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Article

Experimental and Simulation Research on Indoor CO₂ Removal Efficiency and Fresh Air Energy Savings of Living Walls in Office Spaces

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Abstract: Elevated indoor CO₂ levels might have adverse effects on human health. However, the introduction of outdoor air to lower indoor CO₂ concentrations results in significant HVAC energy consumption. Aligning with office hours and the natural light cycle, the utilization of photosynthesis in living walls offers an energy-efficient and sustainable solution for the mitigation of high CO₂ levels in office spaces. This study experimentally investigates the impacts of the carbon fixation pathways, light intensity, and substrate moisture content on the CO₂ removal rate of living walls at the room scale. Furthermore, the fresh air energy-saving effects of living walls under different scenarios are accurately simulated in EnergyPlus. The results demonstrate that choosing C₃ plants over CAM plants in living walls yields higher CO₂ removal efficiency. In a 30-m² office room accommodating 2-3 occupants, living walls can reduce the demand for fresh air by 12.3%-27.8% and decrease fresh air energy consumption by 11.2%-28.2%. The city with the highest energy savings has energy savings that are 4.5 times greater than those of the city with the lowest energy savings. The findings of this research promote the application and development of living walls, thus providing a viable solution for improving indoor air quality.

Keywords: CO₂ concentration; living walls; fresh air energy consumption; Energyplus; indoor air quality

1. Introduction

Poor indoor air quality results in various building-related illnesses that negatively impact human health [1–3]. Extremely high concentrations of CO₂ (>20,000 ppm) have the potential to be fatal [4,5]. In typical indoor environments in non-industrial buildings (<5000 ppm), elevated CO₂ levels impair lung gas exchange function [6], leading to worker fatigue and drowsiness [7], headaches [8,9], and negative effects on cognition and decision-making [10–12]. The acceptable upper limit for the CO₂ concentration in non-industrial buildings in China is 1000 ppm [13], and ASHRAE standard 62.1-2019 recommends that indoor CO₂ concentrations should not exceed this level [14]. The outdoor CO₂ concentration typically remains around 400 ppm [15], posing no harm to human health. However, in enclosed spaces, CO₂ levels can rapidly rise due to human respiration, which can potentially harm human well-being.

The commonly used CO₂ capture technologies are challenging to implement in civil buildings. Absorption is unsuitable indoors due to high energy consumption, the large equipment size, and potential air pollution concerns [16]. Adsorption exhibits reduced CO₂ adsorption capacity in humid indoor conditions due to the effects of surface water vapor [17]. Membrane separation faces obstacles in terms of low efficiency at low CO₂ concentrations and high costs [18]. In contrast, a relatively simple approach involves the introduction of outdoor air to dilute indoor CO₂ concentrations. However, for this approach, the impact of natural ventilation on the thermal comfort of occupants [19] and the potential introduction of outdoor pollutants into indoor spaces [20] must be considered.

Mechanical ventilation, while effective, requires energy consumption [21]. Additionally, HVAC systems are necessary to condition the outdoor air for human thermal comfort, which includes processes such as heating, humidification, cooling, and dehumidification. These HVAC processes contribute to approximately half of the energy consumption in HVAC systems [22,23]. Therefore, it is crucial to explore energy-saving and sustainable approaches for the reduction of indoor CO₂ concentrations.

Phytoremediation provides an effective method for air purification [24–26]. Remarkably, researches in the aerospace field have shown that a closed system integrating plants and humans can achieve a balanced carbon cycle [27–29]. However, in civil buildings, the limited space indoors may not allow for a complete balance between plant absorption and human respiration. Nonetheless, the presence of plants can still significantly decrease indoor CO₂ levels. Pegas et al. [30] placed six plants in a classroom and observed an average decrease in indoor CO₂ concentration from 2004 to 1121 ppm. Tudiwer et al. [31] found that a classroom with living walls exhibited a 3.5% faster reduction in CO₂ concentration compared to a classroom without plants. Meng et al. [32] placed living walls in an air-conditioned room and observed a reduction in indoor CO₂ concentration of approximately 10%. Yungstein et al. [33] observed that the implementation of living walls in actual workplace led to an average decrease of 4.8% in the CO₂ concentration. While qualitative studies have demonstrated the potential of plants in to purify CO₂ in indoor environments, practical implementation remains challenging.

Due to their specific characteristics, office spaces are well-suited for the utilization of the living walls to purify CO₂. In these environments, workers inhale O₂ and exhale CO₂, while plants perform photosynthesis, absorbing CO₂ and releasing O₂, thereby creating a carbon cycle within the confined space [28]. The working hours of employees [34] align with the natural light cycle, thus allowing plants to utilize natural light for air purification. Despite the potential impact on human thermal comfort [35] and equipment aging [36], areas with concentrated solar radiation in offices are viable locations for the installation of the living walls. Temperatures in offices are regulated to ensure the thermal comfort of employees [37,38], and most plants can adapt to air-conditioned rooms [39,40]. Furthermore, interaction with plants can enhance productivity of workers [41]. Shao et al. [42] observed that 100 heads of *pakchoi* could reduce the indoor CO₂ concentration in a 30-m² office by 25.7%-34.3%. However, Pennisi et al. [43] suggested that under office lighting conditions, most plants have limited impact on indoor CO₂ levels. Gubb et al. [44] found that the influences of dry and wet substrates on the plant CO₂ purification capacity under typical office lighting levels could be ignored, as plants generally fail to reach their LCP. In conclusion, living walls are suitable for CO₂ purification in office spaces, but further research on plant selection, lighting design, and irrigation management is needed to provide practical guidelines.

Living walls that absorb CO₂ can reduce fresh air energy consumption. In office spaces, the occupants' respiration releases CO₂, which can lead to a rapid increase in indoor CO₂ concentration. If left uncontrolled, the elevated release of CO₂ from the respiration of occupants in an office setting can potentially lead to air pollution [45]. When the concentration of CO₂ released by workers exceeds the required standards, outdoor air is introduced to dilute the indoor CO₂ levels, which results in energy consumption by the HVAC system to process outdoor air. By absorbing CO₂, living walls can decrease the demand for outdoor air and consequently reduce fresh air energy consumption. Torpy et al. [46] estimated that 15 pots of *Dypsis lutescens* could potentially reduce the outdoor air demand by approximately 6%. Parhizkar et al. [47] found that the incorporation of 5 m² of *Azolla* per person in a typical office building could decrease the outdoor air requirement by about 30%. Shao et al. [42] demonstrated that 100 heads of *pakchoi* could reduce fresh air energy consumption by approximately 12.7%-58.4%. Currently, researchers mainly estimate the fresh air energy-saving impact of the living walls by calculating the balance between the CO₂ absorption rate of plants and the CO₂ generation rate of humans. However, this approach makes it challenging to accurately evaluate the fresh air energy-saving capacity of the living walls. EnergyPlus is a simulation tool capable of calculating the transient building loads required to maintain the temperature and ventilation setpoints for a full year

under specific conditions [48]. Nonetheless, there have been limited studies on the use of EnergyPlus to accurately assess the fresh air energy savings of the living walls.

Based on these reviews, living walls are suitable for indoor CO₂ removal and the reduction of fresh air energy consumption in office spaces. This study consists of two main parts, namely (1) the experimental investigation of the impacts of adjustable factors in office spaces on the indoor CO₂ removal rate of the living walls, and (2) the simulation-based evaluation of the fresh air energy savings achieved by CO₂ removal from living walls under different scenarios. The primary objective of this work is to improve the CO₂ removal efficiency of the living walls in office spaces and accurately assess their fresh air energy savings, with the aim of promoting the practical implementation of the living walls.

2. Materials and Methods

2.1. Experiment

The experiment was conducted with the aim of enhancement the CO₂ removal efficiency of the living walls in office spaces. It consisted of three steps. Firstly, the natural CO₂ infiltration rate in the two experimental rooms was confirmed to be the same, allowing for a comparative analysis. Secondly, the impacts of adjustable factors of office spaces such as carbon fixation pathways, light intensity, and substrate moisture content, on the CO₂ removal rate of the living walls were examined in the two experimental rooms. Finally, the significant factor was further investigated to explore its impact on the CO₂ removal efficiency of the living walls.

