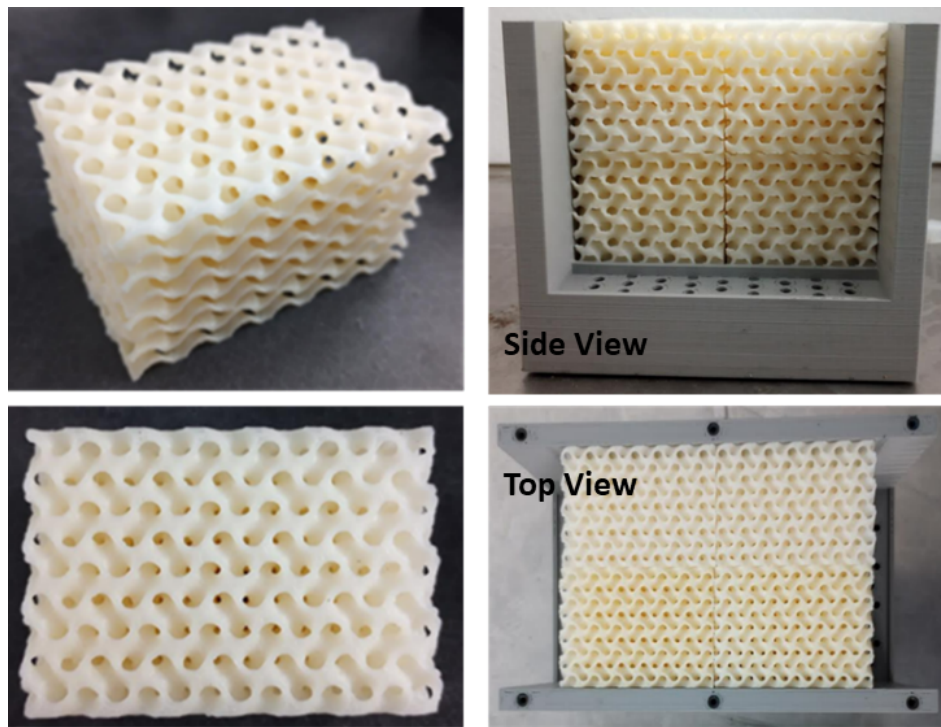
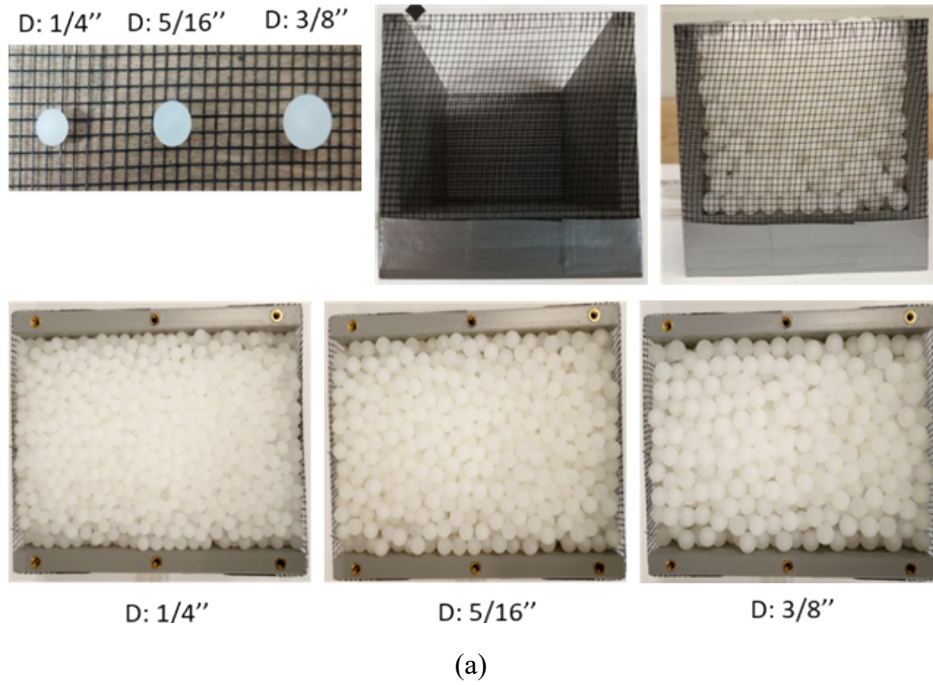
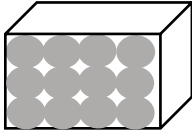
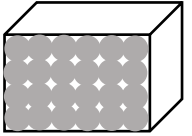
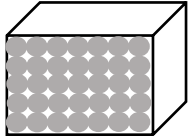
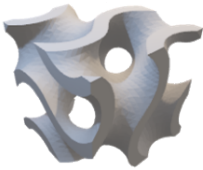


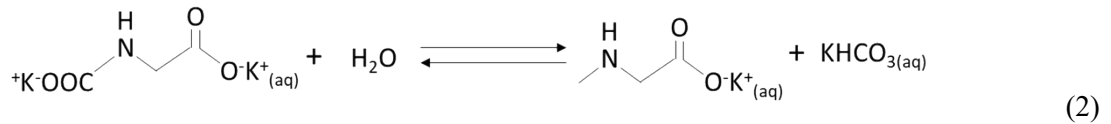
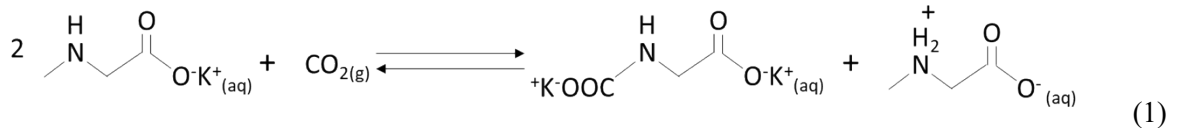
Figure 1. Packing types: (a) conventional packed bed by polypropylene balls, (b) 3D printed Gyroid.

