

#### Article

# Analysis of carbon fixation and humidification ability of indoor cultivation of *Ficus pandurata Hance*

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Copyright © 2025 by author(s). *Eco Cities* is published by Asia Pacific Academy of Science Pte. Ltd. This work is licensed under the Creative Commons Attribution (CC BY) license. https://creativecommons.org/licenses/ by/4.0/ Abstract: To investigate the impact of prolonged exposure to enclosed, low-light environments on carbon fixation and oxygen release in green plants, as well as their capacity to regulate transpiration and humidification, this study utilized *Ficus pandurata Hance*, a common indoor ornamental plant, as the experimental subject to examine the net photosynthetic rate ( $P_n$ ) and transpiration rate ( $T_r$ ). LI-6800 portable photosynthesizer was employed to assess the  $P_n$  and  $T_r$  of Ficus pandurata Hance cultivated under varying temperatures (15 °C, 20 °C, 25 °C, 30 °C, 35 °C) and different CO<sub>2</sub> concentrations (400  $\mu$ mol·mol<sup>-1</sup>, 800  $\mu$ mol·mol<sup>-1</sup>, 1200  $\mu$ mol·mol<sup>-1</sup>) at different parts of the room (indoors or near windows). The results of the light response curve and the CO<sub>2</sub> response curve measurements indicate that the  $P_n$  of Ficus pandurata Hance shows a trend of initially increasing and then decreasing as the light intensity or CO<sub>2</sub> concentration increases. It is noteworthy that under different photosynthetically active radiation (PAR) and  $CO_2$  concentrations, the maximum  $P_n$ of Ficus pandurata Hance cultivated by a window is significantly higher than that of indoorcultivated plants. Under appropriate temperature control (20~30 °C), the  $P_{\rm n}$  and  $T_{\rm r}$  of Ficus pandurata Hance are highest at 800  $\mu$ mol·mol<sup>-1</sup> CO<sub>2</sub> concentration. Under appropriate ventilation conditions (CO<sub>2</sub> concentration  $< 1200 \,\mu \text{mol} \cdot \text{mol}^{-1}$ ), the plants have stronger carbon fixation ability under appropriate temperature conditions and stronger transpiration-induced humidification ability under non-low temperature (T  $\geq$  20 °C) conditions. To sum up, in the case of high CO<sub>2</sub> concentration caused by poor indoor ventilation and dense population, cultivation of Ficus pandurata Hance by the window and proper control of temperature above 20 °C can obtain good ecological benefits of carbon fixation, oxygen release, transpiration and humidification.

**Keywords:** *Ficus pandurata Hance*; net photosynthetic rate; transpiration rate; temperature; carbon dioxide

# 1. Introduction

The indoor thermal environment and air quality are pivotal factors that significantly influence working conditions, as well as the health and productivity of occupants. In contemporary society, the extension of human living and working hours in indoor environments—particularly during and following the COVID-19 pandemic—has resulted in a heightened demand for improved indoor thermal comfort and air quality among occupants [1,2]. The incorporation of vegetation not only enhances aesthetic appeal but also markedly improves indoor environmental quality through the intrinsic capabilities of plants—including carbon fixation, oxygen production, transpiration, and humidification [3,4]. Plants such as *Spathiphyllum wallisii, Hedera helix, Chlorophytum comosum*, and *Epiremnum aureum* can all improve indoor environments to varying degrees [5,6]. In addition to the above

characteristics, many indoor plants can also stagnate and absorb toxic and harmful substances [7].

