Changes in Biomass, Mineral Composition, and Quality of Cardoon in Response to NO$_3^-$:Cl$^-$ Ratio and Nitrate Deprivation from the Nutrient Solution

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Leaf extracts of cultivated cardoon (Cynara cardunculus L. var. altilis DC) are an important source of phenols. Soilless culture represents an important and alternative tool to traditional agriculture, since it allows a precise control of plant nutrition and the maximization of yield and quality of the product. Reducing N supply, while keeping quantity as high as possible is desirable for environmental and health-related reasons, especially that N deficiency can lead to improved concentrations of secondary plant metabolites. Two greenhouse experiments were carried out in order to determine the effect of a decreasing NO$_3^-$:Cl$^-$ ratio (80:20, 60:40, 40:60, or 20:80) and nitrate deprivation (0, 5, 10, or 15 days before harvest) on biomass production, leaf chlorophyll content and fluorescence, mineral composition, and phytochemicals in leaves of cardoon ‘Bianco Avorio’ grown in a floating system. Total phenols, flavonoids and antioxidant capacity increased linearly with Cl$^-$ availability, especially when nitrate was replaced by 80% of chloride (20:80 NO$_3^-$:Cl$^-$ ratio), without having a detrimental effect on yield. Total nitrogen and nitrate concentration in leaves decreased linearly with increasing Cl$^-$ in the nutrient solution. Total phenols and antioxidant capacity recorded after 15 days of nitrate deprivation were higher by 43.1, 42.8, and 44.3% and by 70.5, 40.9, and 62.2%, at 59, 97 and 124 days after sowing, respectively compared to the control treatment. The decrease in leaf nitrate content recorded under N-deprivation occurred more rapidly than the reduction in total nitrogen. Thus, up to 15 days of nitrate withdrawal can lower nitrates without sharply reduce total nitrogen or affecting growth and biomass of cultivated cardoon. The use of N-free nutrient solution prior to harvest or the replacement of nitrates with chlorides could be adopted among growers to improve the quality of the product and enhance sustainability of crop production system.

Keywords: antioxidant capacity, Cynara cardunculus L. var. altilis DC, nitrate/chloride ratio, secondary metabolism, floating system, total phenols
INTRODUCTION

*Cynara cardunculus*, a Mediterranean perennial species within the *Asteraceae (Compositae)* family, includes the two cultivated taxa globe artichoke (var. *scolymus*) and cardoon (var. *altilis* DC), along with their progenitor the wild cardoon also called artichoke thistle [var. *sylvestris* (Lamk) Fiori] (Rottenberg and Zohary, 1996; Portis et al., 2005; Calabrese et al., 2012). The cultivation of cardoon is less diffused than that of globe artichoke; its production is located in the Mediterranean areas of southern Europe (e.g., Italy, France, Greece, Portugal, and Spain; Portis et al., 2005).

Traditionally, cardoon is cultivated for the fleshy leaf petiole and part of the central leaf vein used to prepare typical Italian dishes (Cravero et al., 2012). In addition to the described traditional uses, this botanical variety has been also considered as ‘multipurpose perennial crop’ for several industrial applications due to its high biomass productivity (29.3 t ha⁻¹; Ierna et al., 2012) and high dry matter content (Cravero et al., 2012). Therefore, in the last years this plant has been assessed as ‘biomass crop’ for paper pulp production, domestic heating, and power generation (Fernández et al., 2006; Ciancolini et al., 2013). Besides the high content of cellulose and hemicelluloses cultivated cardoon can be further valorized throughout the utilization of high-value extractable chemical components of the leaves. *Cynara* leaf extracts have been used since ancient times in folk medicine, due to their therapeutic potential as cholereic, hepatoprotective, antidiabetic and anticholestasis activities (Gebhardt and Beck, 1996; Gebhardt, 1997). These physiological properties have been mostly attributed to the content of phenylpropanoids (flavonoids, mono-, and di-cafeoylquinic acids) and sesquiterpene lactones (Lattanzio et al., 2009; Menin et al., 2010; Pandino et al., 2015).

Cardoon for leaf extract production is traditionally cultivated in Italy under open-field conditions with high-density plantation. Despite the relatively high interest of growers and pharmaceutical industries in cardoon leaf extracts, advanced cultivation techniques such as floating system have received little attention so far (Rouphael et al., 2012a; Colla et al., 2013a). Floating system is the most diffused cultivation soilless system for the production of fresh-cut leafy vegetables (Tomasi et al., 2015). Compared with traditional soil-based culture, floating system offers the possibility (i) to shorten the growing cycle with year round production, (ii) to reduce manpower cost, (iii) to avoid soil limitations (i.e., soil-borne pathogens, salinity, pollution), (iv) to grow plants at high density, (v) as well as to make more efficient the use of water and fertilizers (Falovo et al., 2009a,b; Tomasi et al., 2015). Moreover, manipulation of target molecules such as antioxidants and phytochemicals by management of nutrient solution is a research area attracting interest of many scientists (Fanasca et al., 2006a,b). Nutrient solution recipes used in hydroponics have been usually formulated with high concentration of nitrate—the main N source for most crops, to reach maximal yield (Gruda, 2009). Minimizing N supply, while keeping quantity and especially quality as high as possible represents a major sustainability challenge, since many growers are faced with environmental and economical concerns (Colla et al., 2010, 2011). Recently, many investigations have shown that low N availability increases for example the concentrations of phenolics and flavonoids in several leafy vegetables (Stefanelli et al., 2010; Becker et al., 2015). In this perspective, floating system represents an effective way since it permits the precise control of plant nutrition (Rouphael et al., 2012a; Colla et al., 2013a; Borgognone et al., 2014). For instance, nitrate concentration in the nutrient solution can be reduced by replacing nitrate with chloride or by eliminating it several days before harvest (DBH). To our knowledge, no published data is available concerning the effects of NO₃⁻:Cl⁻ ratio and nitrate deprivation before harvesting on cardoon yield and quality. Our hypothesis is that a controlled nitrogen limitation in the nutrient solution by nitrate withdrawal or replacement with chloride may raise the quality of cardoon while maintaining yield.

