

광환경 조절이 인삼의 생육과 진세노사이드 함량에 미치는 영향

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Effect of Controlled Light Environment on the Growth and Ginsenoside Content of *Panax ginseng* C. A. Meyer

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ABSTRACT

Background: The photosynthetic efficiency cool-season, semi-shade ginseng is normal at low morning temperatures, but drops at high afternoon temperatures. Therefore, optimal plant performance would be ensured if it were possible to control daily light transmission rates (LTR).

Methods and Results: Plants were grown in a controlled light environment that replicated 11 AM conditions and comparatively analyzed against plant grown under normal conditions. Growth in the controlled light environment resulted in a 2.81 fold increase in photosynthetic efficiency with no change in chlorophyll content, although LTR were high due to low morning temperatures. Increased aerial plant growth was observed in the ginseng plants adapted to the controlled light environment, which in turn influenced root weight. An 81% increase in fresh root weight (33.3 g per plant on average) was observed in 4-year-old ginseng plants grown in controlled light environment compared to the plants grown following conventional practices (18.4 g per plant on average). With regard to the inorganic composition of leaves of 4-year-old ginseng plants grown in controlled light environment, an increased in Fe content was observed, while Mn and Zn content decreased, and total ginsenoside content of roots increased 2.37 fold.

Conclusions: Growth of ginseng under a favorable light environment, such as the condition which exist naturally at 11 AM and are suitable for the plant's photosynthetic activity creates the possibility of large scale production, excellent-quality ginseng.

Key Words: *Panax ginseng* C. A. Meyer, Controlled Light Environment, Shading Facility, Light Transmission Rate, Photosynthetic Efficiency

INTRODUCTION

Ginseng (*Panax ginseng* C. A. Meyer) does not have stomas on the leaf surface; adaxial, but they are scattered across the backside; abaxial, and their number is very small. In addition, ginseng is one of the semi-shade plants whose chlorophyll development and Rubisco activation are low and whose chlorophyll-protein complex is created slowly, so the CO₂ gas interchange quantity is very small (Yang *et al.*, 1987; Xu and Zhou, 2008).

For this reason, for the ginseng leaves to be less

stressed out, and for the optimal photosynthetic speed to be maintained, the light environment condition of 8-21 klux at 18-21°C should be maintained (Lee, 1988). Since Korea has a big temperature difference according to season and time zone, and the light-blocking net is fixed in the case of a conventional shading facility, however, it is practically difficult to attract the proper amount of light. Most farmers ordinarily try to install only four-layer shading net and customarily add a two-layer shading net during summer with high temperature to lower the amount of light and temperature (RDA, 2009). As the temperature

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increases, the light saturation point falls, going down below 8 klux at over 35°C, with photo-oxidation proceeding quickly. Thus, the photosynthetic vitality of leaves drops (Lee *et al.*, 2010).

If we look at the photosynthetic speed of ginseng by time zone during the day as measured in Korea, it is high until 11 AM when the temperature is around 23°C with the sun, becoming very low as the temperature went up over 30°C after that (Lee *et al.*, 2012; Hyun *et al.*, 1993). If high radiation intensity continues in the morning when the light saturation point becomes low as the temperature becomes high soon, the unnecessary light energy stimulates the leaves; consequently, single oxygen ($^1\text{O}_2$), superoxide (O_2^-), hydrogen peroxide (H_2O_2), and hydroxyl radical ($-\text{OH}$) occur excessively, causing the lipid components of the biological membrane to undergo peroxidation, which destroys the enzyme protein and chlorophyll rapidly (Yang *et al.*, 1990; Ahn *et al.*, 1994).

Damaged leaves cannot recover, so they fall early in the middle of August; in worse cases, root hypertrophy remarkably declines, and defoliation occurs (Mo *et al.*, 2015). On the other hand, if we can control LTR every day, we will be able to draw an ideal result in terms of ginseng growth through the inflow of a lot of light at the beginning of 11 AM as the inflection point of photosynthetic efficiency and by drawing in less light later.

Therefore, this study was conducted to install sun shading that could control the light environment based on 11 AM and examine ginseng's growth and photosynthetic characteristics. It compared/analyzed the amount of ginsenosides as the representative second metabolite of ginseng to provide basic data for the study on improving ginseng's physiology and sun shading facility.