2.1.1. Experimental set up

The experimental site was located in Guangzhou, Guangdong Province, southern China, with geographic coordinates of 23°10' N, 113°21' E. The fluctuating outdoor wind environment and indoor-outdoor temperature difference have significant impacts on the natural CO₂ infiltration rate of the room [49,50], thereby affecting the calculated CO₂ removal rate of the living walls. To minimize the impact of external weather conditions and accurately measure the CO₂ removal rate of the living walls, two experimental rooms with the same structure were built on the rooftop (Figure 1 (a)). Room A, as the experimental group, was equipped with living walls, while room B, as the control group for comparison experiments, was a conventional air-conditioned room without living walls. Both experimental rooms had an exterior-facing west wall with interior walls on the remaining three sides. Each room measured 1.9 m × 1.9 m × 3.0 m (width × depth × height). The walls were constructed with a 2-mm metal sheet, a 100-mm EPS foam board, and another 2-mm metal sheet. The original windows of the rooms were sealed with black polypropylene (PP) sheets to improve airtightness and block natural light. This ensured that all light used in the experiment was provided by a stable artificial light source.

An air conditioner (710 W) was utilized to regulate the air temperature in the room, and was positioned 2.2 m above the ground. The air temperature was set at 25 °C, which is suitable for human thermal comfort [51]. An air quality monitor capable of measuring the air temperature, relative humidity, and CO₂ concentration was centrally placed at a height of 1.2 m, consistent with the breathing height of the occupant. A fan (40 W) on the ground was used to enhance air mixing, and a CO₂ release device, consisting of dry ice and a container, was placed in front of the fan. Dry ice was added to the boiling water in the container to accelerate its sublimation. The layout of the instruments in the room is shown in Figure 1(b), and the specific instruments are listed in Table 1.

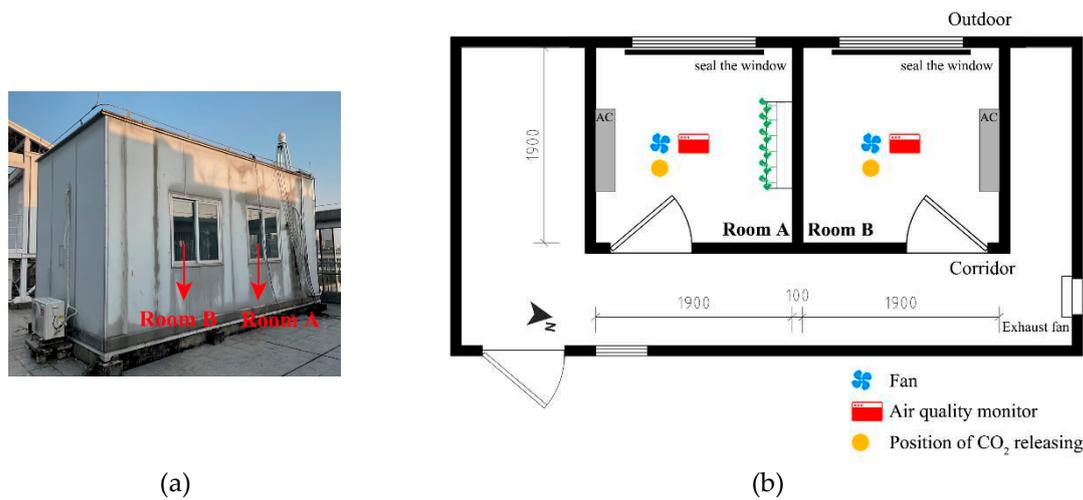


Figure 1. The appearance and plan of the experimental room: (a) appearance of the experimental room; (b) settings of the instruments.

Table 1. The instruments used in the tests.

Instrument	Model	Parameters	Accuracy	Setting Points
Air quality monitor	SenseAir S8	CO ₂ (ppm)	±40 ppm	Indoor: In the center of the room, 1.2 m above the ground
	SensenSHT20	Temperature (°C)	±0.3 °C	
		Relative humidity (%)	±3%	
Photosynthetic photon flux density (PPFD) meter	Apogee MQ-500	PPFD/	±5%	On the top of canopy
	Apogee SQ-520	($\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$)		Outdoor: Roof
CO ₂ sensor	RS-BYH-CO ₂ -M	CO ₂ (ppm)	±40 ppm (25 °C)	Outdoor: Roof
	Campbell 81000	Wind speed (m/s)	±1%	
Weather station	Campbell CS215	Temperature (°C)	±0.3 °C (25 °C)	Outdoor: Roof
		Relative humidity (%)	±2% (25 °C)	
Volumetric water content (VWC) sensor	Decagon Devices 5TE	VWC (m^3/m^3)	±3%	Substrate

2.1.2. Experimental materials

Sansevieria trifasciata and *Epipremnum aureum* were selected as experimental plants in this study due to their widespread use in relevant studies [52] and their potentially high CO₂ removal efficiency. They are commonly used as indoor plants in the local area where field surveys were conducted, and can adapt to low-light conditions indoors. Additionally, they can represent different carbon fixation pathways. Further details about the selected plant species are presented in Table 2.

The plants used in this study were of the same species, with similar health, age, and size characteristics. However, because the plants were purchased from different cultivation sites, there were variations in the composition of the growth substrate and cultivation methods. The substrate can significantly impact the indoor CO₂ concentration and, in some cases, can even surpass the CO₂ absorption of certain plant species [53]. To minimize this effect, the plants were transplanted into a uniform nutrient soil containing natural organic matter, rooting powder, polyoxin, perlite, and vermiculite. The diameter of the flower pot was 10 cm.

After transplantation, the plants were placed in the room for 14 days to acclimate to the new environment. The plants received artificial light ($5\text{--}10 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$) for a 12-hour light/dark cycle (07:00–19:00 light period) to simulate natural light conditions. During the maintenance period of the plants, the room was kept ventilated, and no fertilizers were used. The *E. aureum* was watered every 3 days, while the *S. trifasciata* was watered every 7 days. Watering sessions were performed between 08:00 and 09:00, with each session continuing until water overflowed from the bottom of the flower pot. During the experiment, ventilation was stopped, and humidity was not controlled. The temperature was set to $25 \text{ }^{\circ}\text{C}$ to allow the plants to acclimate to the experimental environment for 24 h. After the experiment, the plants were destructively measured to determine their leaf area.

Living walls were constructed in a modular manner and could hold 48 plants, and had an overall size of $0.84 \text{ m} \times 0.25 \text{ m} \times 1.606 \text{ m}$ (width \times depth \times height). The living walls included a water tank and plant modules (Figure 2). The height of the plant modules was 1.276 m, and the total area of the living walls was 1.07 m^2 .

Table 2. The characteristics of the plant species selected for experiments. The leaf area and plant height ($n = 6$) are presented as the means \pm standard error of the mean (SEM).

Species	Family	Metabolism	Leaf Type	Leaf Area (cm^2)	Plant Height (cm)
<i>Sansevieria trifasciata</i>	Asparagaceae	CAM	succulent	537 ± 77	28 ± 1
<i>Epipremnum aureum</i>	Araceae	C ₃	herbaceous	614 ± 111	22 ± 1



Figure 2. The living walls in the experimental room: (a) the structure of the living walls; (b) photos of the living walls.

2.1.3. Experimental process

2.1.3.1. Confirmation of identical natural infiltration in two rooms

During the experiment, high-concentration CO_2 was released once, after which the indoor CO_2 concentration gradually decreased. The decline in the CO_2 concentration in the room with living walls was caused by two factors, namely the net photosynthesis of the living walls and the natural infiltration of the room. At the room scale, meteorological conditions such as the wind speed, the wind direction, and indoor-outdoor temperature differences can significantly affect the natural CO_2 infiltration rate of the room. These changes could have impacted the CO_2 removal rates of the living walls measured in the experiment.

In a single experimental room, it is challenging to measure the decrease in CO_2 concentration caused by natural infiltration. Multiple experiments conducted under similar and relatively stable outdoor meteorological conditions are necessary to determine the natural CO_2 infiltration rate of the room.

Prior to each experiment, the current outdoor meteorological conditions were assessed. Moreover, after the experiment, the meteorological conditions during the experiment were reviewed to avoid the significant impacts of sudden changes in the meteorological conditions on the measurement of the CO₂ removal rates of the living walls.

A comparative experiment of the two rooms was conducted. Room A, as the experimental group, was equipped with living walls while Room B without living walls was used as the control group. CO₂ was simultaneously released in both rooms, and the same outdoor meteorological conditions were applied to both rooms. The changes in the CO₂ concentration in both rooms were measured, and the decrease in CO₂ concentration in the control Room B during the experiment was used to estimate the decrease caused by natural infiltration in the experimental Room A. The CO₂ removal rates of the living walls were calculated by subtracting the estimated natural infiltration rate from the total decrease in CO₂ concentration in Room A.