To verify the current hypothesis, two greenhouse experiments were carried out in order to assess the effect of a decreasing NO₃⁻:Cl⁻ ratio (Experiment 1) and nitrate deprivation (Experiment 2) on biomass production, leaf chlorophyll content and fluorescence, mineral composition, total phenols, flavonoids and antioxidant activity in leaves of cardoon grown in a floating system.

MATERIALS AND METHODS

Growth Conditions, Treatments, and Nutrient Solutions

Two experiments were conducted, one in 2013 (NO₃⁻:Cl⁻ ratio experiment; Experiment 1) and the other in 2013–2014 (time course N deprivation experiment; Experiment 2), in a 300 m × 300 m polyethylene greenhouse situated at the experimental farm of Tuscia University, central Italy (latitude 42° 25’ N, longitude 12° 08’), altitude 310 m). The greenhouse was maintained at daily temperature between 10 and 30°C and day/night relative humidity 55/80%.

In both experiments, seeds of *Cynara cardunculus* L. var. *altilis* cv. ‘Bianco Avorio’ (La Semiorto Sementi, Lavorate di Sarno, Italy) were sown in polystyrene plug trays on 27 August 2013 (Experiment 1) and 7 December 2013 (Experiment 2). ‘Bianco Avorio’ was selected due to its high productivity, leaf quality and the adaptability to grow in floating raft culture (Rouphael et al., 2012a; Colla et al., 2013a). At two–true-leaf stage [23 and 21 days after sowing (DAS) in Experiments 1 and 2, respectively] cardoon plants were moved to a floating raft growing system. The floating raft system consisted of the polystyrene plug trays floating in plastic tanks with a constant volume of 60 L of aerated nutrient solution by nitrate withdrawal or replacement with chloride may raise the quality of cardoon while maintaining yield.

A randomized complete-block design with four replicates was used to compare four different ratios of NO₃⁻ to Cl⁻ (80:20, 60:40, 40:60, or 20:80) in Experiment 1 and four time course nitrate deprivation (0, 5, 10, or 15 DBH) in Experiment 2. Each experimental unit consisted of polystyrene plug trays (0.1815 m²)
containing 84 plants. In Experiments 1 and 2, the composition of the basic nutrient solution was: 1.5 mmol L$^{-1}$ P, 4.5 mmol L$^{-1}$ K, 6.5 mmol L$^{-1}$ Ca, 2 mmol L$^{-1}$ Mg, 20 µmol L$^{-1}$ Fe, 9 µmol L$^{-1}$ Mn, 0.3 µmol L$^{-1}$ Cu, 1.6 µmol L$^{-1}$ Zn, 20 µmol L$^{-1}$ B, and 0.3 µmol L$^{-1}$ Mo. In Experiment 1, the four NO$_3$$^-$/Cl$^-$ ratios were obtained by adding to the basic nutrient solution different amounts of calcium nitrate and calcium chloride (Table 1). In Experiment 2, nitrates (10 mmol L$^{-1}$ NO$_3$) was removed totally from the nutrient solution at 0, 5, 10, or 15 DBH, and calcium nitrate (15.5% of N) was replaced by calcium chloride (analytical grade) in order to keep constant Ca concentration. In experiments, the electrical conductivity and pH of the nutrient solutions in all treatments were 2.0 ± 0.2 dS m$^{-1}$ and 6.0 ± 0.3, respectively. To prevent large fluctuation in the anions/cations concentrations, electrical conductivity and pH, the nutrient solution in all treatments were renewed from all tanks weekly.

Leaf Biomass Determination
In both experiments, the leaves of cultivated cardoon were mowed three times during the growing cycles (45, 73, and 101 DAS in Experiment 1 and 59, 97, and 124 DAS in Experiment 2) when the plant height reached 25 cm. At each harvest date, the leaf tissues were dried in a forced-air oven at 60°C for 72 h for dry biomass determination. In both experiments, the dried leaf tissue of each harvest was used for mineral and quality analysis.

SPAD Index and Fluorescence Measurements
In the experiments, the Soil Plant Analysis Development (SPAD) index and the fluorescence measurements were recorded at each harvest time. A portable chlorophyll meter (SPAD-502, Minolta Corporation, Ltd., Osaka, Japan) was used to measure the relative leaf chlorophyll concentration as a rational unit. Measurements were made at the central point of the leaflet between the midrib and the leaf margin of the third leaf starting from the apical shoot. Twenty random readings per plot were taken and averaged to a single SPAD value for each treatment.

On the same dates, the chlorophyll fluorescence was recorded on 20 min dark-adapted leaves (in 3rd – 4th from top; three measurements per plant) by means of a chlorophyll fluorimeter Handy PEA (Hansatech Instruments Ltd, King’s Lynn, UK) with an excitation source intensity higher than 3000 µmol photons m$^{-2}$ s$^{-1}$ at the sample surface. The minimal fluorescence intensity (F$_{0}$) in a dark-adapted state was measured in the presence of a background weak light signal (about 2–3 µmol photons m$^{-2}$ s$^{-1}$). The maximal fluorescence level in the dark-adapted state (F$_{m}$) was induced by 0.8 s saturating light pulse (3000 µmol photons m$^{-2}$ s$^{-1}$). The maximum quantum yield of open photosystem II (PSII) (F$_{v}$/F$_{m}$) was calculated as (F$_{m}$ – F$_{0}$)/F$_{m}$, as described by Arena et al. (2005).

Mineral Analysis
In both experiments, the dried leaf tissues of cultivated cardoon, harvested three times during the growing cycles, were ground in a Wiley mill to pass through a 20-mesh screen, then 1 g samples were analyzed for nitrogen and nitrate, and also for chloride but only in Experiment 1. Nitrogen concentration in leaf tissues was determined after mineralization with sulfuric acid in the presence of potassium sulfate and a low concentration of copper by the Kjeldahl method (Bremner, 1965). Nitrate concentration in dry leaf was determined using the salicylic-sulfuric acids method (Caldao et al., 1975) by spectrophotometry (Helios Beta Spectrophotometer, Thermo Electron Corporation, UK). Chloride ion concentration was determined by dry ashing at 400°C for 24 h, dissolving the ash in nitric acid (HNO$_3$, 1:20 w/v), and assaying the solution by titration with AgNO$_3$ in the presence of K$_2$CrO$_4$ (Eaton et al., 1995).

