

Effect of Chlorocholine Chloride on Chlorophyll, Photosynthesis, Soluble Sugar and Flavonoids of *Ginkgo biloba*

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Abstract

The flavonoids content determines the quality characteristics of *Ginkgo biloba* extract that could be increased by using of plant growth regulators. The objective of study was to investigate the effect of chlorocholine chloride (CCC), an anti-gibberellin growth retardant, on photosynthesis, chlorophyll, soluble sugar, total amino acids and phenylalanine contents, flavonoid accumulation, and flavonoids enzyme activity in *G. biloba* leaves. The ginkgo seedlings were grown in the greenhouse conditions with foliar applications of 0 (control), 0.5, 1.0 and 2.0 g l⁻¹ CCC. Results showed that 0.5, 1.0 and 2.0 g l⁻¹ CCC treatments significantly increased photosynthetic rates of leaves, the contents of chlorophyll, soluble sugar, total amino acids and phenylalanine in ginkgo leaves. Total polyphenols, flavonoids, anthocyanins content, phenylalanine ammonia-lyase (PAL), chalcone synthase (CHS) and chalcone isomerase (CHI) activities were all significantly increased by 1.0 and 2.0 g l⁻¹ CCC treatments. Foliar treatment with CCC therefore might be a useful means of improving pharmacological properties of *G. biloba* leaves.

Keywords: chlorocholine chloride, chlorophyll, flavonoids, *Ginkgo biloba*, photosynthesis, soluble sugar,

Introduction

The flavonoids, one kind of polyphenolic compounds, have a broad spectrum of action in plants. They protect against UV-B radiation and pathogen attack, act as attractants for pollinators, act as signal molecules for initiating plant-microbe symbiotic associations (Winkel-Shirley, 2002) and have been used as colorants, and for allelopathy (Hossain *et al.*, 2009). The studies on flavonoids in *G. biloba* had been more and more popular due to their antioxidative constituent that beneficial for human health. The flavonoids extracted from *G. biloba* leaves showed antioxidant activity and protection against the damages caused by free radicals, which play an important role in many diseases, such as cerebral hypoxia, cardiovascular disease and cancer (Ding *et al.*, 2009; Iwashita *et al.*, 2000; Kawaii *et al.*, 1999; Sachin *et al.*, 2000). Standardized preparations of *G. biloba*, Egb761, contain 24% flavonoids, which have several pharmaceutical properties for human health (van Beek, 2002). In *G. biloba*, flavonoids located primarily in the leaf are important determinants of its pharmacological properties and determine the quality characteristics of ginkgo tea or extract. Therefore, much interest is currently focusing on increasing the medicinal value of ginkgo leaves by increasing their flavonoid contents. *In vivo* experiments on ginkgo leaves indicated that externally supplied CCC,

ABA, ethephon, and metal ions could increase flavonoid contents (Cheng *et al.*, 2004; Wang *et al.*, 2007).

Chlorocholine chloride (CCC, 2-chloroethyltrimethylammonium chloride) is an antigibberellin growth retardant, with its mechanism based on the restraint of gibberellin biosynthesis in plant tissues (Wang and Xiao, 2009). It is well known that CCC induces changes in the growth rate of grasses and the morphogenesis of potato plants cultured *in vitro*. CCC recently has been widely used in potato production to increase yield and quality. Application of CCC causes an increase in dry matter accumulation in sweet potato tubers (Wang *et al.*, 2009). CCC might induce higher tuber production by causing an increase in chlorophyll content and photosynthesis (Sharma *et al.*, 1998a; Wang *et al.*, 2009). In addition, CCC treatment can enhance plant nutrient uptake from soil, improve water balance and protein synthesis in growing organs (Grossmann, 1990; Wang *et al.*, 2010). Furthermore, CCC was also found to promote anthocyanin synthesis in red cabbage and spiderwort leaves (Grebinskii and Khmil, 1980). Anthocyanin biosynthesis could be enhanced by feeding of precursors of both the moieties namely, phenylalanine, trans-cinnamic acid, glucose, sucrose and acetate (Guruprasad and Laloraya, 1980; Jain and Guruprasad, 1989). CCC also causes accumulation of carbohydrates in the tissues of wheat plants (Pikush, 1974).