To measure the CO₂ removal rates of the living walls, it is necessary to confirm that the two identically constructed rooms had the same natural infiltration rate of CO₂ before conducting the experiment. Before placing living walls in Room A, three random experiments were conducted in both empty rooms without selecting specific outdoor meteorological conditions. The initial CO₂ concentration and duration of the experiments were consistent with those of the measurement experiment. After each experiment, the decrease in CO₂ concentration in both rooms without living walls was compared, and adjustments were made to ensure that the difference in CO₂ concentration reduction was within an acceptable range. Once confirmed, living walls were placed in Room A, and the measurement experiment began.

2.1.3.2. Optimization of the impacts of the adjustable factors of office spaces on CO₂ removal efficiency

The carbon fixation pathways, light intensity, and substrate moisture content were selected as controlled variables to investigate their impacts on the CO₂ removal efficiency of the living walls. The reasons for this selection are as follows.

1. Plants absorb CO₂ via photosynthesis, which is influenced by both internal and external factors. Internal factors refer to the carbon fixation pathways of plants (C₃, C₄, and CAM), while external factors include light, the environmental CO₂ concentration, the temperature, mineral elements, and water [54]. In office spaces, the temperature is usually set to ensure thermal comfort, while humidity is often difficult to regulate. Therefore, in the experiment, the temperature was controlled at 25 °C but the humidity was not regulated.
2. Due to the unsuitability of C₄ plants for indoor environments, C₃ and CAM plant species were selected as the experimental subjects. This selection was based on their distinct carbon fixation efficiency and adaptability to different environmental conditions.
3. Supplemental lighting was installed to ensure the function of the living walls due to the variability of the natural light intensity influenced by factors such as the weather, season, and orientation. Thus, it was necessary to examine the effect of light intensity on the living walls and adjust the supplementary illumination.
4. The substrate moisture content is closely related to the irrigation system and maintenance of the living walls, and insufficient and excessive watering are common issues in the management of the living walls [44]. The investigation of the effects of different substrate moisture levels on the CO₂ removal efficiency of the living walls can help to optimize the maintenance procedures and reduce associated costs.

The following four light intensities were set during the experiment: dark (0 μmol·m⁻²·s⁻¹), low light (10 μmol·m⁻²·s⁻¹), medium light (50 μmol·m⁻²·s⁻¹), and high light (400 μmol·m⁻²·s⁻¹). Low light was chosen to simulate the typical lighting conditions in office spaces, which commonly have an illuminance level of not less than 300 lx to support the productivity of workers [55]. Moderate light is sufficient for most plants to lower indoor CO₂ concentrations [46], as their net photosynthesis rate surpasses the CO₂ release rate from substrate microorganisms. High light is the maximum light

intensity achievable on the plane under laboratory conditions. The maximum light intensity was provided by eight high-efficiency and energy-saving LEDs (300 W, 6500 K white light), while the other light intensities were achieved by adjusting the number of lamps and their distance from the living walls. Four points on the living walls were evenly sampled, and the photosynthetic photon flux density (PPFD) was measured by the Apogee MQ-500 instrument placed at the highest point of the canopy, with the average value used to determine the experimental light intensity.

In the experiment, the substrate moisture content was divided into two moisture levels: dry and wet. Pre-experimental findings revealed that watering plants daily greatly increased the risk of root rot and death. Therefore, watering was carried out every 3 days. During this period, the lowest volumetric water content (VWC) measured for *S. trifasciata* and *E. aureum* was $0.15 \text{ m}^3/\text{m}^3$, which was defined as a wet substrate. If the soil was not watered for a certain period of time, it would become dry and clumpy, and the plants would not wilt immediately. The highest VWC measured during this period was $0.05 \text{ m}^3/\text{m}^3$, which was defined as a dry substrate. During the wet substrate experiment, three randomly selected plants on the living walls were measured using a 5TE VWC sensor before the start of each experimental day. If the VWC was below $0.15 \text{ m}^3/\text{m}^3$, watering was carried out. After the wet substrate experiment, no further watering was performed until the soil was dry for the dry substrate experiment.

The experiment lasted for 2 hours. The duration of the experiment was determined based on several factors. If the duration was too brief, the impact of the living walls on indoor CO_2 removal would be insignificant, which would also lead to potential errors in data analysis and measurement. On the other hand, if the duration was too long, the decrease in the indoor CO_2 concentration could lead to the reduction of the CO_2 removal rate of the living walls. Previous studies have suggested a duration of 40 minutes for the measurement of the CO_2 removal rate of plants, as within this period, the decrease in the CO_2 concentration is linear [46,56,57]. However, in the present study, the experimental scale was expanded, and pre-experiments revealed that even under high light conditions, the indoor CO_2 concentration changed nearly linearly within 2 hours. Thus, the change in the indoor CO_2 concentration over 2 hours was measured to calculate the CO_2 removal rate of living walls.

To prepare for the experiment, the living walls were exposed to the experimental light intensities for 15 minutes to adapt to the experimental conditions. A CO_2 generator was created by placing dry ice in boiled water, and the fan was turned on for 10 minutes to evenly mix the indoor air. Once the indoor CO_2 concentration reached 1000 ppm on the air quality monitor, the CO_2 generator was removed, and the outer door was tightly closed. The same process was repeated for the control room. The initial CO_2 concentration set in this experiment was the standard limit of 1000 ppm. Due to the disorderliness of gas diffusion, the actual initial CO_2 concentration was 1000 ± 100 ppm. Once the indoor gas was uniformly mixed and the indoor CO_2 concentration no longer exhibited sudden changes, the timer was started. Each experiment lasted for 2 hours, and each condition was repeated three times.

Theoretically, the CO_2 removal rate from photosynthesis is influenced by the environmental CO_2 concentration. However, the range of the CO_2 concentration changes caused by living walls within 2 hours is limited, and thus the impact can be considered negligible. The decay of the indoor CO_2 concentration over 2 hours was regarded as linear. The data were analyzed using Origin 2021 software to perform linear regression analysis and minimize the impact of outliers. The CO_2 removal rate of the living walls was calculated. To illustrate the trend of CO_2 concentration changes and facilitate comparisons between different conditions, the indoor CO_2 concentration variation was expressed as a percentage of the initial concentration.

Statistical analysis was conducted on the data using the Shapiro-Wilk test to check for a normal distribution. If the data met the assumptions, analysis of variance (ANOVA) was performed. For non-normal data, a rank transformation was applied, and non-parametric tests were used. The Mann-Whitney test was used for the analysis of two independent samples, while the Kruskal-Wallis test was used for the analysis of more than two independent samples. A p-value above 0.05 indicates no significant differences between the groups.

2.1.3.3. Optimization of the impact of the significant factor of office spaces on the CO₂ removal efficiency

Among the adjustable factors of office spaces, light intensity significantly affects the CO₂ removal rate of the living walls. To analyze its influence, the light response curve of the living walls was measured in this study. The light saturation point (LSP) and light compensation point (LCP) of the plants were determined from the curve, offering valuable insights for the design of supplemental lighting.

The *E. aureum* living walls with a wet substrate were selected for the experiment, as they exhibited better performance in the previous experiments. The light intensities were set as, 0, 10, 50, 100, 200, 300, and 400 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$. The CO₂ removal rate of the living walls was measured using consistent experimental and data processing methods from previous experiments. The light response curve was fitted using a rectangular hyperbolic model described by the following equations [58,59]:

$$P_n = \frac{\alpha I P_{n\max}}{\alpha I + P_{n\max}} - R_d \quad (1)$$

where P_n is the net photosynthesis rate, I is the light intensity, α is the initial slope of the light response curve at $I=0$, which is known as the initial quantum efficiency, $P_{n\max}$ is the maximum net photosynthesis rate, and R_d is the dark respiration rate.

To estimate the light compensation point (I_{sat}), a linear equation was fitted to the data under low light conditions ($\leq 100 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$) to obtain the apparent quantum efficiency (AQE). The following equation was then solved [59].

$$P_{n\max} = AQE \times I_{sat} - R_d \quad (2)$$

2.2. Simulation

A simulation was carried out with the aim of accurately assessing the potential fresh air energy savings of the living walls in office spaces and providing a reference for practical applications. The optimal CO₂ removal rate of the living walls determined in the experimental research was utilized in the simulation. EnergyPlus was employed to simulate the reduction in energy consumption associated with fresh air intake when implementing living walls in office spaces. The simulation included various numbers of occupants and mechanical ventilation systems across seven climate regions in China.