Analysis of Total Phenolics and Flavonoids
Dried cardoon leaf sample (100 mg) was extracted sequentially three times in 10, 5, and 5 mL of fresh ethanol/water (80:20 v/v) containing 1 mL L$^{-1}$ of 370 g L$^{-1}$ HCl (total volume 25 mL). The extraction mixture was stirred for 10 min and then solids were removed by centrifugation at 503 × g for 5 min. The total content of phenolic compounds in each extract was determined by the Folin–Ciocalteu method (Singleton and Rossi, 1965). A calibration curve was prepared using aliquots of gallic acid ethanolic solutions, and results were expressed as gallic acid equivalents (mg gallic acid g$^{-1}$ dry weight). All reagents and solvents used to prepare standard solutions and to extract phenols were of analytical grade and were purchased from Sigma Aldrich (Milan, Italy).

Total flavonoids were determined following the colorimetric aluminum chloride method reported by Dutta and Maharia (2012). A calibration curve was prepared using aliquots of quercitin (Sigma Aldrich, Milan, Italy), and results were expressed as quercetin equivalents (mg quercetin g$^{-1}$ dry weight).

Antioxidant Activity Determination
Antioxidant activity was determined by the ferric reducing-antioxidant power (FRAP) following the method described by Pellegrini et al. (2003). A calibration curve was prepared with increasing concentrations of FeSO$_4$ and results were expressed as µmol FeSO$_4$ g$^{-1}$ dry weight. All reagents were of analytical grade and were purchased from Sigma Aldrich (Milan, Italy).

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| Treatment NO$_3$-Cl$^-$ | NO$_3$ mM | Ca(NO$_3$)$_2$ mg l$^{-1}$ | CaCl$_2$ mg l$^{-1}$ |
|-------------------------|-----------|--------------------------|---------------------|
| 80: 20                  | 9.6       | 867.1                    | 133.2               |
| 60: 40                  | 7.2       | 650.3                    | 266.4               |
| 40: 60                  | 4.8       | 433.5                    | 399.6               |
| 20: 80                  | 2.4       | 216.8                    | 532.8               |
Statistical Analysis

Analysis of variance (ANOVA) of the experimental data was calculated using the software package, SPSS 10 for Windows, 2001 (SPSS Inc., Chicago, IL, USA). In both experiments, orthogonal linear and quadratic polynomial contrasts were used to compare the different ratios of NO$_3^-$ to Cl$^-$ and the nitrate deprivation effects at different DBH on selected parameters (Gomez and Gomez, 1983).

RESULTS

NO$_3^-$:Cl$^-$ Ratio Experiment

In Experiment 1, no significant difference in leaf dry biomass was observed between treatments when harvested at different DAS (data not shown). The mean leaf dry biomass of cardoon plants was 0.80, 0.63, and 0.42 g plant$^{-1}$ at 45, 73, and 101 DAS respectively, whereas the total biomass of cardoon recorded at the end of the experiment was 1.86 g plant$^{-1}$ DW (data not shown).

Similarly to dry biomass, the SPAD index and the maximum quantum use efficiency of the PSII in dark-adapted state measured as F$_{m}$/F$_{m}$ ratio were not affected by the NO$_3^-$:Cl$^-$ ratio. The mean SPAD index in leaves was 29.9, 31.4, and 26.6 at 45, 73, and 101 DAS, respectively, while the mean F$_{m}$/F$_{m}$ ratio in cardoon leaves was 0.84, 0.82 and 0.84, at 45, 73, and 101 DAS, respectively (data not shown).

The nitrogen (N), nitrate and chloride concentrations of cardoon ‘Bianco Avorio’ were significantly affected by the NO$_3^-$:Cl$^-$ ratio (Table 2). At 73 and 101 DAS, the N and nitrate concentration in leaves decreased linearly with increasing Cl$^-$ in the nutrient solution. The chloride concentration increased linearly with the increase of Cl$^-$ in the nutrient solution (Table 2).

Total phenolics (TP), total flavonoids (TF), and antioxidant activity (FRAP) were significantly affected by the NO$_3^-$:Cl$^-$ ratio, except for TF and FRAP at 101 DAS, where no significant effect was observed between treatments (Table 3). Cardoon leaf extract quality (i.e., TP, TF, and FRAP) increased linearly with Cl$^-$ availability, especially when nitrate was replaced by 80% of chloride (20:80; Table 3). However, for chloride concentration between 40 and 60%, the TP, TF, and FRAP remained unchanged. Finally, irrespective of the NO$_3^-$:Cl$^-$ ratio, the highest values of TP, TF, and FRAP were recorded at 101 DAS (Table 3).

Nitrate Deprivation Experiment

In Experiment 2, the dry mass of cardoon leaves and total biomass values were not significantly affected by nitrate deprivation from the nutrient solution (data not shown). The average leaf biomass production was 0.40, 0.82, and 0.46 g plant$^{-1}$ DW at 59, 97 and 124 DAS, respectively, whereas the mean total leaf dry biomass was 3.09 g plant$^{-1}$ (data not shown).

At the last harvest, 124 DAS, the SPAD index decreased linearly with the increase in the number of days of nitrate deprivation (Table 4). Moreover, there were no statistical differences between treatments in the fluorescence measurements (Table 4).

The N concentration in leaf tissue decreased linearly at 59 and 124 DAS and quadratically at 97 DAS, with increase in number of days of nitrate deprivation from the nutrient solution (Table 5). Similarly, the nitrate concentration in leaves of cardoon plants, harvested at different DAS decreased linearly with nitrate withdrawal from the nutrient solution with a sharp reduction (83–87%) observed after 15 days of nitrate deprivation in comparison to control treatment (Table 5).

The bioactive compounds (i.e., TP and TF) and antioxidant activity of cardoon leaf extract harvested at different DAS increased linearly, with the highest values recorded in the 15 days of nitrate deprivation treatment (Table 6). Finally, irrespective to nitrate withdrawal from the nutrient solution, the highest values of TP, TF, and FRAP were recorded at 97 DAS (Table 6).