MATERIALS AND METHODS

1. Operation of light environment control shade facility and growth condition

The experiments were performed in the experimental field at the department of Herbal Crop Research located in Eumseong, Chungbuk from March 28, 2014 to October 15, 2015. Pipes with diameter of 48 mm were inserted into the ground at a depth of 0.6 m, and height measurement was adjusted to 2.5 m; doublers with width of 30 cm were installed in blind form to control LTR. The ceiling was

opened in the morning and closed after that at 11 AM every day. Of course, the ceiling was closed in the morning when it rained to prevent the occurrence of spread of disease such as alternaria blight by *Alternaria panax*, anthracnose by *Colletotrichum gloeosporioides*, and phytophthora blight by *Phytophthora cactorum* (Park *et al.*, 2012).

Four- and two-black layer polyethylene shading nets were put on the timber A-type slant shade facility as a control plot. The test areas were 121 m² for each treatment plot, and sudan grasses were sowed at the beginning of May 2013, harvested in the middle of July, and plowed up to 12 times at a 10-day interval with organic compost of 1,500 kg/10 a to manage the preparation field. 2-year-old violet-stemmed ginseng (2.5 ± 0.2 g) was planted at 72 plants/3.3 m² in the east/west furrow direction, and disease/pest prevention, weed removal, and other management were done based on the ginseng GAP standard cultivation guideline (RDA, 2009).

The external and internal radiation intensities of facilities were measured for 2 minutes and 3 times by LI-250A quantum sensor (LI-COR, Lincoln, NE, USA) at the height of the leaves to establish LTR following the light environment control; a thermometer (Thermo Recorder TR-72Ui, T&D Corp., Tokyo, Japan) with data save function was hung around the leaves during the high-temperature period from July to August 2014 to measure the temperature.

2. Growth research and photosynthetic characteristics

The growth of the aboveground part was surveyed on July 21, 2014 and July 20, 2015; the underground part growth was researched for 30 weeks at random when all leaves fell by years. The leaf area was measured by the WindDIAS image analysis system (Delta-T Devices Ltd., Cambridge, England), length of stem and root, by general ruler, diameter of stem and root, by Digimatic calipers (Caliper 500-182, Mitutoyo, Kawasaki, Japan), and fresh root weight, by electronic scale (RE260, CAS, Seoul, Korea).

Data on chlorophyll content was printed out by putting the chlorophyll meter (SPAD-502, Konica Minolta, Tokyo, Japan) at the center of the leaf three times before harvesting the ginseng (Markwell *et al.*, 1995).

The net CO_2 assimilation rate (A_n) and stomatal

conductance (g_s) were measured by putting the LI6400 portable photosynthesis system (LI-COR, Lincoln, NE, USA) to which a 6 cm² leaf chamber was attached at the center of the leaf vertically at 9:00 AM and 3:00 PM on July 22, 2014. The internal conditions of the leaf chamber were set as follows: air influx rate of 500 $\mu\text{mol/s}$ and CO₂ concentration of 400 μmol . The temperature was opened in order to interlock with the outside (Lee *et al.*, 2012).

3. Analysis of leaf inorganic component

3-year-old ginseng leaves were harvested on July 20, 2014, cleansed with distilled water, and freeze-dried for 72 hours at -80°C . They were decomposed to 0.5 g pulverized dry samples by using nitric acid, perchloric acid, and hydrochloric acid in that order and diluted and filtered to a fixed quantity. P₂O₅, K₂O, MgO, CaO, Na₂O, Cu, Mn, Fe, and Zn were quantified (NIAST, 2000) by the ICP analyzer (GBC Scientific Equipment, Braeside, Australia).