2.2.1. Simulation conditions

The office room was modeled in OpenStudio 3.5.1 and imported into EnergyPlus 22.2.0. The work schedule for occupants was set from 08:00 to 17:00, 5 days a week [34]. An ideal HVAC system was used, and its equipment parameters and capacities were determined based on the corresponding Chinese Standard Weather Data (CSWD). The outdoor CO₂ concentration was set as 400 ppm [60], and the indoor CO₂ concentration was maintained below the standard limit of 1000 ppm [14]. The design parameters for indoor air were set to 26 °C in summer, 20 °C in winter, and a relative humidity of 60% [61]. In the simulation, a 1-minute time step was used to calculate fresh air energy consumption for each minute of the year. This study focused on fresh air energy consumption, which refers to the energy consumed by the HVAC system for heating, cooling, humidification, and dehumidification of outdoor air. The building orientation, envelope construction, and HVAC equipment were set to default parameters as they were not relevant.

2.2.2. Modeling description

An office room measuring 3.25 m × 7.70 m × 4.00 m (width × depth × height) and located on the middle floor of an office building, was considered as the case study (Figure 3). The room had one exterior wall and three interior walls, with two windows on the exterior wall measuring 2.75 m × 2.80 m (width × height). Two living walls were installed in the room, each measuring 1.2 m × 2.5 m (width

× height). The room was a typical mechanically ventilated office with natural infiltration set to 0.175 air changes per hour (ACH) [62].

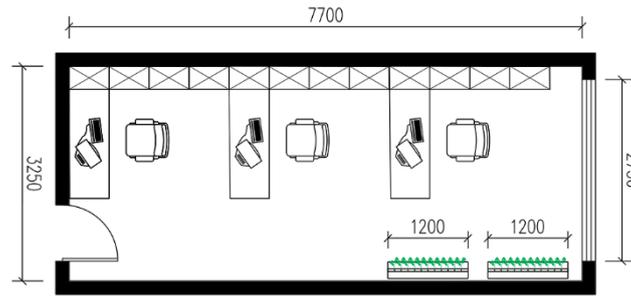


Figure 3. The plan of the simulated office.

2.2.3. Indoor CO₂ concentration

Assuming that the air is uniformly mixed in a room with a volume of V and there are no chemical reactions of contaminants, the CO₂ emission rate is \dot{m} , the outdoor CO₂ concentration is C_{out} , the indoor CO₂ concentration is C_{in} , and the ventilation rate of the room is Q . According to the law of conservation of mass, the differential equation for the change in indoor CO₂ concentration C after a very short time τ is as follows [63].

$$V \frac{dC}{d\tau} = Q(C_{out} - C_{in}) + \dot{m} \quad (3)$$

The analytical solution for the transient process of Equation (3) is given by the following [48,63]:

$$C_2 = C_1 \exp\left(-\frac{Q}{V}\tau\right) + \left(\frac{\dot{m}}{Q} + C_{out}\right) \left[1 - \exp\left(-\frac{Q}{V}\tau\right)\right] \quad (4)$$

where C_2 is the CO₂ concentration in the space after time τ , and C_1 is the initial CO₂ concentration in the room.

When $\tau \rightarrow \infty$, Equation (4) can be written as follows.

$$C_2 = C_{out} + \frac{\dot{m}}{Q} \quad (5)$$

Equation (5) indicates that the indoor CO₂ concentration reaches a steady state and becomes a constant value after a long period of time. Therefore, when the CO₂ emission rate (\dot{m}) and the acceptable indoor CO₂ concentration (C_{eq}) are known, the minimum ventilation rate (Q_m) of the room can be calculated as follows [64].

$$Q_m = \frac{\dot{m}}{C_{eq} - C_{out}} \quad (6)$$

The CO₂ emission rate (\dot{m}) in an office with occupants and living walls can be calculated as

$$\dot{m} = n \cdot V_{hum} - V_{lw} \quad (7)$$

where V_{hum} is the CO₂ generation rate from human respiration, n is the number of occupants in the room, and V_{lw} represents the CO₂ removal rate from the net photosynthesis of the living walls.

The CO₂ generation rate from human respiration can be calculated as follows [65]:

$$V_{hum} = \varepsilon \frac{0.202RQ \cdot M \cdot H^{0.725} \cdot W^{0.425}}{21(0.23RQ + 0.77)} \quad (8)$$

where ε is the correction coefficient, $\varepsilon = 0.85$ for adult males and $\varepsilon = 0.75$ for adult females. Moreover, RQ is the respiratory quotient, which is taken as 0.83 for light or sedentary activities in adults. M is the metabolic rate, which is 60 W/m² for sedentary activities, and H and W are respectively the height and weight of the occupant.

According to the 2022 report [66], the average height and weight of Chinese males aged 20-24 are 172.6 cm and 70.4 kg, respectively. This gives a CO₂ generation rate from human respiration of 452.4 mg/min.

In an office, occupants are the only source of CO₂ emissions. Before entering, indoor and outdoor CO₂ concentrations are equal. As occupants breathe, the indoor CO₂ concentration gradually rises. When there is a higher difference between the CO₂ concentrations of the indoor and outdoor environments, there is a faster exchange of gases. This exchange leads to a gradual slowing of the rate at which the indoor CO₂ concentration increases, until it reaches a steady state. The steady-state indoor CO₂ concentration can be calculated by Equation (5). Figure 4(a) presents the change in indoor CO₂ concentration without a ventilation system. Without any control measures, natural infiltration in an office with occupants leads to the steady-state CO₂ concentration exceeding the standard limit. For example, when three occupants work in the office, the steady-state CO₂ concentration is 2996 ppm, which is approximately three times higher than the standard limit.

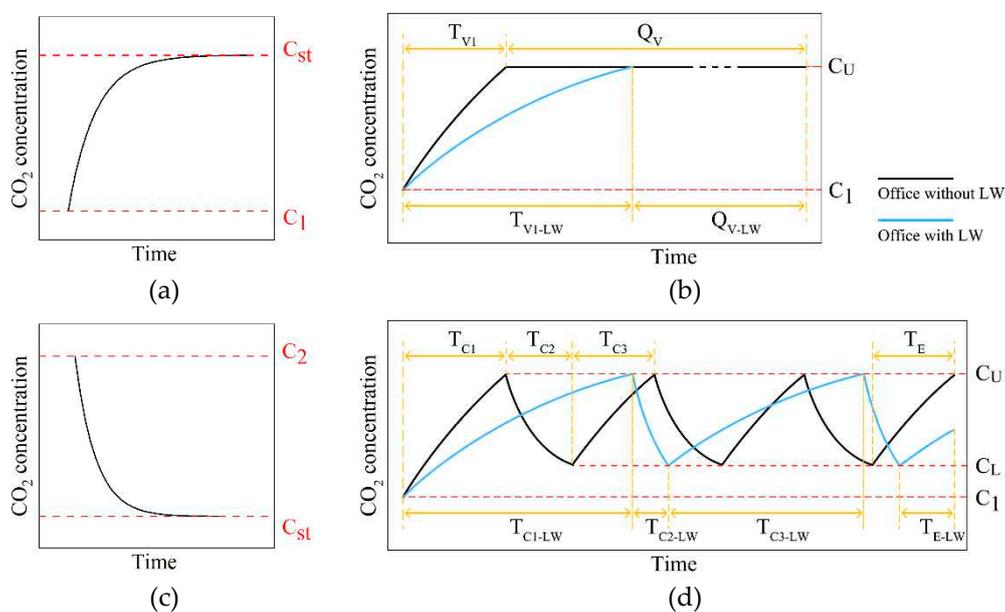


Figure 4. The changes in the indoor CO₂ concentration (a) before the ventilation turns on, (b) during working hours with a VAV system, (c) after the CAV system turns on, (d) and during working hours with a CAV system. Note: C_1 is the initial CO₂ concentration, C_{st} is the steady-state CO₂ concentration, and C_U and C_L respectively denote the upper and lower limits of the CO₂ concentration of the CAV system.

2.2.4. Mechanical ventilation types

To maintain a healthy indoor working environment, a ventilation system can be employed to introduce outdoor air into the office, thereby diluting the indoor CO₂ concentration to an acceptable level. Demand-controlled ventilation (DCV) based on CO₂ is an intelligent ventilation strategy that can improve indoor air quality and reduce energy consumption. This strategy entails the installation of CO₂ concentration sensors in the room, which activate the ventilation system when the CO₂ concentration level exceeds the standard limit [67]. Generally, DCV automatically adjusts ventilation rates based on the indoor CO₂ concentration, which requires the support of a variable air volume (VAV) system [68]. However, mechanical ventilation in many individually controlled or split air-conditioned offices is often designed with a constant air volume (CAV) [69]. Because living walls might be used with both systems, both types of mechanical ventilation systems, namely VAV and CAV, were investigated.