DISCUSSION

Management of the growing conditions, in particular the nutrient solution composition, is one of the most important aspect for successful leafy crop production in floating raft culture (Fallovo et al., 2009b; Rouphael et al., 2012a). In the current experiments, we have illustrated that minimizing the nitrate supply either by partial substitution of nitrate with chloride, or by nitrate deprivation several DBH, could be proposed as an effective tool to raise the quality of the cardoon leaves while maintaining yield.

It is well established that biomass production and marketable yield decrease with increasing salinity, as it has been demonstrated in many sensitive to moderately salt-tolerant

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**TABLE 2** | Effect of NO$_3^-$:Cl$^-$ ratio on total nitrogen (N), nitrate (NO$_3^-$), and chloride (Cl$^-$) contents in leaves of cardoon plants (*Cynara cardunculus* L. var. *altillis* DC) harvested at different days after sowing (DAS); cardoon plants were grown in a floating system under greenhouse conditions (Experiment 1).

| Treatment | N (g kg$^{-1}$ DW) 45 DAS | N (g kg$^{-1}$ DW) 73 DAS | N (g kg$^{-1}$ DW) 101 DAS | NO$_3^-$ (g kg$^{-1}$ DW) 45 DAS | NO$_3^-$ (g kg$^{-1}$ DW) 73 DAS | NO$_3^-$ (g kg$^{-1}$ DW) 101 DAS | Cl$^-$ (g kg$^{-1}$ DW) 45 DAS | Cl$^-$ (g kg$^{-1}$ DW) 73 DAS | Cl$^-$ (g kg$^{-1}$ DW) 101 DAS |
|-----------|--------------------------|--------------------------|--------------------------|------------------|--------------------------|--------------------------|------------------|--------------------------|--------------------------|
| 80:20     | 42.48                    | 41.50                    | 46.91                    | 16.78            | 16.63                    | 10.53                    | 29.03            | 27.97                    | 27.23                    |
| 60:40     | 44.32                    | 40.30                    | 43.91                    | 12.70            | 14.36                    | 8.67                     | 31.72            | 35.02                    | 33.38                    |
| 40:60     | 41.89                    | 36.76                    | 42.23                    | 10.36            | 10.67                    | 7.01                     | 38.07            | 41.10                    | 35.09                    |
| 20:80     | 41.53                    | 35.96                    | 37.11                    | 8.82             | 7.56                     | 7.66                     | 48.97            | 40.49                    | 38.83                    |
| Significance | NS                      | L**                      | L***                     | L**              | L**                      | L***                     | L***             | NS                      | NS                      |

$L$, linear; $Q$, quadratic; $Ns$, non-significant; $^{*}P < 0.05$; $^{**}P < 0.01$; $^{***}P < 0.001$. 
TABLE 3 | Effect of NO$_3^-$:Cl$^-$ ratio on total phenols, total flavonoids and ferric-reducing antioxidant power (FRAP) in leaves of cardoon plants (Cynara cardunculus L. var. altilis DC) harvested at different days after sowing (DAS); cardoon plants were grown in a floating system under greenhouse conditions (Experiment 1).

| Treatment | Total phenols (mg gallic acid g$^{-1}$ DW) | Total flavonoids (mg quercitin g$^{-1}$ DW) | FRAP (µmol FeSO$_4$ g$^{-1}$ DW) |
|-----------|---------------------------------|---------------------------------|--------------------------------|
| NO$_3^-$:Cl$^-$ | 45 DAS | 73 DAS | 101 DAS | 45 DAS | 73 DAS | 101 DAS | 45 DAS | 73 DAS | 101 DAS |
| 80: 20 | 7.63 | 13.97 | 17.22 | 3.27 | 21.36 | 38.16 | 92.52 | 217.20 | 290.97 |
| 60: 40 | 9.03 | 15.04 | 19.34 | 6.97 | 24.69 | 45.89 | 117.11 | 247.37 | 318.04 |
| 40: 60 | 9.51 | 16.35 | 19.74 | 7.26 | 29.94 | 44.79 | 125.46 | 283.88 | 322.17 |
| 20: 80 | 10.59 | 22.09 | 21.03 | 8.82 | 49.55 | 54.62 | 144.65 | 390.45 | 371.98 |
| Significance | L$^*$ | L$^{**}$ | L$^*$ | NS | L$^*$ | L$^{**}$ | NS | L$^*$ | L$^{***}$ | NS |

L, linear; NS, non-significant; *P < 0.05; **P < 0.01; ***P < 0.001.

TABLE 4 | Effect of nitrate deprivation at different days before harvest (DBH) on SPAD index and maximum quantum use efficiency of PSII ($F_v/F_m$) in dark-adapted state of cardoon plants (Cynara cardunculus L. var. altilis DC) harvested at different days after sowing (DAS); cardoon plants were grown in a floating system under greenhouse conditions (Experiment 2).

| Nitrate deprivation | SPAD index | $F_v/F_m$ |
|---------------------|------------|-----------|
| DBH                 | 59 DAS     | 97 DAS    | 124 DAS   | 59 DAS | 97 DAS | 124 DAS |
| 0                   | 28.5       | 32.6      | 34.7      | 0.80  | 0.84  | 0.83   |
| 5                   | 25.4       | 34.2      | 33.5      | 0.81  | 0.84  | 0.83   |
| 10                  | 27.1       | 36.5      | 30.8      | 0.81  | 0.84  | 0.83   |
| 15                  | 25.4       | 33.7      | 28.0      | 0.77  | 0.84  | 0.83   |
| Significance         | NS         | NS        | L$^{***}$ | NS   | NS   | NS     |

L, linear; NS, non-significant; ***P < 0.001.