Table 1. Specifications of each packing types for DAC application.

				
	Packed bed #1	Packed bed #2	Packed bed #3	Gyroid
Particle size/Unit cell size (in)	0.375	0.312	0.25	0.42
Voidage (%)	40.3	39.4	38.7	70
Total number of balls/unit cells	1,615	2,981	5,706	1,052
Specific surface area (m ² /m ³)	360	438	578	579
Actual surface area (m ²)	0.46	0.56	0.71	0.71

2.2. Solvent preparation and characteristics

The required chemicals, namely sarcosine (99% purity) and potassium hydroxide pellets (99% purity), were procured from Sigma Aldrich. To prepare a 1M K-SAR solution, a mixture of 89.1 g sarcosine and 56.1 g KOH was added to a 1-liter solution of deionized water. Mixing the components resulted in the formation of a 1-liter volume of 1M K-SAR solution. One of the key advantages of using amino acid solutions for CO₂ capture is the ability to regenerate the loaded solution. The dynamic viscosity, μ , and density, ρ , of 1M K-Sar are 0.01 kg/m-s and 1100 kg/m₃ respectively. The governing chemical reaction with atmospheric gas phase CO₂ and 1M K-Sar are shown in Equations 1 and 2. Regeneration is achieved by directly applying heat to the fluid, which facilitates the desorption of CO₂ and restores the amino acid to its basic form. This study explores the potential of utilizing aqueous amino acid solutions, specifically K-SAR, for CO₂ capture. The unique properties and regenerability of these solutions make them promising candidates for sustainable carbon capture technologies.



2.3. Air contactor design and wind tunnel application

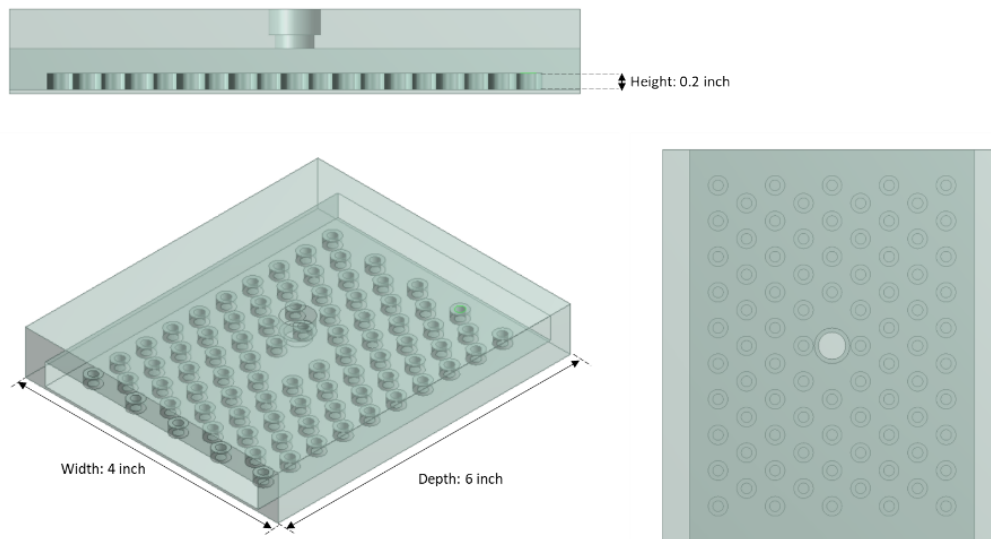
The design of the air contactor involves the implementation of a horizontal air flow that passes through the contactor. In this particular setup, a micro- to millimeter thickness layer of liquid sorbent is applied to a vertical packing material[24]. The design of a crossflow air contactor for this purpose is influenced by factors such as gas-solvent contact, solvent flow rate, and energy requirements[13, 28]. High surface area packing structures are utilized to enhance the interaction between the gas and liquid phases. Typically, stainless steel is used in conventional cooling tower packing structures. However, employing advanced packing structures potentially reduces the pressure drop across the contactor that can result in lower operational costs[10]. To optimize solvent flow rate and packing structure coverage while minimizing capital and operational expenses, the Gyroid packing has been proposed. The energy requirements associated with cross-flow contactors arise from the pumping of the solvent and the operation of fans to overcome the pressure drop across the contactor. For an annual capture rate of 1 Mt CO₂ per year, these contactors are estimated to be 20 m tall, 200 m long, and have a depth ranging from 5 to 8 m[14]. The depth directly affects the pressure drop across the contactor and the percentage of CO₂ captured from the inlet air stream, making its impact on total energy requirements complex.

Liquid solvent-based CO₂ absorption in structured packings was conducted using a modular absorber column that combined 3D-printed components with different packing structures. The purpose of this setup was to achieve the efficient absorption of CO₂. To ensure an even distribution of the solvent flow throughout the system, a specially designed top distributor was created. This distributor, which is featured in Figure 2(a), was 3D-printed and incorporated evenly spaced circular orifices at the top of the liquid flow inlet. Each orifice was designed to have a height of 0.2 in, allowing the liquid solution to collect at the top and smoothly flow down through the distributor.