Therefore, CCC application might be a promising practice for improving the secondary metabolite production in ginkgo leaves, such as flavonoids (Cheng *et al.*, 2009). A study by Cheng *et al.* (2011) showed that CCC treatment could promote flavonoid accumulation in ginkgo leaves. However, the underlying physiological mechanisms of CCC treatment enhancing the flavonoids content have not yet been elucidated. In this study, the effects of foliar application of different concentrations of CCC on photosynthesis, contents of chlorophyll, carbohydrate and activities of enzymes involved in flavonoid were investigated. Furthermore, total polyphenols, flavonoids, and anthocyanins accumulation in ginkgo leaves, which could be promoted by application of CCC were also determined. The objective of the present study was to evaluate the physiological action and to provide a theoretical basis of CCC application for high-yield flavonoid content in ginkgo leaves.

Materials and methods

Plant materials and CCC treatments

Three-years seedlings of *G. biloba* cv. 'Jiafoshou' were sown in the pots (25 cm diameter, 40 cm height, 1 plant/pot) filled with sandy soil (sand: humus soil 1:2). All seedlings were placed in a greenhouse with 12 h photoperiod (25°C; day/night, 500 $\mu\text{mol m}^{-2} \text{s}^{-1}$) at 60-70% RH, and irrigated daily with 1/2 Hoagland solution. After the fourth leaves were fully expanded, all seedlings were treated by foliar sprayed with aqueous solutions of CCC (containing 0.01% Tween 20, pH 5.8) at four concentrations of 0 (the control), 0.5, 1.0, and 2.0 g l^{-1} . Twenty mL of solution was applied once/month for three months. Six replications (9 plants/replication) were used per treatment and sample pooled for each determination. All the measurements were made 4 months after the initial application of CCC.

Measurement of photosynthetic rates, Chlorophyll a+b, soluble sugar, total amino acids and phenylalanine contents

The photosynthetic parameters were determined on the fifth leaf using a LiCor-6400 portable photosynthesis system (LiCor, U.S.A.) according to the instruction manual. Measurement periods were in the morning from 8:00 to 11:00 am when leaves of *G. biloba* were under a light intensity of 1000 $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ and humidity of 60% (Adamski *et al.*, 2011).

Chlorophyll a+b content was determined on fully expanded leaves. Leaf material was homogenized in cold (4°C) aqueous 80% (v/v) acetone and kept in the dark and centrifuged at 1000 $\times g$ for 5 min to remove the leaf debris. Chlorophyll a+b content was determined for each sample with a spectrophotometer (DU 730, Beckman Coulter Inc., USA) according to the methods of Graan and Ort (1984).

The soluble sugars of fresh leaf tissue were extracted with boiling ethanol/water and determined by the method of Sharma *et al.* (1998a), using an ultraviolet spectrophotometer (DU 730, Beckman Coulter Inc., USA). The concentration of soluble sugars was expressed as a percentage of fresh weight (w/w, FW).

The quantitative determination of total free amino acid was carried out by using spectrophotometer method (Rosen, 1957). Free amino acids were extracted from 0.1 g plant samples by homogenizing at 4°C in 5 mL of 80% (v/v) ethanol, diluted to a volume of 20 mL with 80% ethanol, kept in a water bath at 70°C for 30 min, and then centrifuged at 12000 $\times g$ for 10 min. The supernatants were filtered and then the filtrates were separated by two-dimensional paper chromatography (Whatman No. 1) according to Stepka (1957). The paper was loaded with 0.1 mL of the acetone extract. Phenol:water (100:39 v/v) was used as first and butanol: acetic acid: water (100:22:55 v/v) as second solvent. Papers were developed by spraying with 0.5% ninhydrin in acetone and heating the dry paper at 65-70°C for 20 min. Identification of the separated amino acids was done using the reference standard. The spots were eluted with 4 mL of 75% EtOH, respectively and then 0.2 mL eluent was made up to 2 mL deionized water and 1 mL ninhydrin solution was added. The absorbance at 570 nm was measured with a spectrophotometer (DU 730, Beckman Coulter Inc., USA) against a blank containing 2 mL deionized water and 1 mL ninhydrin solution. The concentration of each amino acid was determined from each amino acid standard curve. The phenylalanine concentration was estimated with equally handled phenylalanine standards. The total amino acid concentration was the amount of each amino acid concentration.