In offices with VAV systems with living walls, plants absorb the CO₂ from occupants through photosynthesis, which lowers the rate of increase of indoor CO₂ concentration (Equation (7)). As a result, the time it takes for the indoor CO₂ concentration to reach the standard limit is postponed

(Equation (4)) (in Figure 4(b), the time is postponed from T_{V1} to T_{V1-LW}), and the required ventilation rates to maintain indoor CO_2 concentration within the standard limit decreases (Equation (6)) (in Figure 4(b), the ventilation rate decreases from Q_V to Q_{V-LW}). This results in a decrease in the required outdoor air volume during working hours and a reduction in the energy consumption of the ventilation system.

In offices with CAV systems, it is necessary to set upper and lower limits of the indoor CO_2 concentration (Figure 4(d)). The upper limit is the standard limit, while the setting of the lower limit should be determined based on the actual needs, energy consumption, and equipment durability [69]. Market research shows that there are two CAV systems suitable for 30-m² offices, with ventilation rates of 120 m³/h and 210 m³/h, respectively. According to the derivation of Equation (4), a larger difference between indoor and outdoor CO_2 concentrations leads to a faster decay rate of the indoor CO_2 concentration (Figure 4(c)); thus, setting a higher lower limit can achieve greater energy reduction. To ensure the durability of the equipment, the CAV system should not be turned on and off with an interval of less than 10 minutes [70]. Considering these three requirements, a CAV system with ventilation rates of 120 m³/h and a lower CO_2 concentration limit set to 900 ppm was set in the simulation. The ventilation system operated with a minimum interval of 12 minutes between on and off cycles.

In offices with CAV systems with living walls, the CO_2 concentration changes similarly to that in offices with VAV systems before the ventilation system is activated. The ventilation system operates at a constant ventilation rate and reduces the indoor CO_2 concentration. When the CO_2 concentration reaches the lower limit, the ventilation system stops, and the indoor CO_2 concentration subsequently rises to the upper limit. One cycle is completed when the indoor CO_2 concentration changes from the upper limit to the lower limit and back to the upper limit. After this, a new cycle begins. Living walls prolong the time needed to reach the upper limit (in Figure 4(d), the time is prolonged from T_{C1} to T_{C1-LW} and from T_{C3} to T_{C3-LW}) and shorten ventilation system operation (in Figure 4(d), the time is shortened from T_{C2} to T_{C2-LW}). Living walls may also reduce the number of cycles within working hours. Notably, it is more energy-efficient if the working hours finish when the indoor CO_2 concentration reaches the standard limit and the CAV system is about to turn on (in Figure 4(d), $T_E = T_{C3}$). Living walls may alter CO_2 levels at the end of working hours, thereby impacting energy efficiency. EnergyPlus can calculate the difference in fresh air energy consumption in 1-minute intervals.

2.2.5. Climate and locations

Typical cities were selected based on the Climate Region of Building (Figure 5), and their corresponding meteorological conditions are provided in Table 3. In the field of civil buildings in China, there are two climate classification methods, namely the Climate Region for Building Thermal [71] and the Climate Region of Building [72]. The former divides China into five zones based solely on the average temperatures of the coldest month (January) and the hottest month (July), while the latter considers both these factors and the average relative humidity in July, resulting in a division of the country into seven zones. Fresh air energy consumption in a building is influenced by both the outdoor temperature and humidity. Therefore, fresh air energy consumption in the selected typical cities was calculated in EnergyPlus based on the Climate Region of Building.

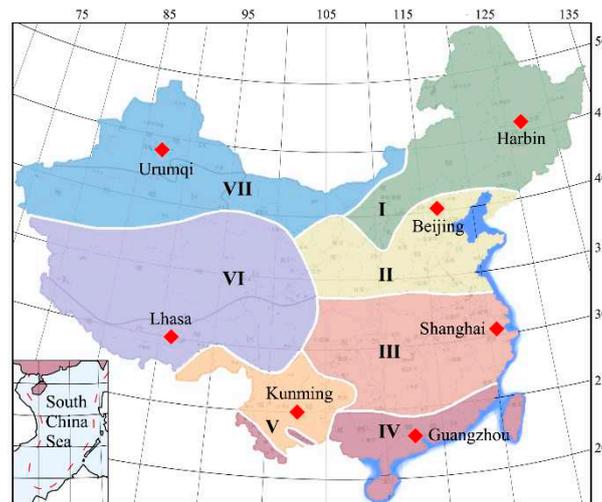


Figure 5. The division of the Climate Region of Building in China and typical cities.

Table 3. The outdoor meteorological parameters of typical cities in different climate regions [51].

Climate Region	City	Average Temperature in Winter (°C)	Dry Bulb Temperature in Summer (°C)	Relative Humidity in Summer	Atmospheric Pressure in Summer (kPa)	Moisture Content in Summer (g/kg)	Enthalpy in Summer (kJ/kg)
I	Harbin	-27.1	30.7	62%	98.8	17.7	76.2
II	Beijing	-9.9	33.5	61%	100.0	20.2	85.7
III	Shanghai	-2.2	34.4	69%	100.5	24.1	96.5
IV	Guangzhou	5.2	34.2	68%	100.0	23.6	95.0
V	Kunming	0.9	26.2	68%	80.8	18.3	73.0
VI	Lhasa	-7.6	24.1	38%	65.3	11.0	52.4
VII	Urumqi	-23.7	33.5	34%	91.1	12.2	65.2

3. Results

3.1. Experimental research

3.1.1. Confirmation of identical natural infiltration in two rooms

The confirmation results are reported in Table 4. The indoor CO₂ concentration in the two experimental rooms decreased at different rates within 2 hours due to variable weather conditions, including the wind speed, the wind direction, and indoor-outdoor temperature differences. These factors affected the calculation of the CO₂ removal rate of the living walls. Nevertheless, the changes in the indoor CO₂ concentration between Rooms A and B were relatively small during the same period. The accuracy of the CO₂ sensor (SenseAir S8) used in the experiment is ± 40 ppm, and the difference in indoor CO₂ concentration changes between the two rooms was within this range. Therefore, it was assumed that the two rooms had the same natural infiltration rate of CO₂. The decrease in the CO₂ concentration in Room B without living walls can be used as a substitute for the natural infiltration rate of CO₂ in Room A with living walls. By subtracting the natural infiltration rate in Room A from the total decrease in the CO₂ concentration, the CO₂ removal rate of the living walls could be calculated.

The comparative experiment in the two rooms helped to reduce the impact of variable outdoor meteorological conditions on the experimental results, resulting in more precise outcomes. Furthermore, the experiment was not constrained by season, weather, or local solar time, which enhanced the flexibility of the selection of the experimental time.

Table 4. The confirmation results of the two experimental rooms.

Date	Wind Speed (m/s)	Leading Wind Direction	Outdoor and Indoor Temperature Difference (°C)		CO ₂ Concentration (ppm)		CO ₂ Concentration Difference (ppm)	
			Room A	Room B	Room A	Room B		
			1	2022/10/14 11:00-13:00	2.3	WSW/SSW/W NW		4.1
2	2022/10/15 0:00-2:00	2.3	WSW/SSW	4.6	3.5	47	49	2
3	2022/10/16 10:30-12:30	4.6	WSW/SSW	4.9	3.8	164	182	18

3.1.2. Optimization of the impacts of the adjustable factors of office spaces on CO₂ removal efficiency

The change in the indoor CO₂ concentration within 2 hours is shown in Figure 6. Among the 16 experimental conditions, the indoor CO₂ concentration decreased in eight conditions, remained stable under two conditions (the range of CO₂ concentration change is within the reference line of $y=100\%$), and increased under six conditions. The *E. aureum* living walls respectively exhibited the highest and lowest indoor CO₂ removal rates within a 2-hour period under high light intensity and dark conditions with a wet substrate. Additionally, the highest CO₂ removal rate was associated with a slower reduction in the indoor CO₂ concentration, potentially indicating a decrease in the net photosynthesis rate as the indoor CO₂ concentration decreased.