TABLE 5 | Effect of nitrate deprivation at different days before harvest (DBH) on nitrogen (N), and nitrate (NO$_3^-$), contents in leaves of cardoon plants (Cynara cardunculus L. var. altilis DC) harvested at different days after sowing (DAS); cardoon plants were grown in a floating system under greenhouse conditions (Experiment 2).

| Nitrate deprivation | N (g kg$^{-1}$ DW) | NO$_3^-$ (g kg$^{-1}$ DW) |
|---------------------|--------------------|--------------------------|
| DBH                 | 59 DAS | 97 DAS | 124 DAS | 59 DAS | 97 DAS | 124 DAS |
| 0                   | 49.17 | 38.56 | 43.79 | 19.87 | 10.91 | 8.89   |
| 5                   | 41.54 | 42.17 | 34.96 | 17.83 | 5.63  | 5.76   |
| 10                  | 41.49 | 43.73 | 35.83 | 14.53 | 3.91  | 4.66   |
| 15                  | 39.00 | 29.34 | 31.56 | 11.74 | 1.42  | 1.52   |
| Significance         | L$^*$   | Q$^*$   | L$^{**}$ | L$^{***}$ | L$^{***}$ | L$^{***}$ |

L, linear; Q, quadratic; NS, non-significant; *P < 0.05; **P < 0.01; ***P < 0.001.

TABLE 6 | Effect of nitrate deprivation at different days before harvest (DBH) on total phenols, total flavonoids and ferric-reducing antioxidant power (FRAP) in leaves of cardoon plants (Cynara cardunculus L. var. altilis DC) harvested at different days after sowing (DAS); cardoon plants were grown in a floating system under greenhouse conditions (Experiment 2).

| Nitrate deprivation | Total phenols (mg gallic acid g$^{-1}$ DW) | Total flavonoids (mg quercitin g$^{-1}$ DW) | FRAP (µmol FeSO$_4$ g$^{-1}$ DW) |
|---------------------|---------------------------------|---------------------------------|--------------------------------|
| DBH                 | 59 DAS | 97 DAS | 124 DAS | 59 DAS | 97 DAS | 124 DAS | 59 DAS | 97 DAS | 124 DAS |
| 0                   | 10.00  | 23.18  | 15.23   | 10.67  | 38.55  | 12.21   | 130.11 | 401.82 | 194.05 |
| 5                   | 11.27  | 20.34  | 17.11   | 14.80  | 25.34  | 18.08   | 157.58 | 343.02 | 231.51 |
| 10                  | 11.10  | 25.03  | 20.09   | 14.42  | 42.20  | 30.30   | 151.25 | 438.93 | 294.07 |
| 15                  | 14.31  | 33.12  | 21.99   | 25.21  | 75.51  | 31.85   | 221.81 | 566.17 | 314.74 |
| Significance         | L$^*$  | L$^*$  | L$^{***}$ | L$^{**}$ | L$^*$  | L$^{***}$ | L$^*$  | L$^{***}$ |

L, linear; *P < 0.05; **P < 0.01; ***P < 0.001.

greenhouse vegetables such as cucumber, melon, pepper, watermelon and zucchini squash grown hydroponically (Colla et al., 2006a,b,c, 2012, 2013b; Rouphael et al., 2006, 2012c). Ions, mainly Na$^+$ and/or Cl$^-$ in high concentration generates oxidative stress in plant tissues as revealed by the production of reactive oxygen species (ROS; Sreenivasulu et al., 2000) leading...
to several morphological, physiological and biochemical changes (Gratton and Grieve, 1999). High levels of Na\(^+\) and/or Cl\(^-\) in the root zone decreased the net nitrate uptake rate and nitrate translocation to the shoot in many vascular plants. This was the case in the current study (Experiment 1), since replacing nitrate by 80% of chloride (20:80) decreased nitrates in cardoon leaves as demonstrated in previous studies on leafy vegetables grown hydroponically (Urrestarazu et al., 1998; Neocleous et al., 2014).

Touraine et al. (1994) reported that salinity can reduce the growth rates of the plants, leading to an indirect downregulation of net nitrate uptake rate via a lower internal demand for N. This was not the case of Experiment 1 since shoot dry biomass was unaffected to any level of chloride in the nutrient solution. Rubinigg et al. (2003) found that root to shoot translocation of N was severely inhibited relative to the net nitrate uptake ratio at all NaCl concentrations applied. Anion channels with similar permeability for both NO\(_3\)\(^-\) and Cl\(^-\) have been found in xylem parenchyma cells from barley (Köhler and Raschke, 2000). These channels were suggested to play an important role in the xylem loading of NO\(_3\)\(^-\) and Cl\(^-\). In the presence of high Cl\(^-\) levels, root to shoot translocation of NO\(_3\) could therefore be decreased at the site of entrance into the xylem via competition for the same channel (Rubinigg et al., 2003). Moreover, the decrease of nitrogen could also be due to a reduction in nitrate influx resulting from an interference of external chloride with one or more types of transporters at the root plasma. Accumulation of chloride in leaf tissue at the highest Cl\(^-\)/NO\(_3\)\(^-\) ratio did not reduce the total leaf biomass, confirming that cardoon can be considered as a salt-tolerant crop (Ksouri et al., 2012; Pace et al., 2012).

It has been generally accepted that the critical toxicity concentration of chloride in leaf tissues ranges from 4 to 7 and from 15 to 50 g kg\(^{-1}\) on dry weight basis for Cl-sensitive and Cl-tolerant species, respectively (Marschner, 2012). In the present work chloride concentration in cardoon leaves recorded in the 20:80 NO\(_3\)\(^-\)/Cl\(^-\) treatment was below the critical threshold (e.g., 50 g kg\(^{-1}\) DW; Table 2); this could explain the lack of biomass reduction recorded under high chloride concentration in the nutrient solution.

High chloride concentration in the nutrient solution interferes with several aspects of plant physiology including pigment biosynthesis and the activity of the photosystem (PS) II (Colla et al., 2010). Since the response of NO\(_3\)\(^-\)/Cl\(^-\) ratio depends on the cardoon ability to counteract the effects induced by the four nitrate to chloride ratios, the lack of changes between treatments on the leaf dry biomass should be also reflected at physiological level. This was indeed the case, since with no significant difference between the four nitrate to chloride ratios was observed for chlorophyll content (SPAD index) and the maximum quantum efficiency of PSII (\(F_V/F_M\)), frequently used as an indicator of the photoinhibition or stress damage to the PSII (Calatayud and Barreno, 2004; Colla et al., 2013b).