4. Analysis of root ginsenoside

10 kinds of ginsenosides were analyzed by finally harvesting 3-year-old and 4-year-old ginseng roots. The 10 kinds of ginsenoside standard samples were Rb₁, Rb₂, Rb₃, Rc, Rd, Re, Rf, Rg₁, Rg₂, and Rh₁ (ChromaDex, Irvine, CA, USA), the solvent for analysis was GR-grade. For the ginsenoside analysis, 0.2 g ginseng powder sample and 1 ml of 70% MeOH were put into a 2 ml tube to mix them well. The mixture was then put in ultrasonic bath for extraction by ultrasonic wave at 50°C for 30 minutes. The supernatant obtained by centrifugal separation (4°C , 15 min) at 13,000 rpm was put into a 2 ml tube, and 1 ml was refined by the Sep-Pak C₁₈ cartridge. The refined extract was filtered by a 0.45 μm membrane filter for use as assay sample (Kim *et al.*, 2008), and the ginsenoside content was measured using the Agilent 1100 series HPLC system (Agilent Technologies, Santa Clara, CA, USA). 10 μl of the extract was injected using the Halo RP-amide column (4.6×150 mm, 2.7 μm , Wilmington, DE, USA) and was analyzed at mobile flow velocity of 0.5 ml/min, column temperature of 50°C , and UV detector wavelength of 203 nm.

5. Statistical treatment

The results obtained from this experiment on the control

group and treatment plot were analyzed via student's *t*-test using the SAS software package (SAS v9.2, SAS Institute Inc., Cary, NC, USA), and the difference was proven to be statistically significant when the *p*-value was under 0.05.

RESULTS AND DISCUSSION

1. Comparison of cultivation environment and photosynthetic characteristics

The LTR of the conventional shading facility in 2014 was 7.0%, which was in the optimal LTR 5-20% range for ginseng growth; it coincided with the light environment in the inclined facility installed by most farmers and researchers (Cheon *et al.*, 1991a, b).

In controlling the light environment, LTR was 17.9% before 11 AM; the light inflow rate was higher than the conventional shading facility, but it did not exceed 20% and became 3.4% when the light was almost blocked after 11 AM (Table 1). The average temperature during the day at the time (6 AM-8 PM) the sun was up from July 1 to August 31 was 1.2°C lower than conventional practice (27.1°C) when the light environment (25.9°C) was controlled. This was because the hot air was not stagnant but escaped between the sun visors (Fig. 1). The lower temperature than the conventional shading facility and LTR control following the temperature rise had a positive effect on the morning photosynthetic efficiency of 3-year-old ginseng (Fig. 2). In the morning, A_n was $5.9 \mu\text{mol/m}^2/\text{s}$, and g_s was $0.08 \text{ mol/m}^2/\text{s}$; these were about 2.8 or 2 times higher than conventional practice, as the radiation intensity did not reach the light saturation point at a relatively low temperature than the afternoon.

Being put in a light environment wherein photosynthetic action could be carried out smoothly seemed to be the

Table 1. Light transmission rates of control and blind-type shading facilities controlled at 11:00 AM.

| Treatment | Open field | Control | Blind-type | |
|------------------------------|------------|---------------|-------------------------|-----------------------|
| | | | Opening before 11:00 AM | Closed after 11:00 AM |
| Light transmission rates (%) | 100 | 7.0 ± 2.1 | 17.9 ± 6.2 | 3.4 ± 1.9 |

Mean values \pm SD from triplicate separate experiments are shown.

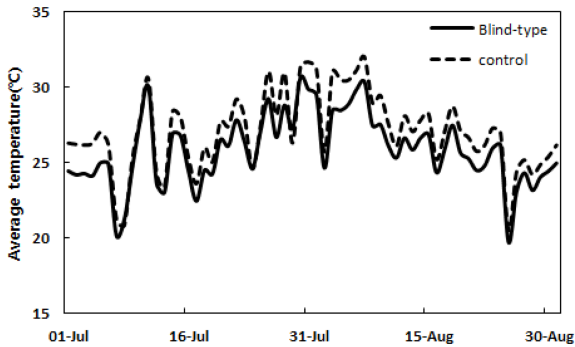


Fig. 1. Average temperature at 6 AM - 8 PM of control and blind-type shading facilities from July 1, 2014 to August 31, 2014.