The results of the CO₂ removal rates of the living walls are shown in Figure 7. The statistical analysis indicated no significant differences ($p > 0.05$) in the impacts of CAM plants represented by *S. trifasciata*, C₃ plants represented by *E. aureum*, and dry/wet substrates on the CO₂ removal rates. However, significant differences ($p < 0.05$) were observed between the medium/high light intensity and low light intensity/dark conditions, highlighting the greater influence of light intensity on CO₂ removal rates as compared to the carbon fixation pathways, and substrate moisture level.

The maximum CO₂ removal rates for plants with different carbon fixation pathways were observed under high light intensity, with the removal rates of *S. trifasciata* and *E. aureum* reaching 795 and 1176 mg/h, respectively. As the light intensity increased from medium to high, the CO₂ removal rates exhibited varying degrees of improvement. For *S. trifasciata*, the rates respectively increased by 134.9% and 20.3% under the dry and wet substrate conditions. For *E. aureum*, the rates respectively increased by 91.8% and 98.8% in the dry and wet substrate conditions. However, when transitioning from the dark conditions to low light intensity, the CO₂ removal rate of the *E. aureum* living walls increased, while that of the *S. trifasciata* living walls decreased. This observation may be attributed to the CAM carbon fixation pathway, in which the photo-related signal inhibits the CAM metabolism [73].

The maximum CO₂ removal rates for plants with different carbon fixation pathways occurred under varying substrate moisture levels. The C₃ plant *E. aureum* thrives in a well-hydrated soil environment, while the CAM plant *S. trifasciata* can adapt to drought conditions. Due to insufficient moisture, under the medium/high light intensity conditions, the CO₂ removal rates for the *E. aureum* living walls decreased by 18.4% and 21.3%, respectively. Under medium light intensity, the *S. trifasciata* living walls exhibited a 22.2% reduction in the CO₂ removal rate due to water deficiency. Surprisingly, under high light intensity, *S. trifasciata* living walls showed a 52.0% increase in the CO₂ removal rate despite an inadequate substrate moisture supply. CAM is a typical ecophysiological adaptation of plants to arid conditions [74], and the experimental results confirm the drought tolerance of CAM plants.

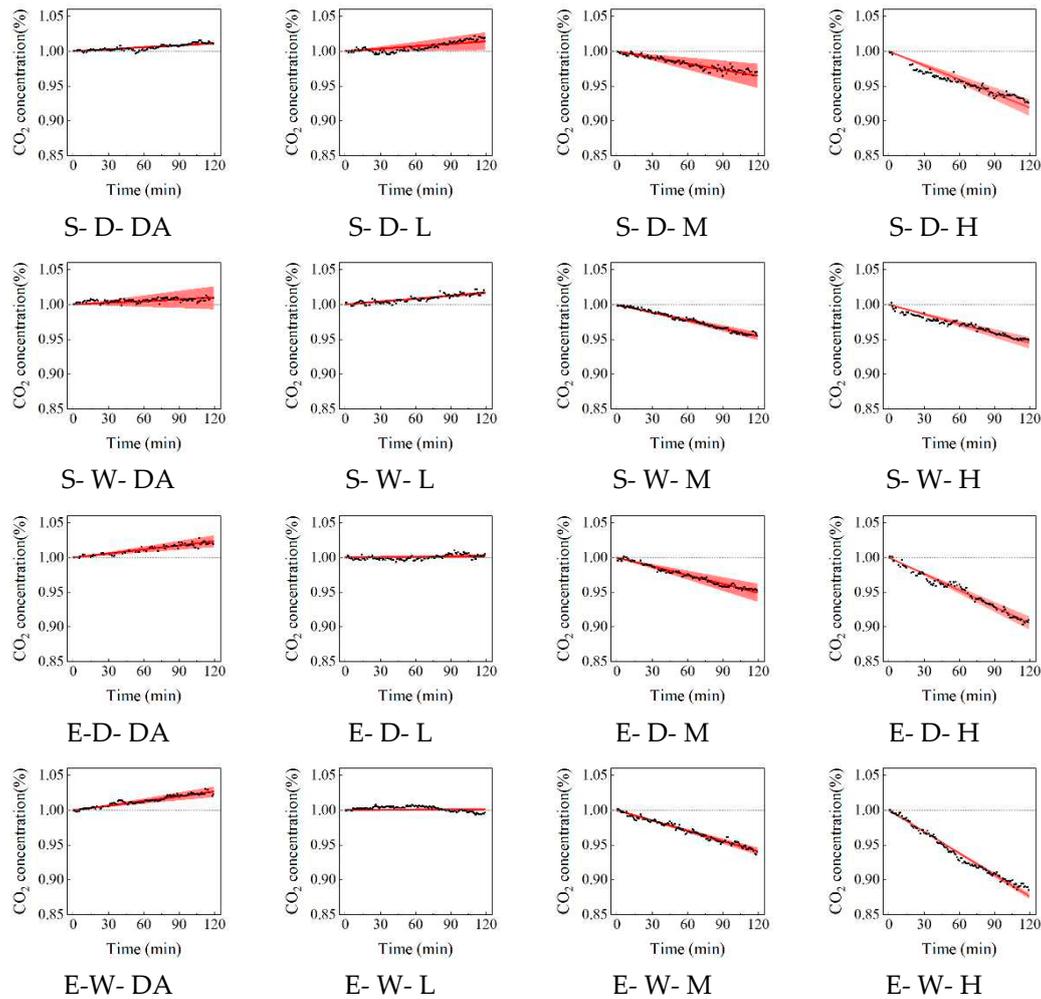


Figure 6. The mean CO₂ concentration (%) changes from the input concentration (1000±100 ppm) within 2 hours under different conditions: *S. trifasciata* (S), *E. aureum* (E), Dry (D), Wet (W), High light intensity (H), Medium light intensity (M), Low light intensity (L), Dark (DA). Shaded areas represent the SEM (n = 3).

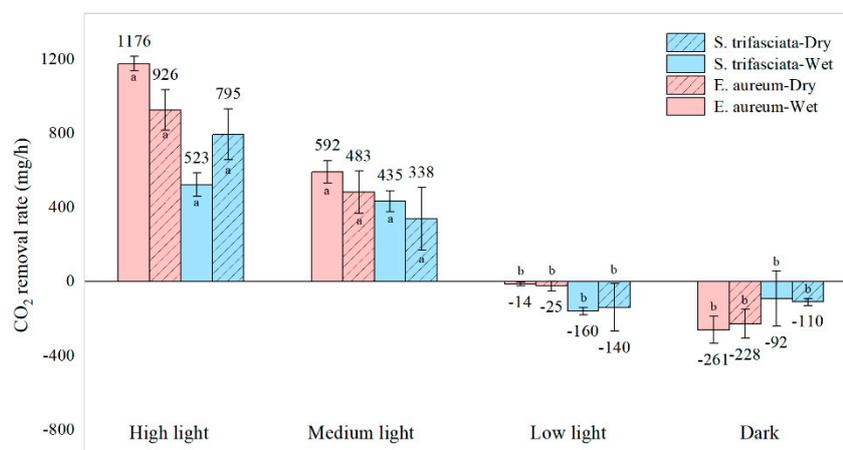


Figure 7. The CO₂ removal rates are presented as the mean ± SEM (n=3). Notes: lowercase letters indicate significant differences (p=0.05).

3.1.3. Optimization of the impact of the significant factor of office spaces on the CO₂ removal efficiency

The light response curve results of the *E. aureum* living walls in a wet substrate are shown in Figure 8. The calculation of the net photosynthetic rate (P_n) in relation to light intensity (I) is given by Equation (9).

$$P_n = \frac{37 \times I \times 1620}{37I + 1620} - 312 \quad (R^2=0.99267) \quad (9)$$

The linear regression result of net photosynthetic rate (P_n) in relation to light intensity (I) for plants under low-light conditions ($\leq 100 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$) were obtained by Equation (10).

$$P_n = 11I - 144 \quad (R^2=0.88337) \quad (10)$$

Based on Equation (2), the estimated the LSP is $176 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$, and the LCP is $13 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$.

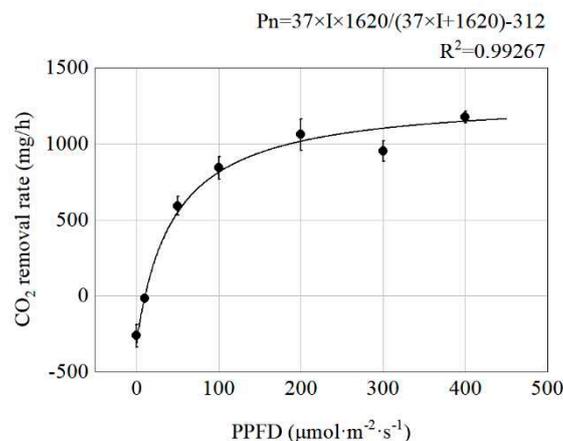


Figure 8. The light response curve of the *E. aureum* living walls in a wet substrate.