Nutraceutical and pharmaceutical properties of cardoon leaves are linked to the high concentration of antioxidant molecules in particular phenolic acids and flavonoids, as well as their high antioxidant activity (Rouphael et al., 2012a; Colla et al., 2013a; Borgognone et al., 2014). Moreover, the concentration of bioactive compounds (e.g., flavonoids and phenolic acids) varies significantly in relation to genetic, environmental, agronomic, and biotic/abiotic factors (Pandino et al., 2010; Rouphael et al., 2012b). Under stress conditions plants activate a series of counteracting measures including physiological and molecular mechanisms that consent acclimation to a suboptimal environment (Atkinson and Urwin, 2012). Among these measures, the accumulation of specific molecules and secondary metabolites with multiple functions has a crucial role of ensuring plant growth under unfavorable conditions. These antioxidants molecules are thought to contribute in ROS scavenging and cellular water homeostasis (Bandyopadhyay et al., 2012; Sharma et al., 2012). More interestingly, the highest phenolics and flavonoids concentration recorded under high chloride concentration (Table 3) are also important to human health, and therefore they attribute an added value (compared to the control) to the nutraceutical properties of cardoon leaf extract. These results are very important because several pharmacological studies have demonstrated that cardoon extracts are supposed to exert antimicrobial, anti-inflammatory, anti-oxidative, anti-carcinogenic, bile-expelling and urinative activities, as well as the ability to inhibit low-density lipoprotein (LDL) oxidation and cholesterol biosynthesis (Fallah et al., 2008; Kukic et al., 2008; Kammoun et al., 2010; Mileo et al., 2012).

Horticultural crops are usually supplied with high concentrations of inorganic nutrients, particularly nitrates which are the major anions responsible for biomass accumulation (Le Bot et al., 2001). Although this practice prevents the growth from being limited by mineral supply, it can lead to excessive plant vigor, decreasing quality, causing delay in harvest, nitrate accumulation in leaf tissues and high environmental impact (Stewart et al., 2001; Chen et al., 2004; Becker et al., 2015). An effective and sustainable fertigation strategy to limit nitrate waste in the environment while maintaining yield could be to terminate nitrate supply in the nutrient solution several days prior to final harvest. Nitrate withdrawal in the solution few DBH has been successfully used for lowering nitrate content in leafy vegetables with or without affecting crop productivity (Santamaria et al., 1998; Gonnella et al., 2004; Gent, 2012). In fact, the results of Experiment 2 clearly demonstrated that it is possible to withdraw nitrates in the nutrient solution from 5 up to 15 DBH without any negative effects on leaf dry biomass. Therefore, the standard nutrient recipes normally used in floating raft culture appear to be luxurious (Le Bot et al., 2001). Our results are in line with those of Santamaria et al. (1998) and Gonnella et al. (2004), who observed that reducing and/or eliminating nitrates in the nutrient solution few days (2–5) before harvesting can significantly reduced leaf nitrate concentration without affecting yield of chicory, rocket and lamb’s lettuce. ‘This acclimation appears the result of upregulation of nutrient uptake by plant roots as well as increasing in root biomass’ (Siddiqi et al., 1998) and may also depend on the plant species. Moreover, nitrogen limitation can negatively affect photosynthesis by decreasing available chlorophyll and disturbing photosynthesis owing to starch accumulation (Stewart et al., 2001). This was partially observed in the Experiment 2 since the chlorophyll content was...
not influenced by nitrate deprivation at 59 and 97 DAS, but was only affected at 124 DAS (Table 4).

The decrease in leaf nitrate content observed under N-deprivation was more pronounced than the reduction in total nitrogen (Table 5). Thus, up to 15 days of nitrate withdrawal could lower nitrate without sharply reducing total nitrogen or affecting growth and biomass of cultivated cardoon. These results are consistent with the findings of Gent (2012) who demonstrated that tissue nitrate changed more rapidly than total N over a 6-day interval of nitrogen depletion without affecting yield of hydroponic lettuce. The lowest nitrate concentration in cultivated cardoon leaves after 15 days of N-deprivation would suggest that a significant part of nitrate was assimilated in the ‘metabolic pool’ since the external NO₃⁻ is exhausted, while little would have remained available to be stored in cell vacuoles or ‘reserve pool’ as osmotic regulator (Santamaria et al., 1998). Contrarily, in the control treatment, the highest nitrate concentration in leaf tissue indicates that a greater part of nitrate was stored in the ‘reserve pool’ (Izmailov et al., 1992).

In the present study, it was also found that nitrate application can be eliminated 10–15 DBH with a significant increase in cardoon leaf quality as expressed by total phenols, ferric-reducing antioxidant power (FRAP) and total flavonoids (Table 6). It has been demonstrated that nutrient deficiency resulted in enhancing phenolic compounds in different leafy vegetables due to upregulation of the phenylpropanoid pathway, whose key enzyme, phenylalanine ammonia-lyase (PAL) is the link between primary and secondary metabolism (Oh et al., 2009; Becker et al., 2015). In a recent review, Stefanelli et al. (2010) elucidated how high nitrogen availability could inhibit phenolic production and subsequently antioxidant capacity, in several leafy vegetables. In line with our findings, highest total plant phenolics (TPP) in Chinese cabbage were observed in plants grown in soil without N fertilization while higher N application (Nguyen and Niemeyer, 2008; Becker et al., 2015). Moreover, highest phenolic content in basil, and leaf lettuce grown hydroponically was also found after minimal nitrogen application (Nguyen and Niemeyer, 2008; Becker et al., 2015). Similarly to total phenols, the total flavonoids in leaf tissue increased linearly after 5, 10, or 15 days of nitrate depletion (Table 6). Increased flavonoids accumulation under low N availability has been reported previously in basil, broccoli and tomato leaves (Stewart et al., 2001; Jones et al., 2007; Nguyen and Niemeyer, 2008). An explanation for increased flavonoids synthesis under nitrogen stress is the possible activation of the flavonoids pathway through a number of enzymes and genes (Bongue-Bartelsman and Phillips, 1995) but does not have a pronounced influence on the shikimate pathway which produce phenylalanine (Lillo et al., 2008). It could be also expected that N deficiency (i.e., deprivation) could increase concentrations of flavonoids as they could be involved in the ROS (i.e., hydrogen peroxide) scavenging cascade (Takahama and Oniki, 1997).