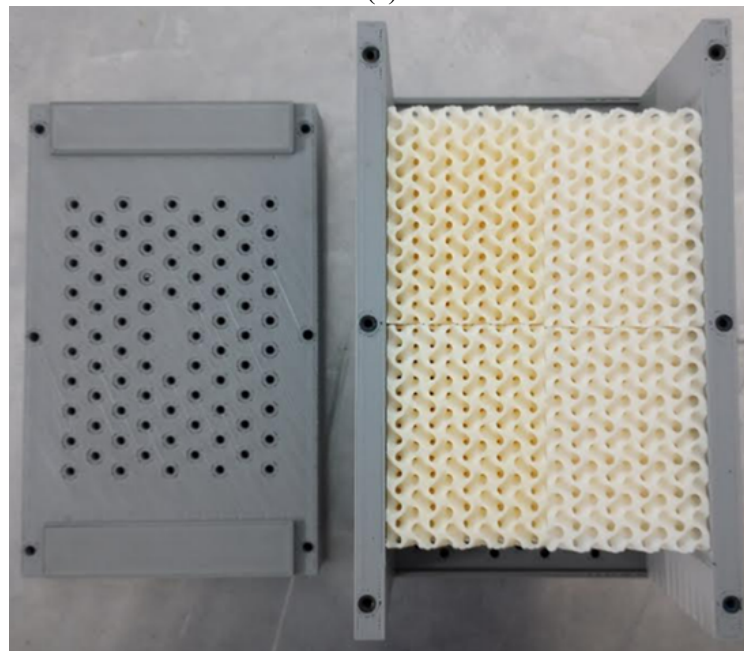
Figure 2(b) shows the 3D-printed contactor containing Gyroid packing, which was specifically chosen for the wind tunnel experiment. Gyroid packing is renowned for its substantial surface area and effective gas-liquid contact, thereby optimizing the absorption process. To assess the performance of the structured packings, a fixed volume of recirculated aqueous Amino Acid solvent (1M K-Sar, a mixture of Potassium Hydroxide and Sarcosine) was employed for CO₂ removal from the surrounding ambient air. A consistent volume of 1.5 liters of the solvent was used, and a continuous recirculation system was implemented. The liquid residue was collected in a designated container and subsequently pumped back to the top of the distributor, ensuring continuous flow and enhanced gas-phase interaction for improved CO₂ absorption.

CO₂ capture rates are observed at different ranges of air flow rates and solvent flow rates to find the optimum operating conditions. The CO₂ capture rate is calculated based on Equation 3:

$$CO_2 \text{ Capture Rate} = \frac{CO_{2,upstream} [ppm] - CO_{2,downstream} [ppm]}{CO_{2,upstream} [ppm]} \quad (3)$$

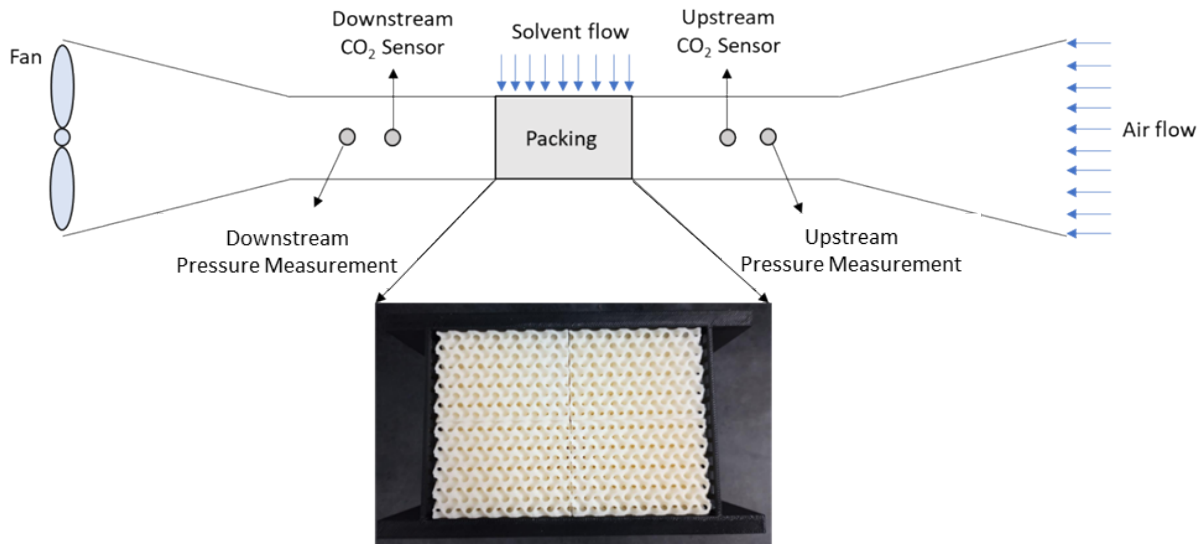


(a)