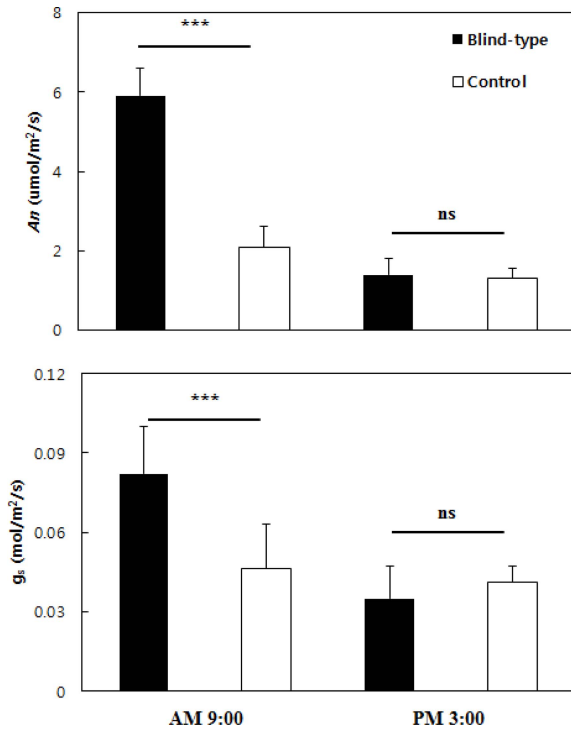


Fig. 2. Net CO₂ assimilation rate *An* and stomatal conductance *g_s* of 3-year-old *Panax ginseng* grown in control and blind-type shading facilities at 9:00 AM and 3:00 PM on July 22, 2014. The vertical error bars represent the standard errors (n = 5). **p* < 0.05, ***p* < 0.01, and ****p* < 0.001 compared to the control. NS; Not significant.

major reason unlike the control plot with limited *An* (Jochum *et al.*, 2007). In addition, ginseng leaves exposed to strong light consistently or intermittently had high light adaptability, so *An* was formed more smoothly (Jang *et al.*, 2015; Mo *et al.*, 2015). This was because the photosynthetic efficiency in the morning was considerably different. Although the conventional facility's LTR was

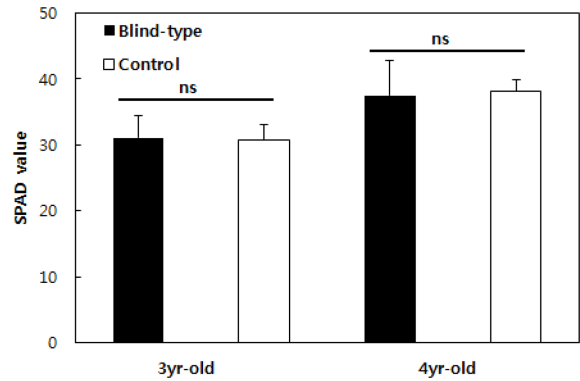


Fig. 3. SPAD value of 3- and 4-year-old *Panax ginseng* grown in control and blind-type shading facilities on July 21, 2014 and July 20, 2015. The vertical error bars represent the standard errors (n = 30). **p* < 0.05, ***p* < 0.01, and ****p* < 0.001 compared to the control. NS; Not significant.

more than 2 times higher in the afternoon, no statistical difference was found not only in *An* but also in *g_s* due to the effect of the temperature rise.

The chlorophyll content of ginseng was higher in the 4-year-old one than in the 3-year-old one. This is because the formation of the leaves is ecologically stable, and mesophyll cells become thick in the 4-6-year-old aged roots (Fig. 3). In the ginseng leaves, the reduction of chlorophyll content by oxidized stress is distinct if light quantity exceeding the light saturation point enters (Yang *et al.*, 1994). Although there is a difference according to the environment, if LTR exceeds 20%, this phenomenon becomes more serious because, if the leaves receive more light than the ones to be used, the response center of the light system II becomes inactive due to the photo-inhibition phenomenon (Ort and Baker, 1988).

As seen in the data, however, although LTR was high in the morning when the light environment was controlled, there was no difference from the control in terms of chlorophyll content. Dynamic photo-inhibition is observed at moderate light. Although the quantum efficiency decreases, the photosynthetic speed does not change. The reason for the dynamic photo-inhibition is that the absorbed light energy is transferred to the heat dissipation, and quantum efficiency declines (Tyystjarvi *et al.*, 1991). Since this reduction is temporary, if the photon flux drops below the saturation level, it can be restored to the original high value.