The building exhibits a variety of characteristics that influence indoor environmental conditions, including inadequate air circulation, inconsistent lighting levels, and temperature differentials between indoor and outdoor spaces resulting from the use of air conditioning in summer and heating in winter. Additionally, factors such as occupancy density, ventilation conditions, and the spatial arrangement of green plants also play significant roles [8]. Collectively, these elements affect variations in indoor light intensity, temperature fluctuations, and CO<sub>2</sub> concentration to differing extents. Consequently, the ecological benefits derived from carbon fixation, oxygen release, transpiration, humidification, and cooling provided by indoor green plants are similarly impacted [9]. And light, temperature, and other environmental factors are important factors affecting plant growth and survival [9,10]. Therefore, identifying optimal indoor environmental conditions that align with the growth requirements of green plants is essential for maximizing their ecological contributions. In response to these challenges, this study focuses on Ficus pandurata Hance as a research subject to investigate its gas exchange responses under varying light intensities, temperatures, and  $CO_2$  concentrations. The objective is to determine the ideal cultivation environment that enhances its photosynthetic carbon fixation capabilities alongside oxygen release while effectively contributing to transpiration humidification and cooling processes. This research aims to provide empirical data supporting the maximization of ecological benefits offered by Ficus pandurata Hance within indoor settings.

### 2. Materials and methods

# 2.1. Site

The experiment was conducted from early April to mid-May. The experimental site is situated in the south-facing office of the school, with dimensions of 7.5 m × 3.5 m × 2.5 m. This office features a single floor-to-ceiling window located on its southern wall, measuring 1.2 m × 2.5 m. The photosynthetically active radiation (PAR) near the window was approximately 400  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>, while that within the room measured around 10  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>. The average indoor daytime temperature was approximately 25 °C, and the average nighttime temperature was about 18 °C.

### 2.2. Material

The commonly utilized green plant, *Ficus pandurata Hance*, was selected for laboratory experiments. This small shrub, classified within the genus Ficus of the Moraceae family, is frequently employed as an indoor ornamental plant due to its considerable aesthetic appeal. The selected plants were uniform in size and exhibited optimal growth conditions, being free from dead leaves, decay, or any signs of pest and disease infestation.

#### 2.3. Method

Three pots of the plant were placed near the window for cultivation (windowcultivated), while 3 pots were positioned indoors (indoor-cultivated—placed on the north wall of the office, away from the window). Watering was conducted in accordance with the dry and wet conditions of the cultivated soil. The experiment commenced after a maintenance period of 1 to 2 weeks. LI-6800 portable photosynthesis system (Li-COR, USA) was utilized to measure relevant indicators daily from 08:00 to 12:00 [9]. During the experiment, all doors and windows remained closed, with only one person present indoors to minimize personnel movement. The changes in indoor environmental conditions under these circumstances are presented in **Table 1**.

Table 1. Diurnal variation of CO<sub>2</sub> concentration in a closed chamber.

| Five-day average | CO <sub>2</sub> concentration in the morning | CO <sub>2</sub> concentration at nightfall | The difference of CO <sub>2</sub> concentration between the morning and nightfall |
|------------------|--|--|---|
|                  | $(\mu mol \cdot mol^{-1})$                   | $(\mu mol \cdot mol^{-1})$                 | $(\mu mol \cdot mol^{-1})$  |
|                  | $480.02 \pm 37.94$                           | $828.15\pm56.73$                           | $348.13 \pm 51.68$  |

When determining, select healthy, vigorous, and clean leaves, and set up 3–5 parallel for each treatment.

(1) Light response curve

The measurement period for the light response curve was conducted from 09:00 to 11:00, the airflow rate was maintained at 500  $\mu$ mol·s<sup>-1</sup> [11,12]. Measurements of the light response curve were performed at room temperature with a CO<sub>2</sub> concentration of 400  $\mu$ mol·mol<sup>-1</sup>. Based on preliminary experimental results indicating significant photoinhibition under high light conditions, the initial value of PAR for the light response curve was established at 1500  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>. Following data stabilization, an automated measurement protocol was employed, specifically setting PAR values to 1500, 1300, 1100, 1000, 800, 700, 600, 550, 500, 400, 300, and down to zero  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>. A stabilization time of between 120 and 180 seconds was implemented.