CONCLUSION

The results of Experiments 1 and 2 showed that floating raft culture appears to be a promising tool to obtain valuable cardoon production as well as improving quality aspects of leaf through proper management of nitrate in the nutrient solution. The use of nutrient solution having a NO₃⁻:Cl⁻ ratio of 20:80 could be adopted in the present conditions to improve the quality aspects of cardoon leaves by increasing antioxidant activity, total phenols and flavonoids and by lowering the nitrate concentration, with no negative impact on yield. The results also indicated that it is possible to withdraw nitrate from the nutrient solution 5, 10 and even 15 DBH without having a detrimental effect on yield. In addition, this will increase leaf quality by increasing total phenols, flavonoids and antioxidant capacity, especially when nitrate deprivation occurred for 15 DBH. Finally, it must be pointed out that nitrate withdrawn strategy may be valid provided that enough nitrate are accumulated in the plant tissues during the first part of the growing cycle.

AUTHOR CONTRIBUTIONS

DB conducted the trial as part of her Ph.D. program. YR was involved in the agronomic measurements and interpretation of the results. MC conducted the mineral analysis and interpretation of the results. LL made the analysis of total phenolics and flavonoids. GC set up the research program and he coordinated the research group in conducting the trial, statistical analysis, interpretation of results. All authors were involved in writing the manuscript and statistical analysis.

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Bongue-Bartelsman, M., and Phillips, D. A. (1995). Nitrogen stress regulates gene expression of enzymes in the flavonoid biosynthetic pathway of tomato. Plant Physiol. Biochem. 33, 539–546.

Borgognone, D., Cardarelli, M., Lucini, L., and Colla, G. (2014). Does CaCl$_2$ play a role in improving biomass yield and quality of cardoon grown in a floating system under saline conditions? HortScience 49, 1523–1528.

Bremner, J. M. (1965) “Total nitrogen,” in Methods of Soil Analysis (Agronomy Monograph No. 9, Part 2), eds C. A. Black, D. D. Evans, I. L. White, L. E. Ensminger, and F. E. Clark (Madison, WI: American Society of Agronomy) 1119–1178.

Calabrese, N., Carito, A., Boari, F., Cantineo, V., and Damato, G. (2011). Does nitrate solution concentration and growing season affect yield and quality of Lactuca sativa L. var. acephala in floating raft culture. J. Sci. Food Agric. 89, 1682–1689. doi: 10.1002/jsfa.3641

Fallas, S., Colla, G., Maiani, G., Venneria, E., Battistelli, A., and Colla, G. (2009a). Yield and quality of leafy lettuce in response to nutrient solution composition and growing season. J. Agric. Food Environ. 7, 456–462.

Fallas, C., Rouphael, Y., Rea, E., Battistelli, A., and Colla, G. (2009b). Nutrient solution concentration and growing season affect yield and quality of Lactuca sativa L. var. acephala in floating raft culture. J. Sci. Food Agric. 89, 1682–1689. doi: 10.1002/jsfa.3641

Fanasca, S., Colla, G., Maiani, G., Venneria, E., Azzini, E., et al. (2006a). Changes in antioxidant content of tomato fruits in response to cultivar and nutrient solution composition. J. Agric. Food Chem. 54, 4319–4325. doi: 10.1021/jf0502572

Fallas, S., Colla, G., Rouphael, Y., Saccardo, F., Maiani, G., Venneria, E., et al. (2006b). Evolution of nutritional value of two tomato genotypes grown in soilless culture as affected by macronutrient proportions. HortScience 41, 1584–1588.

Fernández, J., Curt, M. D., and Aguado, P. L. (2006). Industrial applications of Cynara cardunculus L. for energy and other uses. Ind. Crops Prod. 24, 222–229. doi: 10.1016/j.indcrop.2006.06.010

Gebhardt, R. (1997). Antioxidative and protective properties of extracts from leaves of the artichoke against hydroperoxide-induced oxidative stress in cultured rat hepatocytes. Toxicol. Appl. Pharmacol. 144, 279–286. doi: 10.1006/taap.1997.8130

Gebhardt, R., and Beck, H. (1996). Differential effects of garlic-derived organosulfides on cholesterol biosynthesis in primary rat hepatocytes cultures. Lipids 31, 1267–1275. doi: 10.1016/0020-2615(95)00186-0

Gent, M. P. N. (2012). Rate of change of composition of lettuce in response to nitrogen depletion or re-supply. J. Sci. Food Agric. 92, 3007–3015. doi: 10.1002/jsfa.5716

Gomez, A. A. and Gomez, A. A. (1983). “Comparison between treatments means,” in Statistical Procedures for Agricultural Research, 2nd Edn, eds K. A. Gomez and A. A. Gomez (New York, NY: John Wiley & Sons), 187–240.

Gonnella, M., Conversa, G., Santamaría, P., and Serio, F. (2004). Production and nitrate content in lamb’s lettuce grown in floating system. Acta Hortic. 644, 61–68. doi: 10.17660/ActaHortic.2004.644.5

Grattan, S. R., andrieve, C. M. (1999). Salinity-mineral nutrient relations in horticultural crops. Sci. Hortic. 78, 127–157. doi: 10.1016/S0166-4482(98)00192-7

Gruda, N. (2009). Does soilless culture have an influence on product quality of vegetables? J. Appl. Bot. Food Qual. 82, 141–147.

Ierna, A., Mauro, R. P., and Mauricomeale, G. (2012). Biomass, grain and energy yield in Cynara cardunculus L. as affected by fertilization, genotype and harvest time. Biomass Bioenergy 36, 404–410. doi: 10.1016/j.biombioe.2011.11.013

Izmailov, S. F., Drobysheva, N. L., and Ovcharenko, G. A. (1992). Temporal and functional characteristics of saturation and utilization of nitrate pools in pea leaves. Soviet Plant Physiol. 39, 559–564.