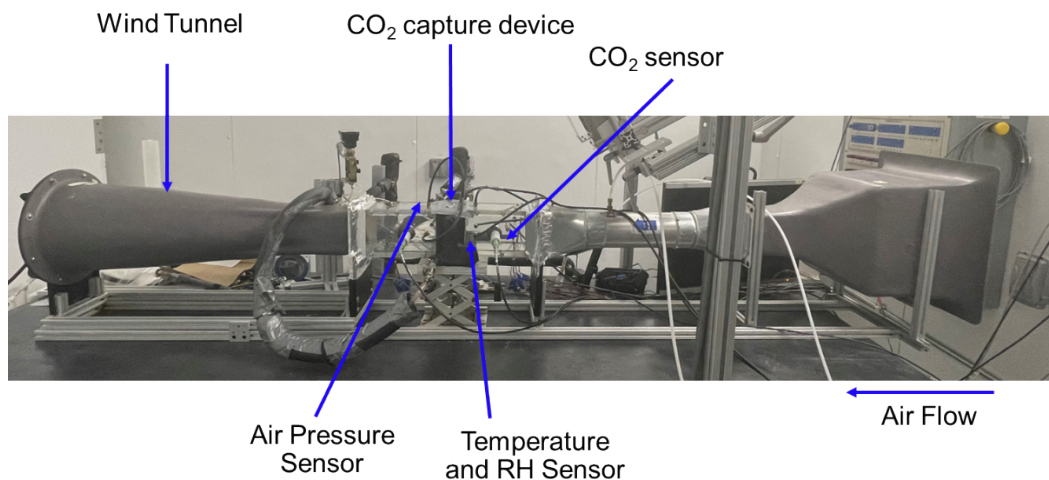


(b)

Figure 2. Liquid-solvent (a) distributor CAD design and (b) 3D-printed contactor with Gyroid packing for wind tunnel experiment.



(a)



(b)

Figure 3. (a) Experimental Schematic and (b) Wind tunnel setup for the DAC application experiment.

To ensure precise control over air temperature and humidity, a controlled environment was established by an environmental chamber. The wind tunnel served as the primary experimental setup for investigating the CO₂ capture performance of different packing types. Situated at the center of the wind tunnel, the various packings were subjected to airflow generated by a downstream fan. A distributor was employed to deliver the 1M K-Sar liquid solvent from the top of the air contactor to the packing, facilitating the gas-liquid interaction. Throughout the experiment, monitoring of the chamber's temperature and humidity levels was conducted to ensure consistency and control. To regulate the mass flowrate of the liquid solvent, a ball valve was utilized. Additionally, a differential pressure transducer was employed to measure the pressure drop across the air contactor, as indicated in Figure 3. The gaseous effluent from the absorber was directed to an Omega digital mass flowmeter, while a portion of it was supplied to an infrared CO₂ sensor (Vaisala GMP252). These instruments facilitated the measurement and analysis of upstream and downstream CO₂

concentrations within the system, providing valuable data for the evaluation of the CO₂ capture performance of the different packing materials.

3. Results and Discussion

In accordance with the outlined methodology, the wind tunnel setup (Figure 3) was utilized to conduct experiments comparing the CO₂ capture performance of conventional packed beds and 3D-printed Gyroid packings. Initially, the focus was on evaluating and discussing the optimal operational conditions of conventional packed beds with varying ball sizes and flowrates of solvent and air, emphasizing their CO₂ capture efficiency. Subsequently, a comprehensive comparison was made between the CO₂ capture performance of the conventional packed bed and the Gyroid packing. Through careful analysis and discussion, the relative effectiveness of these two packing types was assessed, providing valuable insights into their respective capabilities. Furthermore, the CO₂ capture performance of the Gyroid packing was investigated under diverse climatic conditions. The overall operational conditions of the wind tunnel experiment are displayed in Table 2.

Table 2. Operational range of DAC experimental condition.

Air Flowrate (LPS)	0.6 to 3.4
Solvent Flowrate (GPM) (in LPS)	0.06 to 0.16 0.00378 to 0.0101
Temperature (°F)	35, 65, 95
Relative Humidity	80%
Specific surface area (m ² /m ³)	360 to 579
Actual surface area (m ²)	0.46 to 0.71

3.1. Determining the Optimal Operational Parameters for Conventional Packings

Three different ball sizes were employed in conventional packed beds to investigate the impact of SSA on CO₂ capture rate performance. The ball sizes used were 0.25, 0.312, and 0.375 in, corresponding to SSA of 578, 457, and 375 m²/m³ (Table 1), respectively. Figure 4 illustrates the results obtained for the CO₂ capture rates of the three packed beds. The contour plots presented in the results were obtained from a dataset comprising 30 data points. Each data point represents the average values over a 3-minute interval. These data points were utilized to create the contour plots, which provide a visual representation of the variations and trends observed in the parameters being analyzed. Figure 4 reveals that packings with the two larger

ball sizes (0.312 and 0.375 inches) exhibited similar CO₂ capture performance across the range of 0.5-3.5 LPS air flowrate and 0.06-0.16 GPM solvent flowrate. These packings demonstrated CO₂ capture rates ranging from 5% to 25%, primarily dependent on the air flowrate rather than the solvent flowrate. In contrast, the conventional packing with the smallest ball size (corresponding to a SSA of 578 m²/m³) displayed significantly higher CO₂ capture rates, particularly at lower air flowrates, reaching above 50%. For instance, at air flowrates below 1 LPS and with a wide range of solvent flowrates between 0.09 and 0.16 GPM, the CO₂ capture rate exceeded 45%. However, at higher air flowrates, the influence of solvent flowrate became more significant in terms of CO₂ capture rate.

Figure 5 provides a comprehensive depiction of the CO₂ concentrations, both upstream and downstream, under two distinct testing conditions. The representative scenarios were chosen as: 1) Air flowrate: 1 LPS & Solvent flowrate: 0.12 GPM, and 2) Air flowrate: 2 LPS & Solvent flowrate: 0.12 GPM. The selection of these conditions was based on the observed stronger influence of air flowrate on the CO₂ capture rate, compared to the solvent flowrate. The results presented in Figures 5(b) and 6(c) pertain to a packed bed characterized by a SSA of 578 m²/m³. The testing was conducted over a duration of 5 hours, and the obtained outcomes are shown in Figures 5(b) and 6(c).