3.2. Simulation

3.2.1. CO₂ concentration and fresh air volume

Figs. 9 and 10 respectively display the variations of the CO₂ concentration variations and the fresh air volumes in the offices with the VAV or CAV systems, with different numbers of occupants, and with or without living walls on a typical working day.

When two and three occupants are working, living walls delay the activation of the ventilation system by 17 and 9 minutes, respectively. When only one occupant is in the office, living walls make it unnecessary to activate the ventilation system as the indoor CO₂ concentration is within acceptable levels.

In an office with a CAV system and living walls, the number of ventilation system activations remains unchanged with three occupants, but the operating time decreases by 14.3%. With two occupants, the number of activations decreases from 9 to 8, and the operating time decreases by 27.8%.

In an office with a VAV system and living walls, the fresh air volume decreases by 22.2% and 12.3% for two and three occupants, respectively. The ventilation rate required to control the CO₂ concentration within the standard limit (Equation (6)) decreases from 75.6 to 69.5 m³/h for three occupants and from 50.4 to 44.3 m³/h for two occupants.

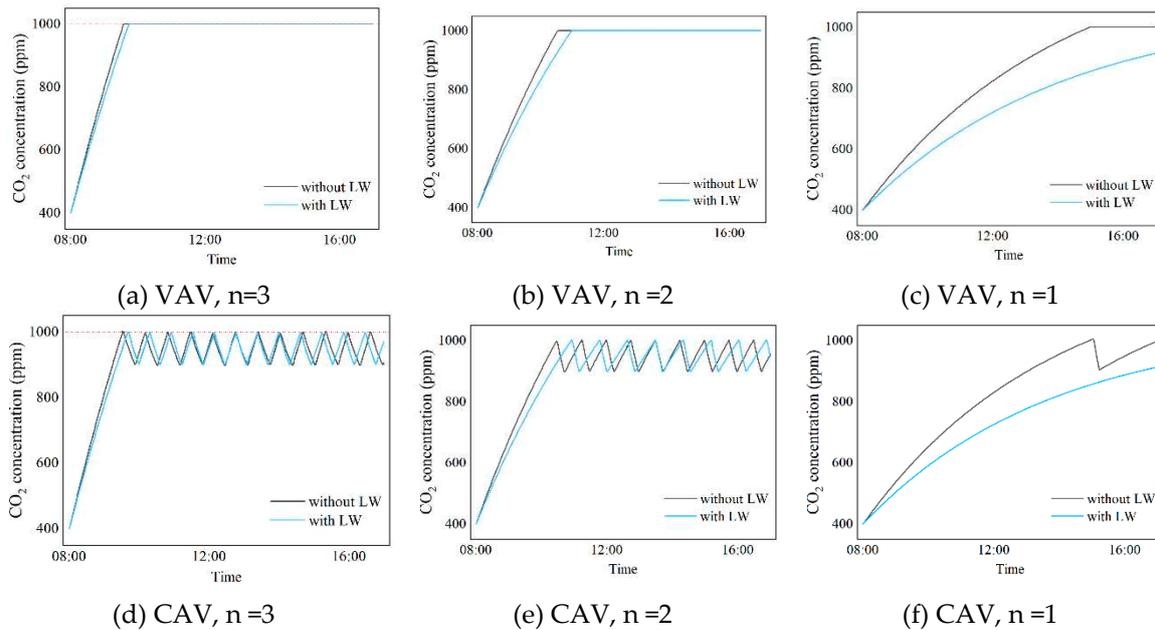


Figure 9. The variations of the CO₂ concentration in the offices with VAV or CAV systems, with different numbers of occupants, and with or without living walls, on a typical working day.

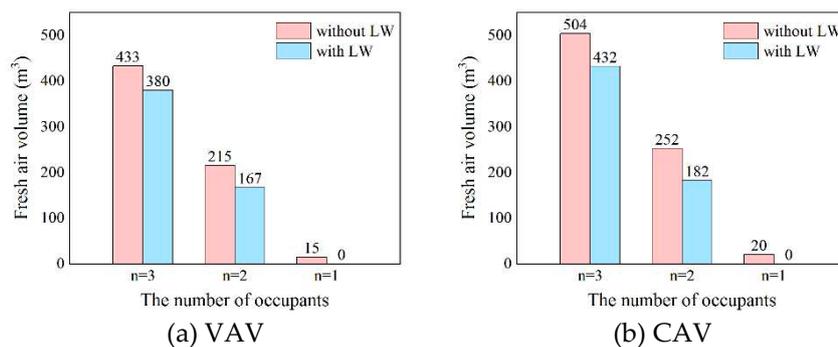


Figure 10. The fresh air volumes in the offices with VAV or CAV systems, with different numbers of occupants, and with or without living walls on a typical working day.

3.2.2. Fresh air energy consumption

Figure 11 presents the annual energy consumption for the cooling and heating of outdoor air in offices with VAV or CAV systems, with different numbers of occupants, and with or without living walls in typical cities. In offices with three occupants, living walls result in an annual reduction in fresh air energy consumption ranging from 17 to 93 kW·h, achieving energy savings of 11.2% to 14.8%. For two occupants, living walls contribute to an annual reduction in fresh air energy consumption ranging from 15 to 72 kW·h, achieving energy savings of 19.8% to 28.2%. Compared to VAV systems, CAV systems exhibit higher energy savings in terms of fresh air energy consumption when using living walls to lower indoor CO₂ concentrations, with CAV systems achieving 1.26 times the energy savings of VAV systems.

The fresh air energy savings achieved by living walls vary significantly due to differences in climate conditions. The city with the highest energy savings is Harbin, with 4.5 times the energy savings of the city with the lowest energy savings, namely Kunming. In humid regions, cities have a higher fresh air energy savings of 17.8% to 29.1% as compared to dry regions at the same temperature. Among cities with extremely cold climates and an average temperature below -10 °C in the coldest month, Harbin and Urumqi have different air humidities [71]. Harbin, with higher air humidity, has a fresh air energy savings of 17.8% to 19.6% higher than those of Urumqi. Beijing and Lhasa both require the consideration of both winter insulation requirements and summer heat prevention [71].

Because Beijing is closer to the ocean than Lhasa, its fresh air energy savings are 18.7% to 29.1% higher than those of Lhasa.

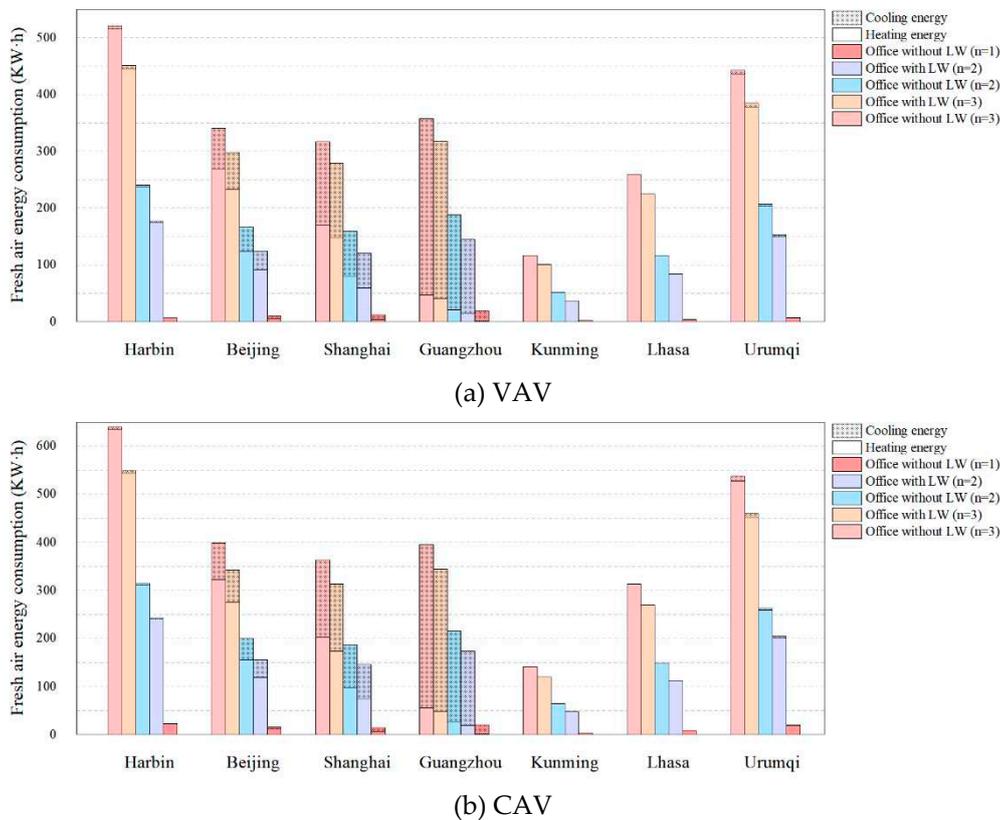


Figure 11. The annual energy consumption for outdoor air cooling and heating in the offices with VAV or CAV systems, with different numbers of occupants, and with or without living walls, in typical cities.