(2) CO<sub>2</sub> response curve

Based on the results of the light response curve analysis (**Figure 1**), PAR was established at 500  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>, with a flow rate maintained at 500  $\mu$ mol·s<sup>-1</sup>. The induction period under a CO<sub>2</sub> concentration of 400  $\mu$ mol·mol<sup>-1</sup> was set to 6–8 min. Following data stabilization, an automated measurement protocol was initiated. Carbon dioxide was supplied from a CO<sub>2</sub> cylinder, and concentrations were adjusted to 400, 300, 200, 100, 50, 25, and zero as well as subsequently to 400, 700, 900, 1000, 1500, and 2000  $\mu$ mol·mol<sup>-1</sup>; the stabilization time for each setting ranged from 120 to 180 s.

(3) The net photosynthesis and transpiration rates of *Ficus pandurata Hance* under different temperatures and  $CO_2$  concentrations

Based on the measurement of light response curves, the correlation measurements under different temperatures and CO<sub>2</sub> concentrations were carried out under 500  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup> PAR, and the flow rate was set to 500  $\mu$ mol·s<sup>-1</sup>. Measure the net photosynthetic rate ( $P_n$ ) and transpiration rate ( $T_r$ ) of *Ficus pandurata Hance* window-cultivated under different CO<sub>2</sub> concentrations (400  $\mu$ mol·mol<sup>-1</sup>, 800

 $\mu$ mol·mol<sup>-1</sup>, 1200  $\mu$ mol·mol<sup>-1</sup>, respectively simulating ventilation, indoor people and ventilation or indoor people and no ventilation, and indoor people and no ventilation), and different temperatures (35 °C, 30 °C, 25 °C, 20 °C, 15 °C, respectively simulating high summer temperature, middle-high summer temperature, spring or autumn or winter with cooling in summer, spring or autumn or winter with moderate temperature ventilation and heating, and winter with low temperature ventilation and heating). The data is stabilized for 3–5 min before counting. For each treatment, 1–2 leaves of similar size, similar growth and basically the same position were selected from each pot of plants for measurement (a total of 3–5 parallel leaves were set for each treatment).

# 3. Results and analysis

This section may be divided by subheadings. It should provide a concise and precise description of the experimental results, their interpretation, as well as the experimental conclusions that can be drawn.

#### 3.1. Light response curve

**Figure 1a** shows  $P_n$  curve of *Ficus pandurata Hance* under light-compensation point at different cultivation positions within the range of 0–100 µmol·m<sup>-2</sup>·s<sup>-1</sup> PAR, in order to better understand the light compensation point of different cultivation positions of *Ficus pandurata Hance* under low light intensity.



**Figure 1.** Light response curves of *Ficus pandurata Hance* at different culture locations: (a) The photosynthetic rate curve from 0 to 100  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>. (The solid line represents the measured value, and the dotted line is the fitting curve [13].

As can be seen from **Figure 1**, the changes of  $P_n$  of *Ficus pandurata Hance* at each culture position in the room showed that with the increase of light intensity, the  $P_n$  increased first and then decreased. When the PAR was lower than 200µmol·m<sup>-2</sup>·s<sup>-1</sup>, the  $P_n$  of *Ficus pandurata Hance* cultured at different locations increased rapidly with the increase of light intensity. When PAR was higher than 200µmol·m<sup>-2</sup>·s<sup>-1</sup>, the plants'  $P_n$  could continue to increase, but the growth rate became slow. The plant windowcultivated exhibited a maximum  $P_n$  of 8.27 µmol·m<sup>-2</sup>·s<sup>-1</sup> at 800 µmol·m<sup>-2</sup>·s<sup>-1</sup> PAR, whereas the indoor-cultivated plant reached its peak rate of 5.09  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup> at a lower light intensity of 500  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>. Subsequently, the  $P_n$  declined with increasing light intensity.