Extraction and analysis of total polyphenols, anthocyanins, and flavonoids

Five gram sample of fresh leaf tissue was frozen with liquid nitrogen, ground into a fine powder, homogenized in 50 mL of methanol containing 1% HCl. The solution was sonicated for 20 min with continual N_2 gas purging and centrifuged for 15 min at 12000 $\times g$ at 4°C. The supernatant was filtered through paper filter. The methanol extract was pooled and concentrated. The aqueous residue was extracted with petroleum ether (3 \times 30 mL), to remove pigments and lipids, and solvent residue was then removed under vacuum. The remaining solution was made up to 25 mL with methanol, and this solution was analyzed for total polyphenols (TP), total flavonoids (TF), and total anthocyanins (TA). Measurements of TP were carried out according to Arnous *et al.* (2001) by using the Folin-Ciocalteu reagent and the values were read at 675 nm. Gallic acid was used as the reference standard and results were expressed as mg gallic acid equivalents (GAE) 100 g^{-1} fresh weight (FW).

The flavonoid contents of *Ginkgo biloba* leaves grown in various concentrations of CCC were measured as de-

scribed by Xu *et al.* (2011). Briefly, one milliliter aliquot of extract, appropriately diluted, was mixed with 0.4 mL distilled water in a 1.5 mL micro centrifuge tube. Thirty microliters of 5% NaNO₂ solution were added and the solution was thoroughly mixed. Five minutes later, thirty microlitre of 10% AlCl₃ was added and the mixture was left to stand for 5 min. Then, two hundred microlitre 1 M Na₂CO₃ and 0.24 mL distilled water were added. After mixing the solution thoroughly, the absorbance was measured immediately at 510 nm using a microplate spectrophotometer. Total flavonoids content was calculated from a calibration curve using rutin as a standard and expressed as milligrams of rutin equivalents (RE) 100 g⁻¹FW.

TA was determined as described by Pirie and Mullins (1976) with minor modification. An aliquot of extract (5 mL) was combined with ethanolic HCl solution (0.25 M) to give a 1:10 dilution. After thorough mixing, the absorbance at 520 nm was read after 5 min. Total anthocyanin contents was determined as milligrams of cyanin (cyanidin 3-O-glucoside) equivalents 100 g⁻¹FW.

Determination of enzyme activity

Enzyme extracts were prepared as described by Xu *et al.* (2011). Protein was determined according to the Bradford method using BSA as a standard (Bradford, 1976). Phenylalanine ammonia-lyase (PAL) (E.C. 4.3.1.5) activity was measured as described by Solecka and Kacperska (2003). One unit of enzyme activity equaled the amount of PAL that produced 1 nmol of cinnamic acid in 1 h. Chalcone synthase (CHS) (E.C. 2.3.1.74) activity was measured according to Xu *et al.* (2007). Chalcone isomerase (CHI) (E.C. 5.5.1.6) activity was measured with the method of Li *et al.* (2006). CHS and CHI activities were both expressed as unit mg⁻¹ protein.

Statistical analysis

Data were statistically analyzed as a completely randomized design with each treatment replicated six times

by analysis of variance (ANOVA) using Data Processing System (DPS, Zhejiang University, Hangzhou, China). Appropriate standard errors of means (SE) were calculated. Duncan's multiple-range test was applied to compare measured parameters from plants that had experienced different treatments.

Results

Effect of CCC on leaf photosynthetic rates, Chlorophyll a+b, soluble sugar, total amino acids and phenylalanine contents

The changes in photosynthetic active of plants induced by spraying CCC were observed. Treatments with 0.5, 1.0 and 2.0 g L⁻¹ CCC resulted in an increase of 1.92, 29.12 and 48.85% in net photosynthesis rate (P_n) compared with the control, respectively. Treatments with 0.5 and 2.0 g L⁻¹ CCC resulted in an increase of 20 and 40% in stomatal conductance (G_s) compared with the control, respectively; while treatment with 1.0 g L⁻¹ CCC decreased the G_s by 10% relative to the control. Treatments with 0.5, 1.0 and 2.0 g L⁻¹ CCC resulted in an increase in intercellular CO₂ (C_i) of 11.52, 23.67 and 9.93% in ginkgo leaves, respectively. Treatments with 0.5, 1.0 and 2.0 g L⁻¹ CCC increased transpiration rate (T_r) by 8.33, 18.33, and 33.33%, respectively. The P_n , G_s , C_i and T_r of leaves were significantly different from each other (Tab. 1).