4. Discussion

4.1. Comparison to other methods of indoor CO₂ removal

Compared to absorption and membrane separation, adsorption has the potential for room-level CO₂ capture [17,75]. In high-density and poorly ventilated environments like airplanes and submarines, where tackling high indoor CO₂ levels is critical, high-cost and efficient CO₂ adsorption methods are required. Living walls are less efficient than adsorption methods for purifying CO₂.

In offices, occupant respiration increases indoor CO₂ levels, thus impacting work efficiency [41]. Ventilation rapidly reduces CO₂ concentrations but consumes significant HVAC energy. Given the synchronization of office occupancy with the natural light cycle, the utilization of natural light for photosynthesis to purify indoor air offers an energy-efficient and sustainable solution to mitigate high indoor CO₂ levels in office environments.

In addition, unlike adsorption, which solely captures CO₂, plants engage in simultaneous CO₂ absorption and O₂ release via photosynthesis. Prolonged work in an enclosed space can lead to decreased indoor O₂ levels, thus affecting cognitive abilities [76]. Plants not only purify indoor air but also generate beneficial air ions [77], remove contaminants such as volatile organic compounds (VOCs) [78,79] and particulate matter (PM) [80], enhance thermal comfort [81], and create a sense of safety and relaxation when in proximity to them [26]. Additionally, plants hold aesthetic value in architectural environments [82].

4.2. Optimization of the CO₂ removal efficiency in office spaces

The plant selection, lighting design, and irrigation management of the living walls are crucial for air purification in office spaces. This study experimentally investigated the impacts of the carbon fixation pathways, light intensity, and substrate moisture level on the CO₂ removal rate of the living walls, providing valuable insights for designers. Botanical research has examined the impacts of light conditions and soil moisture on the net photosynthetic rates of various plant species, as the CO₂ absorption rate reflects their physiological characteristics and productivity [83]. Researchers have explored the potential of plants to reduce indoor CO₂ concentrations in chambers [44] and conducted experiments in actual rooms [56], providing a theoretical basis for the implementation of living walls to lower indoor CO₂ levels. However, further targeted research is needed, particularly for practical applications in real office buildings.

The selection of plant species plays a vital role in the CO₂ removal rate of the living walls, with C₃ plants offering higher rates than CAM plants. Botanical studies have shown that C₃ plants exhibit higher productivity than CAM plants [84]. The experimental results of the present study demonstrate a 48.0% higher CO₂ removal rate in the C₃ plant, *E. aureum*, than in the CAM plant, *S. trifasciata*, under identical high lighting conditions. Moreover, even under similar wet substrate conditions, the *E. aureum* living wall under medium light intensity exhibited a higher CO₂ removal rate than the *S. trifasciata* living wall under high light intensity. This highlights the significant impact of plant selection on the CO₂ removal rate. Additionally, comparing the findings of the present study with those of the study conducted by Torpy et al. [56], the CO₂ removal rate of *Chlorophytum comosum* living walls was found to be 3.2 times higher than that of the *E. aureum* living walls, emphasizing the potential for substantial enhancement in the CO₂ removal rate via appropriate plant selection. While plant selection for the living walls often prioritizes aesthetic considerations [57], the present research provides an alternative perspective for interior designers.

The light intensity significantly influences the CO₂ removal rate of the living walls. Studies have shown that increasing light intensity enhances the CO₂ removal rate of plants [44,56,57,85,86], which is consistent with the findings of the present work. When transitioning from medium to high light intensity, the CO₂ removal rate of the living wall increased by 20.3% to 134.9%. However, plants have a saturation point, beyond which providing additional light no longer increases the CO₂ removal rate. For the *E. aureum* living walls in the wet substrate, the measured LSP was 176 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$. Therefore, when natural light exceeds the saturation point, artificial lighting is unnecessary. Additionally, researches indicate that under typical office lighting, most plants are unable to significantly reduce indoor CO₂ concentrations [43,44]. The results of the present research demonstrate that under typical office lighting conditions, the living walls released CO₂. The LCP of the *E. aureum* living walls in a wet substrate was measured as 13 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$. To prevent plants from adversely affecting indoor air quality, it is necessary to activate supplemental lighting when the natural light falls below the compensation point while people are present in the office.

The utilization of natural light to enhance CO₂ absorption in living walls is an energy-efficient solution for the reduction of indoor CO₂ levels in offices. If artificial lighting is used exclusively, energy-efficient lamps with a photon efficiency of 1.5 $\mu\text{mol}/\text{J}$ [87] would require 117 W and 9 W to reach the LSP and LCP, respectively. The annual lighting energy consumption required to achieve the LSP would be 2.96 times higher than the highest energy savings associated with ventilation. Shao et al. [42] measured the average daily PPFD inside a window sill at 250 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$, exceeding the LSP of the living wall. Therefore, our research suggests that living walls primarily rely on natural light, activating supplemental lighting when the surface light falls below the LCP of 13 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$. While areas with concentrated solar radiation in an office may impact human thermal comfort [35] and lead to equipment aging [36], they are suitable for placing living walls, thus providing a new perspective for interior design based on our research.

Irrigation and maintenance management are key considerations for the operation and management of living walls. Under typical office lighting conditions, substrate moisture has a negligible impact on plant CO₂ removal efficiency [44], as was confirmed in this study. However, water deficit can lead to stomatal closure [88] reducing the CO₂ removal rate of living walls with C₃

plants like *E. aureum* by 18.4% to 21.3% under medium and high light intensities, respectively. Therefore, irrigation management is crucial for C₃ plants used in air purification. In the present experiment, the plants were watered every 3 days to maintain optimal substrate moisture. For CAM plants, the CO₂ removal rate of the living walls remained unaffected by drought under high light intensity but decreased by 22.2% under medium light intensity. Choosing living walls with CAM plants for air purification allows for a reduction in irrigation frequency, thus lowering maintenance costs.

5. Conclusions

The utilization of living walls for CO₂ removal is an energy-efficient and sustainable solution to address high CO₂ levels in office spaces. To promote the practical application of the living walls and provide guidance to designers, this study experimentally investigated the impacts of adjustable factors of office environments on the CO₂ removal rate of the living walls. Additionally, EnergyPlus was innovatively employed to evaluate the fresh air energy-saving potential of the living walls under different scenarios. The following conclusions were drawn.

1. The selection of C₃ plants for living walls was found to achieve higher CO₂ removal efficiency than CAM plants.
2. Natural light is primarily used for CO₂ removal in living walls. The activation of artificial lighting is recommended when the surface light intensity of living walls drops below the LCP of 13 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$.
3. The reduction of the irrigation frequency was found to have no impact on the CO₂ removal efficiency of CAM plant living walls, but decreased the CO₂ removal efficiency by 18.4%-21.3% for C₃ plant living walls.
4. In a 30-m² office with two and three occupants, living walls can reduce the fresh air demand by 12.3%-27.8% and decrease fresh air energy consumption by 11.2%-28.2%.
5. The fresh air energy savings of the living walls varied significantly across climate regions. The city with the highest savings, Harbin, was found to have 4.2 times the energy savings of Kunming, the city with the lowest savings. Humid regions were found to exhibit 17.8%-29.1% higher energy savings as compared to dry regions.

This study focused on the optimization of CO₂ removal via living walls for office spaces. However, further research is needed to explore their application in residential, educational, and commercial buildings. Additionally, while this study accurately calculated the fresh air energy savings of living walls, future research should integrate other lighting analysis software to evaluate the overall energy-saving effects of the living walls.

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