The  $P_n$  of *Ficus pandurata Hance* window-cultivated is significantly higher than that of those grown indoors. Prior to reaching 800 µmol·m<sup>-2</sup>·s<sup>-1</sup> PAR, the  $P_n$  of the window-cultivated plants exhibits continuous growth. Between 800 µmol·m<sup>-2</sup>·s<sup>-1</sup> and 1000 µmol·m<sup>-2</sup>·s<sup>-1</sup>, this rate remains relatively stable. As light intensity continues to increase, the  $P_n$  gradually declines, ultimately aligning with that of indoor-cultivated plants when PAR reaches 1500 µmol·m<sup>-2</sup>·s<sup>-1</sup> (the  $P_n$  value for window-cultivated plants is 1.04 µmol·m<sup>-2</sup>·s<sup>-1</sup>; for indoor-cultivated plants, it is 1.00 µmol·m<sup>-2</sup>·s<sup>-1</sup>).

By fitting the curve with the Ye model [13], it can be seen that the light saturation point (LSP) of *Ficus pandurata Hance* window-cultivated was 718µmol·m<sup>-2</sup>·s<sup>-1</sup>, while the LSP of *Ficus pandurata Hance* indoor-cultivated is 311  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>. This indicates that the LSP of *Ficus pandurata Hance* window-cultivated is significantly higher than that of the indoor-cultivated individuals. As illustrated in Figure 1a, the light compensation point (LCP) of Ficus pandurata Hance window-cultivated (fitted result < 0; observed value: 3.736  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>) is slightly lower than that of individuals indoor-cultivated (LCP, fitted result: 7.786  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>; observed value: 9.668  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>). Under low PAR (0–100  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>), *Ficus pandurata Hance* from different lighting conditions exhibited comparable  $P_n$ , with indoor-cultivated plants showing a marginally higher rate than those window-cultivated. Additionally, from Figure 1, it can be observed that *Ficus pandurata Hance* exhibits a clear photoinhibition phenomenon. The plant window-cultivated can only show photoinhibition at a higher PAR level of 1000 µmol·m<sup>-2</sup>·s<sup>-1</sup>, while the indoorcultivated plant shows a significant decrease in  $P_n$  at a PAR level above 500  $\mu$  mol·m<sup>-2</sup>·s<sup>-1</sup>.

Due to its strong adaptability to light, Ficus pandurata Hance can thrive in both high and low light conditions, making it a popular choice for indoor ornamental cultivation [14]. Experimental results indicate that the light compensation point of Ficus pandurata Hance indoor-cultivated is relatively low (approximately 20  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup> for most shrubs and trees on average [15]). The response of *Ficus* pandurata Hance to light varies significantly depending on its indoor cultivation location. In addition, plants positioned near windows exhibit a higher light saturation point compared to those situated further inside the room, and they also demonstrate marked differences in their responses to high light intensities. This further suggests that the adaptability of *Ficus pandurata Hance* to light varies significantly based on its cultivation location. The light saturation point for Ficus pandurata Hance grown near windows can approximate that of many outdoor shrubs and trees (762.3  $\mu$ mol·m<sup>-2</sup>·s<sup>-1</sup>) [12]. Consequently, when cultivating *Ficus pandurata Hance* indoors, those positioned by windows exhibit enhanced adaptability and efficiency in light utilization, thereby demonstrating superior carbon fixation and oxygen release capabilities.

#### **3.2.** CO<sub>2</sub> response curves

Figure 2b presents P<sub>n</sub> across CO<sub>2</sub> concentrations ranging from 0 to 100

 $\mu$ mol·mol<sup>-1</sup>, facilitating a clearer understanding of the CO<sub>2</sub> compensation points associated with different cultivation locations of *Ficus pandurata Hance*.



**Figure 2.**  $CO_2$  response curves of *Ficus pandurata Hance* at different culture locations: (b) shows the photosynthetic rate when  $CO_2$  concentration is from 0 to 100  $\mu$ mol·mol<sup>-1</sup> (The solid line denoted the measured values and the dashed line represented the fitted curve [16].