2. Growth of ginseng

There is a difference depending on the environment and kind. Proctor *et al.* (2010), who tested on *P. quinquefolius*, reported that, when LTR increased to 30%, the aboveground part and underground part growth increased. Kim *et al.* (1982) claimed that the ginseng that grew in the front row of the shading facility where relatively more light shone grew better than the ones that grew in the rear row. According to our test, where the light environment was controlled, the light influx in the

Table 2. Growth of aerial part of 3- and 4-year-old *Panax ginseng* grown in control and blind-type shading facilities on July 21, 2014 and July 20, 2015.

| Year | Treatment | Shoot length (cm/plant) | Shoot diameter (mm/plant) | Leaf area (cm ² /plant) |
|----------------------|----------------|-------------------------|---------------------------|------------------------------------|
| 2014 (3-year-old) | Control | 12.4 ± 2.0 | 3.4 ± 0.4 | 204.1 ± 47.8 |
| | Blind-type | 14.8 ± 2.4 | 3.9 ± 0.4 | 274.6 ± 59.7 |
| | <i>t</i> -test | ** | ** | *** |
| 2015 (4-year-old) | Control | 41.2 ± 4.7 | 6.1 ± 0.6 | 735.7 ± 73.9 |
| | Blind-type | 46.9 ± 5.2 | 7.5 ± 0.8 | 917.5 ± 60.1 |
| | <i>t</i> -test | ** | *** | *** |

Values are presented as means ± SD (n = 30). **p* < 0.05, ***p* < 0.01, and ****p* < 0.001 compared to the control.

Table 3. Growth of the subterranean part of 3- and 4-year-old *Panax ginseng* grown in control and blind-type shading facilities on the final harvest day.

| Year | Treatment | Root length (cm/plant) | Root diameter (mm/plant) | Fresh root weight (g/plant) |
|----------------------|----------------|------------------------|--------------------------|-----------------------------|
| 2014 (3-year-old) | Control | 22.4 ± 4.5 | 11.2 ± 2.1 | 8.6 ± 2.7 |
| | Blind-type | 24.5 ± 5.4 | 14.3 ± 1.7 | 15.4 ± 4.6 |
| | <i>t</i> -test | * | *** | *** |
| 2015 (4-year-old) | Control | 22.3 ± 4.1 | 15.3 ± 2.4 | 18.4 ± 4.2 |
| | Blind-type | 26.7 ± 2.4 | 19.9 ± 2.6 | 33.3 ± 7.2 |
| | <i>t</i> -test | ** | *** | *** |

Values are presented as means ± SD (n = 30). **p* < 0.05, ***p* < 0.01, and ****p* < 0.001 compared to the control.

Table 4. Inorganic component contents of 3-year-old *Panax ginseng* leaf grown in control and blind-type shading facilities on July 21, 2014.

| Treatment | P ₂ O ₅ | K ₂ O | MgO | CaO | Na ₂ O | Cu | Mn | Fe | Zn |
|----------------|-------------------------------|------------------|-----------|-----------|-------------------|-----------|--------------|--------------|--------------|
| | (%) | | | | | (mg/kg) | | | |
| Control | 0.3 ± 0.1 | 1.8 ± 0.3 | 0.7 ± 0.2 | 1.3 ± 0.3 | 0.01 ± 0.01 | 8.8 ± 1.2 | 768.8 ± 85.2 | 244.7 ± 48.1 | 122.6 ± 23.6 |
| Blind-type | 0.4 ± 0.1 | 1.7 ± 0.2 | 0.6 ± 0.2 | 1.0 ± 0.2 | 0.02 ± 0.01 | 8.5 ± 0.7 | 288.7 ± 37.6 | 410.5 ± 58.2 | 64.3 ± 18.7 |
| <i>t</i> -test | ns | ns | ns | ns | ns | ns | *** | * | *** |

Values are presented as mean ± SD (n = 3). **p* < 0.05, ***p* < 0.01, and ****p* < 0.001 compared to the control. NS; Not significant.

morning could increase the aboveground part growth of ginseng compared to the control (Table 2).