Examining the first testing condition (Air flowrate: 1 LPS & Solvent flowrate: 0.12 GPM), it can be observed that the CO₂ capture rate exhibited variability, ranging from 44% to 50% after a stabilization period of 10 minutes for both upstream and downstream CO₂ concentrations. In contrast, the second testing condition which features a two-fold increase in air flowrate displayed a relatively lower capture rate that ranged from 27% to 31% over the entire 5-hour test duration. These findings underscore the critical role of air residence time in achieving optimal CO₂ capture performance. Prior research has indicated that increasing the depth of the air contactor can lead to capture rates as high as 75%, facilitated by higher incoming air velocities of 1.4 m/s (equivalent volumetric flowrate, 0.0036 m³/s) [14]. However, it is essential to consider tradeoffs in the design process, such as the potential impact on pressure drop and the additional capital costs associated with modifying the contactor to accommodate the extended travel length of the air flow[29-31]. Addressing these considerations is crucial for achieving an optimal balance between CO₂ capture performance and the associated operational and economic considerations.

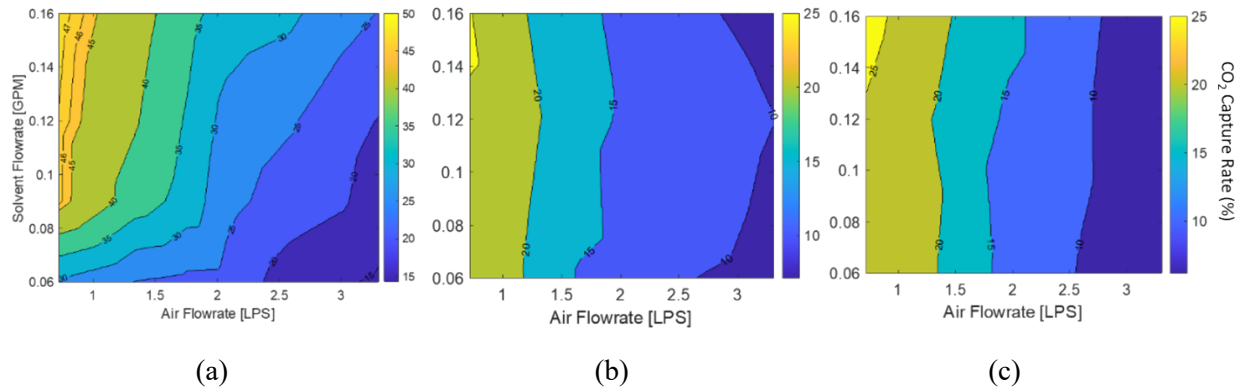
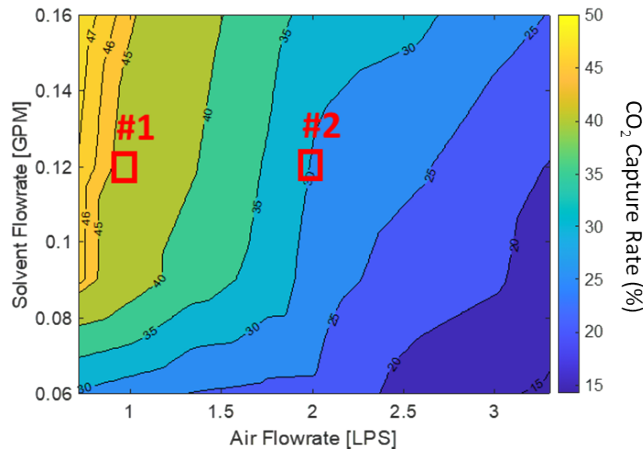


Figure 4. CO₂ capture rate for varying air flowrates and solvent flowrates (a) SSA: 578 [m²/m³] (b) SSA: 457 [m²/m³] (c) SSA: 375 [m²/m³].

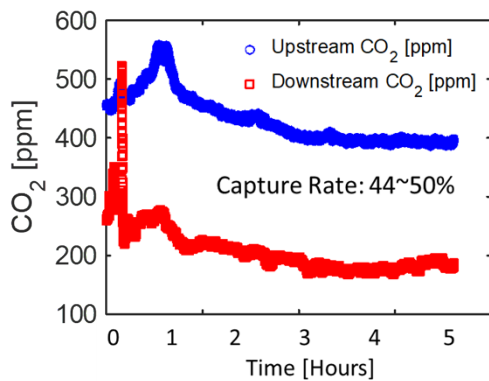
1/4" packed bed (SSA: 578 m²/m³)



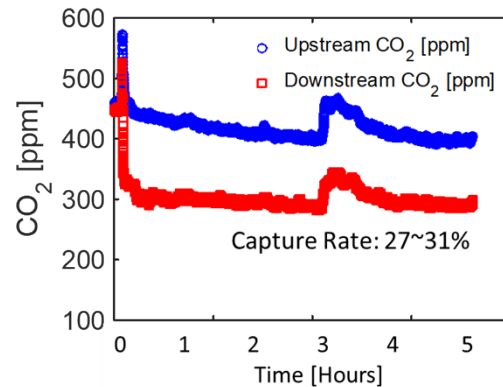
(a)

#1 Air flowrate: 1 [LPS] & Solvent flowrate: 0.12 [GPM]

#2 Air flowrate: 2 [LPS] & Solvent flowrate: 0.12 [GPM]



(b)



(c)

Figure 5. 5-hour testing at two different test conditions: (a) two representative experimental conditions (b) CO₂ upstream and downstream concentration at location #1, (c) CO₂ upstream and downstream concentration at location #2 using conventional packing with a SSA of 578 m²/m³.