As illustrated in **Figure 2**, the  $P_n$  of *Ficus pandurata Hance* cultivated under various environmental conditions demonstrated a pattern where the rate initially increased with rising CO<sub>2</sub> concentrations before subsequently declining. Notably, the maximum  $P_n$  was achieved at a CO<sub>2</sub> concentration of 1500 µmol·mol<sup>-1</sup>, reaching 10.10 µmol·m<sup>-2</sup>·s<sup>-1</sup> for plants window-cultivated, and indoor-cultivated plants was 5.63µmol·m<sup>-2</sup>·s<sup>-1</sup>. When CO<sub>2</sub> concentrations exceed 1500 µmol·mol<sup>-1</sup>, the  $P_n$  of indoor-cultivated plants exhibits a significant decline, whereas the  $P_n$  of plants window-cultivated levels off gradually, with a less pronounced downward trend. Meanwhile, as illustrated in **Figure 2b**, the CO<sub>2</sub> compensation point (LCP; fitted result <0, observed value 9.543 µmol·mol<sup>-1</sup>) for plants window-cultivated is lower than that of those grown indoors (LCP; fitted result 10.999 µmol·mol<sup>-1</sup>, observed value 14.390 µmol·mol<sup>-1</sup>). Throughout the entire range of CO<sub>2</sub> measurement concentrations, the  $P_n$ of plants window-cultivated consistently exceeds that of their indoor counterparts.

Throughout the process of increasing CO<sub>2</sub> concentrations, the  $P_n$  of *Ficus* pandurata Hance exhibited a gradual increase, suggesting that under conditions of limited air circulation indoors, higher CO<sub>2</sub> concentrations correlate with enhanced photosynthetic oxygen release capabilities. Furthermore, the  $P_n$  of plants window-cultivated is consistently significantly higher than that of those indoor-cultivated across the entire range of CO<sub>2</sub> concentrations, indicating that window-side cultivation more effectively enhances the carbon fixation and oxygen-releasing capacity of *Ficus* pandurata Hance. Additionally, the compensation point for plants window-cultivated is lower than that for indoor-cultivated plants, signifying a greater ability to utilize

CO<sub>2</sub>. Therefore, compared to *Ficus pandurata Hance* indoor-cultivated, those grown by windows demonstrate superior carbon fixation and oxygen-releasing abilities.

#### 3.3. Leaf net photosynthetic rate

As depicted in **Figure 3**, with increasing temperatures, the  $P_n$  of *Ficus pandurata Hance* across varying CO<sub>2</sub> concentrations exhibit a pattern of initial increase followed by a decline; however, the magnitude of this change varies. At normal CO<sub>2</sub> concentration (400 µmol·mol<sup>-1</sup>), these plants demonstrate similar adaptability to different temperatures, with only slight increases in  $P_n$  as temperature rises, resulting in minimal differences in  $P_n$  among various temperatures. In contrast, at elevated CO<sub>2</sub> concentrations (800 µmol·mol<sup>-1</sup> and 1200 µmol·mol<sup>-1</sup>), the maximum  $P_n$  occurs at 25°C, accompanied by substantial variability between temperatures (a change rate of 249.88% at 800 µmol·mol<sup>-1</sup> and 70.47% at 1200 µmol·mol<sup>-1</sup>). Under moderate temperature ranges (25–30 °C) and high CO<sub>2</sub> concentrations (800 µmol·mol<sup>-1</sup> and 1200 µmol·mol<sup>-1</sup>) here than that observed under normal CO<sub>2</sub> conditions.



**Figure 3.** Net photosynthetic rate of *Ficus pandurata Hance* window-cultivated under different temperature and CO<sub>2</sub> conditions.