In particular, the area of the leaves as the light acceptor broadened by 34% on 3-year-old ginseng in 2014. The increase of leaf area and *An* in the morning led to the enhancement of the root, which was the storage of metabolite, affecting the growth of the 4-year-old ginseng the following year (2015) because the growth of ginseng that year is greatly influenced by the prolificacy of the previous-year ginseng. Jang *et al.* (2015) reported that the leaf area became broad when LTR increased up to 17% in the case of 2-year-old ginseng; this is because, when excessive light shone after the leaves became broad up to a certain level, the leaves diminished, recognizing it as stress.

In the end, the light environment regulation for 2 years based on 11 AM increased the root weight of the 4-year-old ginseng by about 81% compared to the control (Table 3). Meanwhile, as blind-type shading facility was continuously controlled after the end of the sweltered July, leaf burn of ginseng rapidly happened. It assumed that dry damage was also partially generated because of lowered soil and air humidity.

3. Analysis of leaf inorganic component and root ginsenoside

As shown in Table 4, P₂O₅, K₂O, MgO, CaO, Na₂O, and Cu had no statistically significant difference. Note, however, that there was a big difference in Mn, Fe, and Zn. Fe is a component of hemoprotein and Fe-S protein existing in the redox system, and about 80% exists in chlorophyll.

Since Fe is a component of aconitase, an enzyme that changes the citrate into isocitrate during the Krebs Cycle, if Fe is deficient, organic acid is accumulated, carbohydrate metabolism is affected, and redistribution does not occur in the plant; thus, the chlorophyll content of

Table 5. Ginsenoside contents of roots of 3- and 4-year-old *Panax ginseng* grown in control and blind-type shading facilities on the final harvest day.

| Year | Treatment | Ginsenoside contents of roots (mg/g, root dry weight) | | | | | | | | | | |
|----------------------|------------|---|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|--------------|
| | | Rb1 | Rb2 | Rb3 | Rc | Rd | Re | Rf | Rg1 | Rg2 | Rh1 | Total |
| 2014 (3-year-old) | Control | 1.17 ± 0.09 | 0.54 ± 0.05 | 0.11 ± 0.03 | 0.67 ± 0.07 | 0.25 ± 0.09 | 1.69 ± 0.17 | 0.49 ± 0.05 | 3.35 ± 0.34 | 0.21 ± 0.13 | 0.02 ± 0.01 | 8.50 ± 1.12 |
| | Blind-type | 2.60 ± 0.21 | 0.83 ± 0.10 | 0.17 ± 0.05 | 1.16 ± 0.13 | 0.41 ± 0.12 | 3.27 ± 0.27 | 0.97 ± 0.07 | 8.38 ± 0.62 | 0.41 ± 0.15 | 0.02 ± 0.01 | 18.22 ± 2.49 |
| | t-test | *** | *** | ** | *** | ** | *** | *** | *** | ** | ns | *** |
| 2015 (4-year-old) | Control | 2.57 ± 0.38 | 0.83 ± 0.17 | 0.31 ± 0.03 | 2.65 ± 0.18 | 1.99 ± 0.19 | 2.41 ± 0.58 | 0.99 ± 0.26 | 2.22 ± 0.38 | 0.30 ± 0.04 | 0.04 ± 0.01 | 14.31 ± 3.79 |
| | Blind-type | 5.97 ± 0.34 | 4.78 ± 0.25 | 0.91 ± 0.05 | 3.68 ± 0.25 | 1.49 ± 0.25 | 3.03 ± 0.43 | 1.86 ± 0.32 | 2.75 ± 0.46 | 0.32 ± 0.04 | 0.04 ± 0.01 | 24.84 ± 3.91 |
| | t-test | *** | *** | *** | *** | *** | ** | *** | ** | ns | ns | *** |

Values are presented as means ± SD (n = 30). * $p < 0.05$, ** $p < 0.01$, and *** $p < 0.001$ compared to the control. NS; Not significant.

the young leaves is low, and the leaves become chlorotic (Jeong and Gueriot, 2009). Although the chloroplast source of Fe is not well-known with regard to the synthesis of the photosynthetic cofactors, ferritin, an Fe-saving protein, will be relevant because the buffer action such as saving Fe is done to avoid excessive oxidative stress (Yruela, 2009).