3.2. Advantage of advanced Gyroid packing

We developed and manufactured the Gyroid packing using ABS-30m material to replicate the SSA of a conventional packed bed, which was 578 m²/m³. Initially, we conducted tests within the same solvent and air flowrate range, ranging from 0.06 to 0.16 GPM and 0.5 to 3 LPS, respectively, to assess the CO₂ capture performance and make a comparison with the conventional packed bed. Figure 6 illustrates the CO₂ capture rate and pressure drop of the conventional packed bed (SSA: 578 m²/m³) and the Gyroid packing (SSA: 579 m²/m³).

The overall CO₂ capture performance of the Gyroid packing appears to be similar to that of the conventional PP ball packing. However, significant differences were observed in the lower solvent flowrate range below 1 GPM. In this range, the Gyroid packing exhibited a higher CO₂ capture performance compared to the conventional PP ball packing. For instance, at an air flowrate of 1.5 LPS and a solvent flowrate of 0.06 GPM, the Gyroid packing achieved a 30.5% CO₂ capture rate, which was 20% higher than the conventional

PP packing's CO₂ capture rate of 24.4%. Similarly, at an air flowrate of 2 LPS and a solvent flowrate of 0.06 GPM, the Gyroid packing showed a 26.8% CO₂ capture rate, surpassing the conventional PP packing's CO₂ capture rate of 22.4% by 16.4%. These results can be attributed to the material's surface tension and the fluid dynamic behavior at the interface between the liquid film and the CO₂ gas phase in the air. The surface energy of the material is a critical parameter influencing the wetting efficiency on the column layer surface[10, 24]. As anticipated, the ABS-30m material exhibited a 20.7% higher surface energy (38.5 mJ/m²) compared to the critical surface energy of the conventional PP packed bed (30.5 mJ/m²). Consequently, the contact angles were measured at 80.9° (hydrophilic) and 102.1° (hydrophobic) for ABS-30m and conventional PP, respectively[32].

Another notable result is the pressure drop across both types of packing throughout the entire range of air flowrates, which was from 0.5 to 3 LPS. A common tradeoff and optimization point in liquid absorption is the balance between packing surface area and air flowrate. Increasing either parameter enhances the CO₂ capture rate but also leads to increased pressure drop. As shown in Figure 6, as the air flowrate increased while maintaining a fixed SSA, the pressure drop rose for both packing types. However, across all air flowrates, the Gyroid packing exhibited 33% to 77.8% lower pressure drop compared to the conventional PP packing. This finding aligns with previous studies involving ceramic foams and spherical meshes, as well as 3D printed packing which also demonstrated reduced pressure drop[33-36]. The impact of air movement by fans can be significant. According to a previous study on fan power in relation to pressure drop[12], fan power is directly proportional to pressure drop. This means that the Gyroid packing structure has the potential to reduce operational fan power by 33% to 77% in the air contactor unit. Nevertheless, there are limitations due to factors such as the capital cost of mass production of 3D printed advanced packing, raw material expenses, and the duration of post-processing. The packings differ considerably from each other in terms of cost or pressure drop per unit length, necessitating careful consideration of column height and air flowrates to achieve a balance between the two factors.