Therefore, under conditions of adequate indoor air circulation (atmospheric CO<sub>2</sub> concentration of 400  $\mu$ mol·mol<sup>-1</sup>), *Ficus pandurata Hance* exhibits minimal sensitivity to temperature fluctuations and maintains a relatively stable *P*<sub>n</sub>. Conversely, when indoor air circulation is restricted and CO<sub>2</sub> concentrations are elevated (800  $\mu$ mol·mol<sup>-1</sup> and 1200  $\mu$ mol·mol<sup>-1</sup>), *Ficus pandurata Hance* can sustain a comparatively high *P*<sub>n</sub> provided that the temperature remains within an optimal range; specifically, the *P*<sub>n</sub> can reach 2.35 times and 1.69 times those observed under normal conditions at CO<sub>2</sub> concentrations of 800  $\mu$ mol·mol<sup>-1</sup> and 1200  $\mu$ mol·mol<sup>-1</sup>, respectively. Additionally, it can achieve near-normal *P*<sub>n</sub> even at high/low temperatures. This indicates that *Ficus pandurata Hance* can effectively perform its ecological role of carbon sequestration and oxygen release regardless of the air circulation within the building. Especially under suitable temperature conditions (indoor conditions with air conditioning for temperature control in summer and

heating for temperature control in winter), the air-stagnant indoor environment (high  $CO_2$  concentration conditions) can promote *Ficus pandurata Hance* to carry out photosynthesis and release oxygen. Therefore, *Ficus pandurata Hance* is suitable for indoor cultivation in all seasons and temperatures (such as indoor conditions with air conditioning for temperature control in summer and heating for temperature control in winter) and can play a better ecological role of carbon sequestration and oxygen release.

## 3.4. Leaf transpiration rate

As can be seen from **Figure 4**, with the increase of temperature, the  $T_r$  at different CO<sub>2</sub> concentrations increases first and then decreases. At 15 °C, the  $T_r$  of the plant was lowest under different CO<sub>2</sub> concentrations, and it changed significantly with changes in temperature. When the CO<sub>2</sub> concentration was 800 µmol·mol<sup>-1</sup>, the  $T_r$  was the highest under all temperature conditions. At room temperature (20–25 °C), the  $T_r$  of *Ficus pandurata Hance* under high CO<sub>2</sub> concentration (800 µmol·mol<sup>-1</sup>, 1200 µmol·mol<sup>-1</sup>) was higher than that under normal CO<sub>2</sub> concentration (400 µmol·mol<sup>-1</sup>). At high temperature (30–35 °C), the  $T_r$  of *Ficus pandurata Hance* was the lowest when CO<sub>2</sub> concentration was 1200µmol·mol<sup>-1</sup>.



**Figure 4.** Transpiration rate of *Ficus pandurata Hance* window-cultivated under different temperature and CO<sub>2</sub> conditions.

At a CO<sub>2</sub> concentration of 800  $\mu$ mol·mol<sup>-1</sup>, the *T*<sub>r</sub> of *Ficus pandurata Hance* was highest at the intermediate temperature (25–30 °C), indicating that when the temperature is suitable, *Ficus pandurata Hance* has the strongest evapotranspiration and humidification ability in indoor environments with slightly poor air circulation and moderate population density (refer to **Table 1**, where one person is in a closed indoor environment with a CO<sub>2</sub> concentration of 800  $\mu$ mol·mol<sup>-1</sup>). In indoor environments with poor air circulation and high population density (CO<sub>2</sub> concentration > 800  $\mu$ mol·mol<sup>-1</sup>), *Ficus pandurata Hance*'s evapotranspiration and humidification ability is still better than that in well-ventilated conditions when it is at a suitable temperature (20–25 °C). This indicates that under suitable temperature conditions (indoor conditions with air conditioning for temperature control in summer and heating for temperature control in winter), *Ficus pandurata Hance* can play a better humidification ecological role in indoor environments with slightly poor air circulation. Under high temperature conditions, proper ventilation (CO<sub>2</sub> concentration < 1200  $\mu$ mol·mol<sup>-1</sup>) can enable *Ficus pandurata Hance* to play a better humidification ecological role.

#### 4. Discussion and conclusion

The relatively closed indoor environment is often accompanied by low light level  $(1-50 \ \mu mol \cdot m^{-2} \cdot s^{-1})$ , high CO<sub>2</sub> concentration  $(1000-5000 \ \mu mol \cdot mol^{-1})$ , easy to adjust the temperature (can be properly controlled by heating or air conditioning) and other characteristics[5,6,17]. As one of the important media to improve indoor air quality, indoor green plants (also an important low-carbon method), due to the above indoor environment characteristics are quite different from the original environment of plants, so the changes in indoor light conditions, temperature, CO<sub>2</sub> concentration, etc., will greatly affect the immediate changes in photosynthesis and transpiration, which will affect the plant's carbon fixation and oxygen release, as well as its transpiration and humidification ecological benefits[18,19].