Jones, R. B., Imisc, M., Franz, P., and Tomkis, B. T. (2007). High nitrogen during growth reduced glucoraphanin and flavonol content in broccoli (Brassica oleracea var. italic) heads. Aust. J. Exp. Agric. 47, 1498–1505. doi: 10.1071/EA06205

Kammoun, M., Koubab, I., Ben Ali, Y., Jarrajaa, R., Gargoury, I., Damak, M., et al. (2010). Inhibition of pro-inflammatory secreted phospholipase A2 by extracts from Cynara cardunculus L. Appl. Biochem. Biotechnol. 162, 662–670. doi: 10.1007/s12010-009-8849-8

Köhler, B., and Raschke, K. (2000). The delivery of salts to the xylem. Three types of anion conductance in the plasmalemma of the xylem parenchyma of roots of barley. Plant Physiol. 122, 243–254. doi: 10.1104/pp.122.1.243

Ksouri, R., Ksouri, W. M., Jallali, I., Debez, A., Magné, C., Hiroko, I., et al. (2012). Globe artichoke: a functional food and source of nutraceutical ingredients. J. Funct. Food 1, 131–144. doi: 10.1016/j.jff.2009.01.002
Marschner, P. (2012). *Cardoon Response to Nitrate Deprivation*. Elsevier, 651.

Menin, B., Comino, C., Moglia, A., Dolzhenko, Y., Portis, E., and Lanteri, S. (2010). Identification and mapping of genes related to caffeoylquinic acid synthesis in *Cynara cardunculus*. *Plant Sci.* 178, 338–347. doi: 10.1016/j.plantsci.2010.06.010

Mileo, A. M., Di Venere, D., Abbruzzese, C., and Miccadei, S. (2012). Long term exposure to polyphenols of artichoke (*Cynara scolymus*) exerts induction of senescence driven growth arrest in the MDA-MB231 human breast cancer cell line. *Oxid. Med. Cell. Long.* 2015:11. doi: 10.1155/2015/363827

Neocleous, D., Koukounaras, A., Siomos, A. S., and Vasilakakis, M. (2014). Assessing the salinity effects on mineral composition and nutritional quality of green and red 'Baby' lettuce. *J. Food Qual.* 37, 1–8. doi: 10.1111/jfq.12066

Nguyen, P. M., and Niemeyer, E. D. (2008). Effects of nitrogen fertilization on the flavonoid pathway. *Plant Cell Environ.* 47, 578–593. doi: 10.1111/j.1399-3054.2008.01748.x

Rouphael, Y., Cardarelli, M., Bassal, A., Leonardi, C., Giuffrida, F., and Colla, G. (2012). Improving growth performance under salinity stress by grafting onto *Cucurbita* hybrid rootstocks. *Photosynthetica* 50, 180–188. doi: 10.1007/s11099-012-0002-1

Rubinig, M., Posthumus, F., Ferschke, M., Theo, J., Elzenga, M., and Stulen, I. (2003). Effects of NaCl salinity on 15N-nitrate fluxes and specific root length in the halophyte *Plantago maritima*. *Plant Soil* 250, 201–213. doi: 10.1023/A:1022849928781

Santamaría, P., Elia, A., Parente, A., and Serio, F. (1998). Fertilization strategies for lowering nitrate content in leafy vegetables. *Chicory and rocket salad cases*. *J. Plant Nutr.* 21, 1791–1803. doi: 10.1080/01904169809365524

Sharma, P., Jha, A. B., Dubey, R. S., and Pessarakli, M. (2012). Reactive oxygen species, oxidative damage, and antioxidant defense mechanism in plants under stressful conditions. *J. Bot.* 2012:217037.

Stiddi, M. Y., Kronzucker, H. J., Britto, D. T., and Glass, A. D. M. (1998). Growth of a tomato crop at reduced nutrient concentrations as a strategy to limit eutrophication. *J. Plant Nutr.* 21, 1879–1895. doi: 10.1080/01904169809365530

Singleton, V. L., and Rossi, J. A. (1965). Colorimetry of total phenolics with phosphomolybdic–phosphotungstic acid reagent. *Am. J. Enol. Vitic.* 16, 144–158.

Sreenivasulu, N., Grimm, B., Wobus, U., and Weschke, W. (2000). Differential response of antioxidant compounds to salinity stress in salt tolerant and salt sensitive seedlings of foxtail millet (*Setaria italica*). *Physiol. Plant.* 109, 435–442. doi: 10.1043/1399-3054/2000.100410.x

Stefanelli, D., Goodwin, L., and Jones, R. (2010). Minimal nitrogen and water use in horticulture: effects on quality and content of selected nutrients. *Food Res. Inter.* 43, 1833–1843. doi: 10.1016/j.foodres.2010.04.022

Stewart, A. J., Chapman, W., Jenkins, G. L., Graham, I., Martin, T., and Crozier, A. (2001). The effect of nitrogen and phosphorus deficiency on flavonol accumulation in plant tissue. *Plant Cell Environ.* 24, 1189–1197. doi: 10.1046/j.1365-3040.2001.00768.x

Takahama, U., and Onuki, T. (1997). A peroxidase/phenolics/ascorbate system can scavenge hydrogen peroxide in plant cells. *Physiol. Planta.* 101, 845–852. doi: 10.1043/1399-3054.1997.1010423.x

Tomasi, N., Pinton, R., Dalla Costa, L., Cortella, G., Terzano, R., Mimmo, T., et al. (2015). New solutions for floating cultivation system of ready-to-eat salad: a review. *Trends Food Sci. Technol.* 46, 267–276. doi: 10.1016/j.tifs.2015.08.004

Touraine, B., Clarkson, D. T., and Muller, B. (1994). “Regulation of nitrate uptake at the whole plant level,” in *A Whole Plant Perspective on Carbon-Nitrogen Interactions*, eds J. Roy and E. Garnier (Hague: SPB Academic Publishing), 11–30.

Urestarazu, M., Postigo, A., Salas, M., Sánchez, A., and Carrasco, G. (1998). Nitrate accumulation reduction using chloride in the nutrient solution on lettuce growing by NFT in semiarid climate conditions. *J. Plant Nutr.* 21, 1705–1714. doi: 10.1080/01904169809365515

Zhu, W., Lin, X., Jin, C., Zhang, Y., and Fang, P. (2009). Effects of nitrogen application rates on antioxidant contents and antioxidative activities in Chinese cabbage (*Brassica chinensis* L.). *J. Zhejiang Univ. (Agric. Life Sci.)* 35, 3529–35306.

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