Spraying CCC not only improved the photosynthetic parameters of ginkgo leaves but also affect the contents of chlorophyll a+b, soluble sugar, total amino acids and phenylalanine in ginkgo leaves. The effect of 2.0 g L⁻¹ CCC was more pronounced on chlorophyll a+b, soluble sugar, and total amino acids than other treatments, whereas 1.0 g L⁻¹ CCC was the most effective on phenylalanine content (Tab. 2). With 0.5 and 1.0 g L⁻¹ CCC treatment, the chlorophyll a+b contents were less in ginkgo leaves than the control.

Tab. 1. Effects of CCC on the photosynthetic parameters in ginkgo leaves

CCC treatment	P_n (CO ₂ μmol m ⁻² s ⁻¹)	G_s (H ₂ O mol m ⁻² s ⁻¹)	C_i (CO ₂ μl l ⁻¹)	T_r (H ₂ O mmol m ⁻² s ⁻¹)
0 (Control)	10.44±0.61c	0.20±0.01c	276.75±12.70b	1.20±0.06c
0.5 g l ⁻¹	10.64±0.83c	0.24±0.02b	308.62±33.02b	1.30±0.10c
1.0 g l ⁻¹	13.48±1.41b	0.18±0.03c	342.25±19.36a	1.42±0.04b
2.0 g l ⁻¹	15.54±1.13a	0.28±0.02a	304.23±12.25b	1.60±0.04a

P_n , G_s , C_i and T_r represent net photosynthetic rate, stomatal conductance, intercellular CO₂ concentration, and transpiration rate, respectively. Values are mean ± standard error (SE) of six replications. Different letters within the same column indicate significant differences between treatments according to Duncan's multiple range test at $p < 0.05$ level

Tab. 2. Effect of CCC on the contents of Chlorophyll a+b, soluble sugar, total amino acids, phenylalanine in ginkgo leaves

CCC treatment	Chlorophyll a+b (mg g ⁻¹ FW)	Soluble sugar (mg g ⁻¹ FW)	Total amino acids (ug g ⁻¹ FW)	Phenylalanine (ug g ⁻¹ FW)
0 (Control)	1.47±0.06b	11.66±0.65c	8.81±0.31d	0.16±0.01d
0.5 g l ⁻¹	1.39±0.09b	12.46±0.36	10.95±0.52c	0.22±0.01c
1.0 g l ⁻¹	1.46±0.04b	13.36±0.23b	16.99±1.04b	0.36±0.03a
2.0 g l ⁻¹	1.79±0.02a	14.28±0.40a	21.47±0.89a	0.31±0.01b

Values are mean ± standard error (SE) of six replications. Different letters within the same column indicate significant differences between treatments according to Duncan's multiple range test at $p < 0.05$ level

Soluble sugar contents in all leaves, regardless of treatment, increased gradually (Tab. 2). The sugar contents were significantly ($p < 0.05$) higher in leaves treated with 2.0 g l⁻¹ CCC than in the control. Sugar contents in leaves treated with 0.5 and 1.0 g l⁻¹ CCC conditions were 6.86% and 14.58%, respectively, higher than control.

Enhancement of total amino acids and phenylalanine after CCC treatments (Tab. 2) was observed in leaves. The total amino acids were increased in leaves by 24.92, 92.85, and 143.7% at 0.5, 1.0, and 2.0 g l⁻¹ of CCC treatment compared with the control, respectively. Treatments with 0.5, 1.0 and 2.0 g l⁻¹ CCC resulted in an increase of 37.5, 125 and 93.75% in phenylalanine contents compared with the control, respectively.

Effect of CCC on PAL, CHS, and CHI activities

Changes in three enzymes activities are shown in Tab. 3. Under CCC treatment the activity of PAL in leaves showed an increase compared to control, with the 1.0 g l⁻¹ CCC treatment the activity of PAL was significantly higher (70.88% higher) than control, but treated with 0.5 and 2.0 g l⁻¹ CCC were not significant ($p > 0.05$). The CHS activity of the leaves treated with 0.5 and 2.0 g l⁻¹ CCC was significantly higher ($p < 0.05$) than control, while treated with 1.0 g l⁻¹ CCC was significantly lower (1.99% lower) than control. CHI activity of leaves increased gradually under the CCC treatment. CHS activity was significantly ($p < 0.05$) enhanced (127.16% higher) above the control level by CCC treatment at 2.0 g l⁻¹. The CHI activities of the leaves treated with 0.5 and 1.0 g l⁻¹ CCC were 67.65% and 99.81% higher, respectively, than control (Tab. 3).