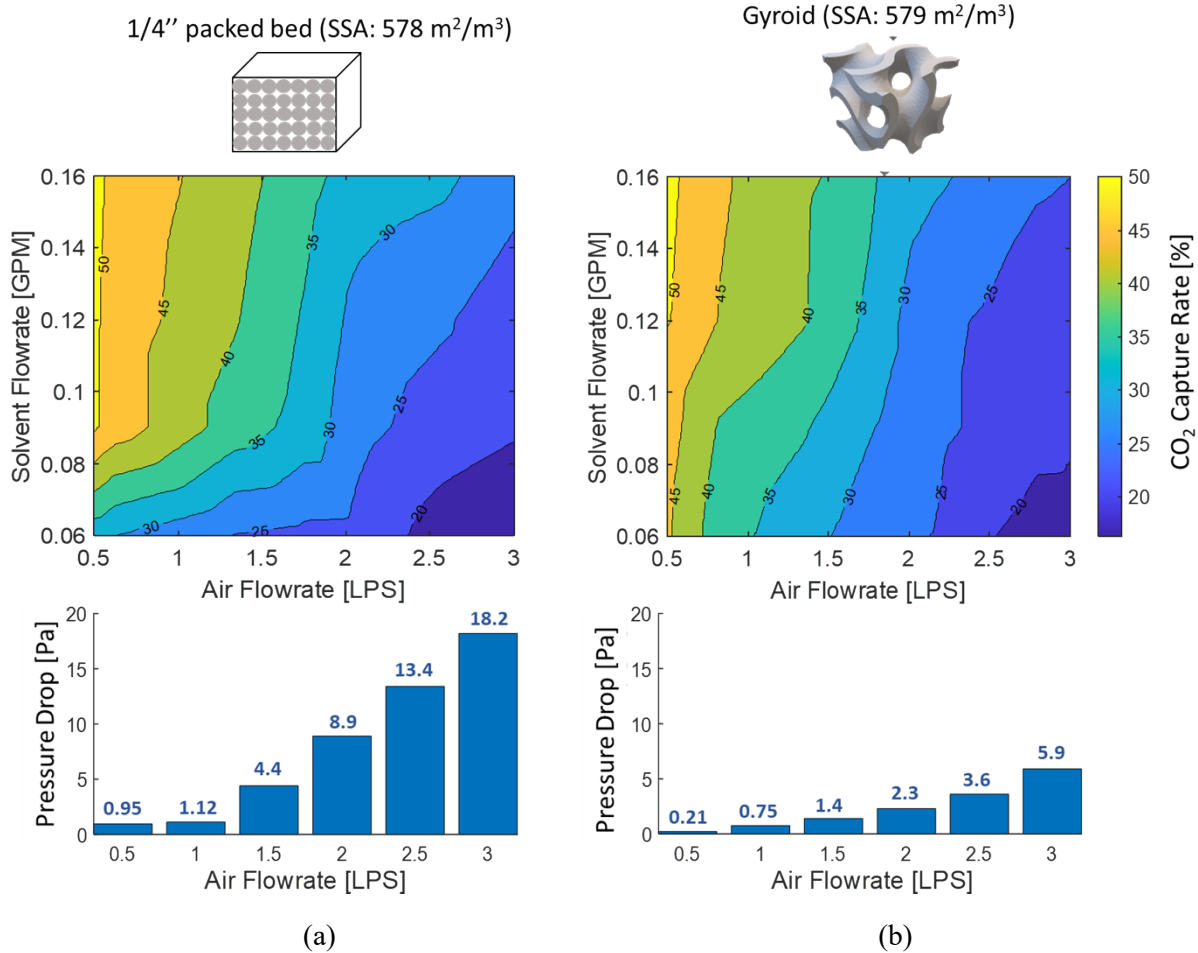


Figure 6. CO₂ capture rate and pressure drop of (a) conventional packed bed (SSA: 578 m²/m³) and (b) Gyroid packing (SSA: 579 m²/m³).

3.3. Climate impact on CO₂ capture performance

In this study, the impact of different climatic conditions on the CO₂ capture performance of the advanced Gyroid packing structure was investigated. The experiments were conducted under fixed operational conditions, including an air flowrate of 1 LPS and a solvent flowrate of 0.12 GPM. Three distinct climatic conditions were considered, namely 35°F, 65°F, and 95°F, with a fixed relative humidity of 80% to minimize water evaporation during the experiments. Each climatic condition was tested for a duration of 1 hour, with data collected at a time resolution of 1 second, resulting in a total of 3,600 data points. The results were analyzed and presented in the form of a whisker plot, as depicted in Figure 7.

The median CO₂ capture rates at the different climatic conditions were observed to be 23.2% at 35°F, 34.9% at 65°F, and 46.8% at 95°F, all at a fixed relative humidity of 80%. It was found that as the atmospheric air temperature increased, the CO₂ capture rates exhibited more stable results. For instance, at an air temperature of 95°F, the CO₂ capture rates were confined within a narrow range of 3.4%. In contrast, at lower temperatures of 65°F and 35°F, the range of CO₂ capture rates increased significantly, with variations of 9.4% and 11.4%, respectively. These findings highlight the influence of climate conditions, particularly

temperature, on the CO₂ capture performance of the Gyroid packing structure, underscoring the importance of considering environmental factors when optimizing and implementing DAC systems.

Previous studies[24, 37] have indicated that the absorption of atmospheric CO₂ into strong aqueous hydroxide solutions is primarily governed by gas-phase resistance and follows a fast pseudo-first-order reaction. The flux of carbon dioxide across the air-liquid interface can be calculated using Equation 4:

$$J_{CO_2} = \sqrt{D_L k [OH^-]} K_H C_{CO_2} \quad (4)$$

where J_{CO_2} represents the flux of CO₂ into the hydroxide solution with a concentration of [OH⁻], per unit area per unit time, D_L denotes the diffusivity of CO₂ in the solution, k is the temperature dependant kinetic constant for the reaction of CO₂ with OH⁻, K_H is Henry's solubility constant for CO₂, and C_{CO_2} represents the concentration of CO₂ in the air. The units and values at each condition are shared in Table 3. Based on Equation 4 and the properties provided in Table 3, the maximum CO₂ flux at an air temperature of 95°F is calculated to be 7.31E-05 mol/m²-s, which is 26% higher than the maximum CO₂ flux at an air temperature of 77°F, which is 5.41E-05 mol/m²-s. The CO₂ capture rate results exhibit a similar trend to the CO₂ flux, with the median CO₂ capture rate at 95°F (46.8%) being 17.9% higher than the median CO₂ capture rate at 77°F (39.6%) based on a linear fit with an R² value of 1. This finding suggests a strong correlation between the capture rate and ambient temperature at controlled CO₂ flux.