Mn and Zn are indispensable mineral elements in photosynthesis in association with the superoxide dismutase; according to our experiment, it seems to have decreased somewhat as the benefit in return, i.e., Fe accumulated on the leaves (Armstrong, 2008).

Well-known as one of the medicinal properties, ginsenoside acts as the indicator that evaluates the quality of ginseng. Ginseng that is inevitably exposed to various kinds of stress for years or dozens of years in one place produces the saponin ingredients for the purpose of protecting itself (Choi *et al.*, 2005).

In this experiment, which controlled the light environment, the amount of ginsenoside consisting of 10 kinds Rb₁, Rb₂, Rb₃, Rc, Rd, Re, Rf, Rg₁, Rg₂, and Rh₁ increased regardless of life in years, although the biomass of ginseng increased. The ginsenoside content increased more in the 4-year-old ginseng than in the 3-year-old ginseng. When the light environment was controlled, the increase range (36%) was 13% higher than the control (23%). Although the study result was opposite to the one (Han *et al.*, 2013) stating that the ginsenoside content was low when the ginseng was heavy in conventional cultivation, light stress is considered one of the environment factors wielding a big effect on the ginsenoside content increase of ginseng (Jang *et al.*, 2015).

In sum, light environment control before and after 11 AM could increase the photosynthetic efficiency without a change in chlorophyll content, although LTR was high because of the low temperature in the morning. The aboveground part growth of the ginseng that adapted to the light environment increased to affect the root weight, and the total ginsenoside content of the 4-year-old ginseng multiplied by 2.37 times compared to the control due to proper light stress. Therefore, if light environment management according to the photosynthetic nature of the ginseng is carried out, it will be possible to produce large-capacity, excellent-quality ginseng.

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REFERENCES

- Ahn JS, Cho BG, Park H and Kim WK. (1994). Changes in chloroplast ultrastructure and thylakoid membrane proteins by high light in ginseng leaves. *Journal of Plant Biology*. 37:285-292.
- Armstrong FA. (2008). Why did nature choose manganese to make oxygen? *Philosophical Transactions of the Royal Society B: Biological Sciences*. 363:1263-1270.
- Cheon SK, Mok SK and Lee SS. (1991b). Effects of light intensity and quality on the growth and quality of Korean ginseng (*Panax ginseng* C. A. Meyer): II. Relationship between light intensity and planting density. *Korean Journal of Ginseng Science*. 15:31-35.
- Cheon SK, Mok SK, Lee SS and Shin DY. (1991a). Effects of light intensity and quality on the growth and quality of Korean