Effect of CCC on total polyphenols, flavonoids and anthocyanins

The contents of TP, TF and TA showed an increase in ginkgo leaves under CCC treatment. A gradual increase in TP, TF and TA in CCC-treated leaves from treatment

with 0.5 and 1.0 g l⁻¹ CCC. The TP, TF and TA contents in CCC treated leaves at 2.0 g l⁻¹ was lower than CCC treated leaves at 1.0 g l⁻¹ CCC, but still higher than control. Furthermore, the TP, TF and TA contents of leaves at 1.0 g l⁻¹ and 2.0 g l⁻¹ CCC application were significantly higher ($p < 0.05$) than their corresponding control, The TP contents were significantly increased 24.46% and 20.7%, respectively, and TF contents were significantly increased 23.37% and 18.46% respectively, while TA contents were significantly increased 16.67% and 13.89%, respectively (Tab. 4).

Discussion

CCC has been reported to benefit photosynthesis by increasing chlorophyll content in potato (Wang *et al.*, 2009), onion (Miroshnichenko and Manankov, 1991), wheat (Stahli *et al.*, 1995) and sunflower (Kumari and Bharti, 1992). Our research found that photosynthetic rates in ginkgo leaves could be significantly enhanced by foliar application of CCC (Tab. 1). The enhanced photosynthetic capacity might have been ascribed to the increased leaf thickness, leaf photosynthetic pigment contents, and stomata activity induced by CCC treatments (Tezuka *et al.*, 1989; Wang *et al.*, 2009).

Our data also showed that the increased chlorophyll a+b contents in ginkgo leaves in the CCC-treated plants, coincided with an increase in the photosynthetic capacity as indicated by the higher P_n , G_s , C_i , and T_r in those plants (Tab. 1 and 2). The increase in chlorophyll a+b might be associated with the increases in anthocyanin and flavonoid contents, as a positive relationship was noted among chlorophyll, anthocyanin, and flavonoid contents (Xu *et al.*, 2011).

In the present experiment, CCC treatments enhanced leaf soluble sugar content, which is in agreement with earlier reports (Sharma *et al.*, 1998a; Wang and Xiao, 2009),

Tab. 3. Effects of CCC on the activities of PAL, CHS and CHI of ginkgo leaves

CCC treatment	PAL(U mg ⁻¹ protein)	CHS(U mg ⁻¹ protein)	CHI(U mg ⁻¹ protein)
0 (Control)	227.29±7.47d	16.61±1.98a	42.63±1.17d
0.5 g l ⁻¹	284.68±8.76c	16.88±0.99a	71.47±3.86c
1.0 g l ⁻¹	388.39±6.66a	16.28±2.30a	85.16±4.52b
2.0 g l ⁻¹	358.90±17.59b	16.88±0.60a	96.84±1.80a

Values are mean ± standard error (SE) of six replications. Different letters within the same column indicate significant differences between treatments according to Duncan's multiple range test at $p < 0.05$ level

Tab. 4. Effects of CCC on the contents of total polyphenols, flavonoids and anthocyanins in ginkgo leaves

CCC treatment	Total polyphenols (mg 100 g ⁻¹ FW)	Flavonoids (mg 100 g ⁻¹ FW)	Anthocyanins (mg 100 g ⁻¹ FW)
0 (Control)	220.99±12.92c	149.79±6.32b	0.36±0.02b
0.5 g l ⁻¹	243.13±6.77b	157.95±9.91b	0.37±0.02b
1.0 g l ⁻¹	275.05±17.35a	184.79±5.29a	0.42±0.02a
2.0 g l ⁻¹	266.74±5.27a	177.44±3.83a	0.41±0.02a

Values are mean ± standard error (SE) of six replications. Different letters within the same column indicate significant differences between treatments according to Duncan's multiple range test at $p < 0.05$ level

where addition of CCC increased the carbohydrate in the tissues in potato. However, our result is not in agreement with the reports of oat caryopses (Foley, 1987) and radish seedlings (Jain and Guruprasad, 1989), where the carbohydrate decreased after CCC treatment. This promotive effect of CCC on soluble sugar contents was likely resulted from the increase in photosynthetic activity of the CCC treated plants due to a higher chlorophyll content of the leaves (Sharma *et al.*, 1998b).