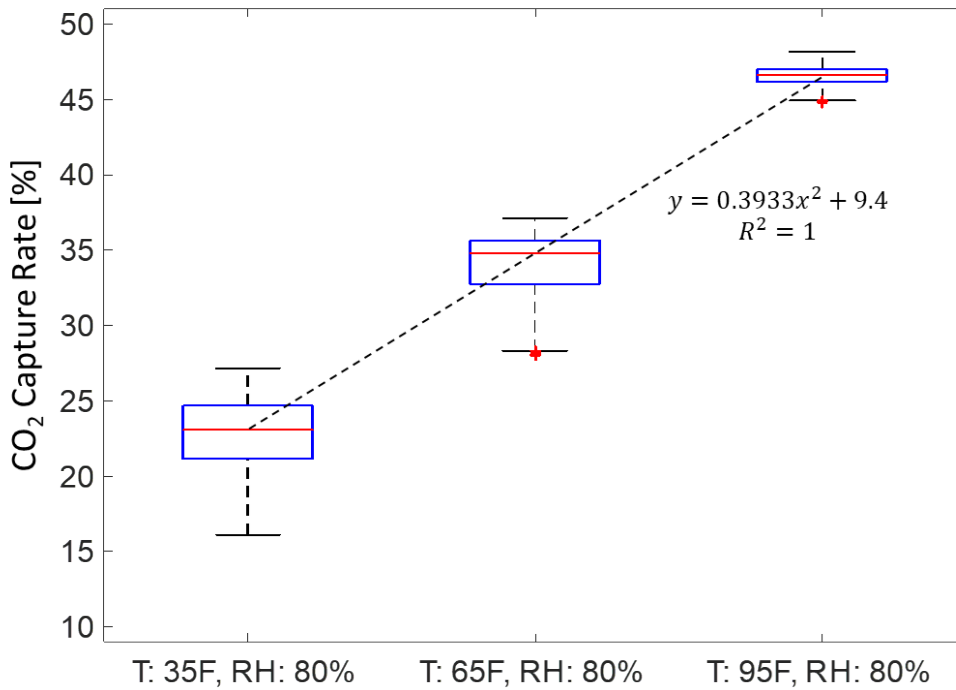


Figure 7. CO₂ capture rate at three different climate conditions: T: 35°F & RH: 80%, 65°F & RH: 80%, 95°F & RH: 80% (with linear fit curve with R²=1).

Table 3. Properties to calculate CO₂ flux at the air and solution interface.

	T: 25°C (=77°F)	T: 35°C (=95°F)	Reference/Note
D_L [m ² /s]	1.54E-09	1.89E-09	[38]
k [L/mol-s]	1.86E+04	4.17E+04	[39]
OH ⁻ [mol/L]	1	1	-
K_H	5.46E-01	4.45E-01	[38]
C_{CO_2} [mol/m ³]	0.0185	0.0185	Equivalent to 415 ppm
J_{CO_2} [mol/m ² .s]	5.41E-05	7.31E-05	-

4. Conclusions

This study focused on optimizing DAC technology using a crossflow air-liquid contactor with a 3D printed Gyroid packing structure and a 1M K-Sar solvent. By investigating various operational parameters, the study aimed to enhance the design and efficiency of DAC systems. The results demonstrated that the Gyroid packing exhibited comparable CO₂ capture performance to conventional packed bed packing, while significantly reducing pressure drop. This finding highlights the potential of the Gyroid design as an efficient and cost-effective solution for gas-liquid contactors in DAC, contributing to the advancement of scalable and sustainable carbon capture systems.

A notable aspect of this study is the use of aqueous amino acid solutions, particularly K-SAR, as solvents for CO₂ capture. These solutions offer advantages over traditional amine solvents, including higher loading capacities, faster absorption kinetics, lower volatility and corrosivity, and greater chemical stability. This study analyzed the climate impact on CO₂ capture performance by subjecting the Gyroid packing to different environmental conditions. The results showed that higher air temperatures led to more stable CO₂ capture rates. This can be attributed to the increased surface energy of the ABS-30m material used in the Gyroid packing, facilitating improved wetting efficiency at the air-solvent interface. Therefore, understanding the influence of climatic conditions on DAC systems is crucial for effective deployment and optimization.

In summary, this study contributes to the advancement of DAC technology by optimizing operational parameters and investigating the potential of advanced packing materials. The Gyroid packing offers comparable CO₂ capture rates (close to 50% at lower air flowrates) to conventional packed bed packing while significantly reducing pressure drop from 33% to 77.8%, indicating its potential as an efficient and cost-effective solution. Our investigation yielded results indicating that the Gyroid packing achieved a similar level of CO₂ capture performance as the conventional packed bed packing, but with a significantly lower pressure drop (3 to 4 times lower). These findings suggest that the Gyroid design holds promise as an efficient and effective solution for gas-liquid contactors in the chemical absorption of CO₂ from air. The Gyroid structure, which is inspired by nature, features periodic surfaces that minimize area and facilitate smooth fluid flow, contributing to its superior performance. Additionally, the Gyroid packing demonstrated resilience and adaptability across different environmental conditions, highlighting its versatility for practical applications. Moreover, considering the climate impact on CO₂ capture performance provides valuable insights for the design and optimization of DAC systems. Lastly, these findings contribute to the development of scalable and sustainable carbon capture solutions, thereby supporting global efforts to address climate change.

Nomenclature

DAC: Direct Air Capture

NET: Negative Emission Technology

CCSU: Carbon Capture and Storage/Utilization

CDR: Carbon Dioxide Removal

PP: Polypropylene

K-SAR: Potassium Sarcosinate

TPMS: Triply Periodic Minimal Surface

LPS: Liter per Second

GPM: Gallon per Minute

SSA: Specific Surface Area

D_L : Diffusivity

K_H : Henry's Constant

C_{CO_2} : CO₂ Concentration

J_{CO_2} : CO₂ Flux

μ : Dynamic Viscosity

ρ : Liquid Phase Density

Acknowledgements

This research is supported by the US Department of Energy (DOE), Building Technologies Office and Laboratory Directed Research and Development and under Contract DE-AC05-00OR22725 with UT-Battelle LLC. This research used resources at the Building Technologies Research and Integration Center (BTRIC), a DOE Office of Science User Facilities operated by the Oak Ridge National Laboratory.

Data Availability Statement

The data that supports the findings of this study is available in the Supplementary Information.

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