There is often a problem with low light levels in the room. Studies have shown that increasing the indoor light intensity to  $300 \,\mu mol \cdot m^{-2} \cdot s^{-1}$  can significantly increase the CO<sub>2</sub> assimilation ability of indoor plants, and the increase of assimilation ability also increases the relative humidity of air[6]. In addition, 'CO2 fertilization effect' shows that many plants have higher carbon assimilation ability under high  $CO_2$ concentration conditions [20]. In this study, by measuring the light response curves and CO<sub>2</sub> response curves of Ficus pandurata Hance at different culture locations, we can see that the light saturation point for Ficus pandurata Hance window-cultivated is 718 µmol·m<sup>-2</sup>·s<sup>-1</sup>, accompanied by a lower light compensation point. This suggests that plants grown near windows exhibit a broader adaptability and enhanced capacity for light energy utilization compared to those cultivated in low-light environments. Furthermore, Ficus pandurata Hance grown near windows displays both a lower CO<sub>2</sub> compensation point and a higher saturation point, further underscoring its superior ability to utilize CO<sub>2</sub> resources in indoor settings. Consequently, *Ficus pandurata Hance* is particularly well-suited for cultivation near windows, where adequate light is available while minimizing the risk of excessive light intensity leading to photoinhibition [21]. In conclusion, cultivating Ficus pandurata Hance in window areas can significantly enhance its ecological functions related to carbon fixation and oxygen release.

At the same time, by investigating the physiological responses of plants across a range of temperatures, light intensities, and CO<sub>2</sub> concentrations, it is possible to accurately determine the optimal growth conditions for indoor plants to maximize their ecological contributions. This study demonstrates that under optimal indoor cultivation conditions, the  $P_n$  of *Ficus pandurata Hance* can approximate that of broadleaf tree species growing outdoors (4.357 µmol·m<sup>-2</sup>·s<sup>-1</sup>) [22]. The closed architectural design of buildings frequently results in insufficient ventilation, combined with a high density of indoor occupants, which can contribute to elevated

CO<sub>2</sub> concentrations within the indoor environment (**Table 1**) [5,17]. *Ficus pandurata Hance* when cultivated indoors can demonstrate notable capabilities for carbon sequestration and oxygen release, as well as transpiration and humidification ecological functions under conditions of high CO<sub>2</sub> concentration (**Figures 2–4**). For example, at optimal temperatures (25–30 °C) and a CO<sub>2</sub> concentration of 800  $\mu$ mol·mol<sup>-1</sup>, the ecological benefits associated with carbon sequestration and oxygen release-alongside transpiration and humidification-are particularly pronounced; similarly, at room temperature (20–25 °C) with a CO<sub>2</sub> concentration of 1200  $\mu$ mol·mol<sup>-1</sup>. Moreover, under high-temperature conditions with adequate ventilation (CO<sub>2</sub> concentration < 1200  $\mu$ mol·mol<sup>-1</sup>), *Ficus pandurata Hance* continues to make significant contributions to humidification ecological benefits. This result also fully reflects the 'CO<sub>2</sub> fertilization effect' [20].

In conclusion, to optimize the ecological benefits of *Ficus pandurata Hance* when cultivated indoors and enhance its positive impact on the working and living environment of indoor occupants, it is recommended that the plant be positioned near a window. Furthermore, under conditions of effective temperature regulation (indoor conditions with air conditioning for temperature control in summer and heating for temperature control in winter) [23], both stagnant air and moderate airflow can enable *Ficus pandurata Hance* to demonstrate enhanced carbon sequestration, oxygen release, transpiration, and humidification capabilities compared to scenarios involving complete ventilation.

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