Significant enhancement in the contents of total amino acids and phenylalanine were observed in response to CCC in the treated ginkgo leaves (Tab. 2), and similar with the literature (Ibrahim and Nermin, 2001) that the protein and amino acid contents were higher in seedlings of CCC treated maize seeds than those of untreated ones. The synthesis of amino acids is coupled to uptake of nitrogen and carbon, and sugars produced by photosynthesis are used as sources of carbon and energy for the synthesis of amino acids (Miyashita and Miyazaki, 1992). Therefore, the increase in total amino acids might be associated with the increases in soluble sugar contents, as a positive relationship was noted among chlorophyll, anthocyanin, and flavonoid contents. That is, the increased net photosynthesis may have produced carbohydrate in excess to what is need for ginkgo growth because of the CCC application.

This study results showed phenylalanine content increased in the CCC-treated ginkgo leaves. A similar conclusion has been reached in buckwheat seedlings (Margna *et al.*, 1969), where CCC increased the level of phenylalanine. Jain and Guruprasad (1989) found that the CCC mediated promotion of anthocyanin synthesis with or without GA₃ supplied can be correlated with an increase in the total free amino acids, in particular phenylalanine. External application of phenylalanine along with CCC further enhanced the synthesis of anthocyanin. In the CCC-treated ginkgo leaves, total anthocyanins contents were also increased. In contrast to our results, phenylalanine did not promote CCC-induced anthocyanin synthesis in red cabbage and spiderwort tissues (Grebinskii and Khmil, 1980). This difference may be due to the presence of phenylalanine in saturating levels in cabbage and spiderwort tissues after application of CCC.

Polyphenols, flavonoids, and anthocyanins are all involved in the quality characteristics of ginkgo leaves and their processed products, like GBE761 (van Beek, 2002). The present study showed notable increase in total polyphenols, flavonoids, and anthocyanins contents following CCC-treatment. CCC treatments also significantly induced the activities of PAL, CHS, and CHI in a concentration-dependent manner in ginkgo leaves. Recent findings demonstrated that PAL, CHS and CHI were both key regulatory enzymes in flavonoid biosynthesis in *G. biloba* leaves (Cheng *et al.*, 2011; Xu *et al.*, 2007, 2008) and observed that CCC could induce the transcription of *chalcone isomerase* gene (*GbCHI*) in ginkgo leaves (Cheng *et al.*, 2011). The transcript levels of several genes encod-

ing the enzymes involved in flavonoid and anthocyanin biosyntheses, such as PAL, CHS, and CHI, are positively correlated with the accumulation of flavonoids and anthocyanins (Fischer *et al.*, 2007; Jaakola *et al.*, 2002).

Total polyphenol, flavonoid, and anthocyanin contents have a positive relationship with the activities of PAL, CHS and CHI in many plants (Ju *et al.*, 1995; Mato *et al.*, 2000; Obinata *et al.*, 2003). Our recently work also demonstrated that CHS and PAL were two key enzymes involved in flavonoid biosynthetic pathway in *G. biloba* (Cheng *et al.*, 2012). For these reasons, increase in the total polyphenol, flavonoid, and anthocyanin contents have been assumed to be due to enhanced PAL, CHS and CHI activities as a result of CCC treatment. In addition, one recent report suggests that lemon balm leaves were exposed to high levels of sucrose, and in response, the anthocyanins and flavonoids accumulated (Hossain *et al.*, 2009). Sugar-induced anthocyanin accumulation has been reported in other plant species (Nagira *et al.*, 2006; Solfanelli *et al.*, 2006). The accumulation of total polyphenol, flavonoid, and anthocyanin contents in CCC-treated ginkgo leaves might also has a few relationships with the extra soluble sugar content, which was enhanced by CCC treatment. Our present data indicated 1.0 g l⁻¹ CCC application has most promotive effects on total polyphenol, flavonoid, and anthocyanin contents and on PAL, CHS and CHI activities.

Conclusions

From present studies it may be concluded that CCC at concentrations of 1.0 g l⁻¹ and 2.0 g l⁻¹ greatly promotes chlorophyll, soluble sugar content, total amino acids and phenylalanine, which are benefit for the accumulation of total polyphenol, flavonoid and anthocyanin contents in ginkgo leaves. Total polyphenol, flavonoid, and anthocyanin contents and PAL, CHS and CHI activities are much more induced by 1.0 g l⁻¹ CCC in ginkgo leaves. It should be suggested that CCC was an appropriate choice for treating ginkgo plants with the objective of increasing the production of flavonoids.

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