- ginseng(*Panax ginseng* C. A. Meyer): I. Effects of light intensity on the growth and yield of ginseng plants. Korean Journal of Ginseng Science. 15:21-30.
- Choi DW, Jung JD, Ha YI, Park HW, In DS, Chung HJ and Liu JR.** (2005). Analysis of transcripts in methyl jasmonate-treated ginseng hairy roots to identify genes involved in the biosynthesis of ginsenosides and other secondary metabolites. Plant Cell Report. 23:557-566.
- Han JS, Tak HS, Lee GS, Kim JS, Woo RJ and Choi JE.** (2013). Comparison of ginsenoside content and ratio of root tissue according to root age and diameter in *Panax ginseng* C. A. Meyer. Korean Journal of Medicinal Crop Science. 21:342-347.
- Hyun DY, Hwang JK, Choi SY and Jo JS.** (1993). Photosynthetic characteristics of *Panax ginseng* C. A. Meyer: I. Photosynthetic response to changes of light intensity and leaf temperature. Korean Journal of Ginseng Science. 17:240-245.
- Jang IB, Lee DY, Yu J, Park HW, Mo HS, Park KC, Hyun DY, Lee EH, Kim KH and Oh CS.** (2015). Photosynthesis rates, growth, and ginsenoside contents of 2-yr-old *Panax ginseng* grown at different light transmission rates in a greenhouse. Journal of Ginseng Research. 39:345-353.
- Jeong J and Guerinot ML.** (2009). Homing in on iron homeostasis in plants. Trends in Plant Science. 14:280-285.
- Jochum GM, Mudge KW and Thomas RB.** (2007). Elevated temperatures increase leaf senescence and root secondary metabolite concentrations in the understory herb *Panax quinquefolius* (Araliaceae). American Journal of Botany. 94:819-826.
- Kim GS, Hyun DY, Kim YO, Lee SE, Kim YC, Lee SW, Son WD, Lee MJ, Park CB, Park HK, Cha SW and Song KS.** (2008). Extraction and preprocessing methods for ginsenosides analysis of *Panax ginseng* C. A. Meyer. Korean Journal of Medicinal Crop Science. 16:446-454.
- Kim JM, Lee SS, Cheon SR and Cheon SK.** (1982). Relationship between environmental conditions and the growth of ginseng plant in field: I. Productive structures as affected by planting positions and ages. Korean Journal of Crop Science. 27:94-98.
- Lee CH.** (1988). Effect of light intensity and temperature on the photosynthesis and respiration of *Panax* spp. Korean Journal of Ginseng Science. 12:11-29.
- Lee JS, Lee DY, Lee JH, Ahn IO and In JG.** (2012). Photosynthetic characteristics of resistance and susceptible lines to high temperature injury in *Panax ginseng* Meyer. Journal of Ginseng Research. 36:461-468.
- Lee JS, Lee JH and Ahn IO.** (2010). Characteristics of resistant lines to high-temperature injury in ginseng(*Panax ginseng* C. A. Meyer). Journal of Ginseng Research. 34:274-281.
- Markwell J, Osterman JC and Mitchell JL.** (1995). Calibration of the Minolta SPAD-502 leaf chlorophyll meter. Photosynthesis Research. 46:467-472.
- Mo HS, Jang IB, Yu J, Park HW and Park KC.** (2015). Effects of enhanced light transmission rate during the early growth stage on plant growth, photosynthetic ability and disease incidence of above ground in *Panax ginseng*. Korean Journal of Medicinal Crop Science. 23:284-291.
- National Institute of Agricultural Sciences and Technology (NIAS).** (2000). Methods of soil chemical analysis. Rural Development Administration. Suwon, Korea. p.108-149.
- Ort DR and Baker NR.** (1988). Consideration of photosynthetic efficiency at low light as a major determinant of crop photosynthetic performance: Quantum yield, coupling factor, electrochemical potential, antenna chlorophyll complexes. Plant Physiology and Biochemistry. 26:555-565.
- Park YH, Lee SG, Ahn DJ, Kwon TR, Park SU, Lim HS and Bae H.** (2012). Diversity of fungal endophytes in various tissues of *Panax ginseng* Meyer cultivated in Korea. Journal of Ginseng Research. 36:211-217.
- Proctor JTA, Palmer JW and Follett JM.** (2010). Growth, dry matter partitioning and photosynthesis in North American ginseng seedlings. Journal of Ginseng Research. 34:175-182.
- Rural Development Administration(RDA).** (2009). Ginseng standard cultivation textbook. Rural Development Administration. Suwon, Korea. p.47-139.
- Tyystjarvi E, Koivuniemi A, Kettunen R and Aro EM.** (1991). Small light-harvesting antenna does not protect from photoinhibition. Plant Physiology. 97:477-483.
- Xu Z and Zhou G.** (2008). Responses of leaf stomatal density to water status and its relationship with photosynthesis in a grass. Journal of Experimental Botany. 59:3317-3325.
- Yang DC, Chae Q, Lee SJ, Kim YH and Kang YH.** (1990). Effects of light and photosynthetic electron transport system on the generation of singlet oxygen(¹O₂) in ginseng thylakoid membrane. Korean Journal of Ginseng Science. 14:57-62.
- Yang DC, Kim YH, Yang DC and Hong YN.** (1994). Effects of antioxidants on the photosynthesis and carbohydrates/saponin contents in *Panax ginseng* leaves. Korean Journal of Ginseng Science. 18:175-181.
- Yang DC, Yoo HS and Yoon JJ.** (1987). Investigation on the photooxidation of pigment in leaf-burning disease of *Panax ginseng*: II. Investigation and analysis of physiological reaction mechanism on the chlorophyll bleaching phenomenon. Korean Journal of Ginseng Science. 11:101-110.
- Yruela I.** (2009). Copper in plants: Acquisition, transport and interactions. Functional Plant Biology. 36:409-430.