Effects of cultivars and coco-substrates on soilless strawberry production in Cyprus

Damianos Neocleous∗
Agricultural Research Institute, Ministry of Agriculture, Natural Resources and Environment, Nicosia, Cyprus

Received 12 March 2012; accepted 24 March 2012

Abstract. This experiment was conducted to generate up-to-date, practical and location specific information for soilless strawberry production in Cyprus. Commercial ‘fresh’ strawberry plants (Fragaria × ananassa Duch) of three cultivars (‘Camarosa’, ‘Festival’, and ‘Ventana’) were cultivated from October 2010 to June 2011, using coco-substrates from three different brands (BVB, Wonder Soil, Pelemix) in a 2 levels-pyramid layout. Harvesting started in January and lasted until June. The total yield obtained was similar in all cultivars and of about 439 g plant\(^{-1}\). However, with ‘Ventana’ and ‘Festival’ ‘extra’ category fruits and berry mass were advanced. Early yield (Jan–April) was higher in ‘Festival’ compared with ‘Camarosa’ and ‘Ventana’ was in between. Amount of water consumed to produce one kg of fruit fresh weight (WUE), was lower in ‘Ventana’ and ‘Festival’ compared to ‘Camarosa’. No differences were observed in productive characteristics, early yield and WUE according to the substrate. Regarding quality, fruits of ‘Festival’ and ‘Ventana’ retained higher soluble solids to acid ratio, whereas fruits of ‘Camarosa’ performed higher levels of bioactive compounds. The results from this study suggest ‘Festival’ and ‘Ventana’ as interesting alternative cultivars.

Keywords: Fragaria × ananassa, coconut fibre, yield, water use, quality

1. Introduction

During the last years, the interest in soilless culture in Mediterranean countries is increasing due to exhausted soils, soil disinfection, and water quality and availability [18, 20]. Strawberries are the most important berry crop cultivated in Europe and one of the most important crops cultivated in soilless systems [21]. Moreover, strawberry production and growth areas increase each year not only because of its highly consumable fruits, but also for the nutritional value of the fruit-rich in antioxidants [21, 26]. It is estimated that soilless culture in Cyprus represents 45.7 ha (10% of the total greenhouse cultivation area). This is relatively low and likely due to growers having knowledge of greenhouse production and the use of substrates [19]. The cultivated strawberry is the most important berry crop in Cyprus accounting for 32 ha and 1840 ton production [7]. Moreover, it is one of the most important crops cultivated under soilless conditions, with ‘Camarosa’ consisting the prevailing cultivar the last decade, using fresh plants that are autumn planted. Soilless culture aims to achieve an efficient growing system for strawberries using as substrate slabs of coconut fibre or rockwool. Coconut fibre is widely used in many countries as it is easy to use, produces good yield, is a renewable resource that can be incorporated to the soil and re-use is possible [17]. However, the need of the growers for improving production, earliness and quality of the fruits is essential to enhance fresh consumption.

∗ Corresponding author: Damianos Neocleous, Agricultural Research Institute, Ministry of Agriculture, Natural Resources and Environment, Nicosia, Cyprus. Tel.: +357 22 403115; Fax: +357 22 316770; E-mail: d.neocleous@arinet.ari.gov.cy.

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The aim of this experiment was to generate up-to-date, practical and location specific information for soilless strawberry production in Cyprus. Particularly the effects of three cultivars (‘Camarosa’, ‘Festival’, ‘Ventana’) and coconut fibre substrates from three different brands (BVB, Pelemix, Wonder Soil) on plant productivity, earliness, water use efficiency and fruit quality was evaluated.

2. Materials and methods

2.1. Plant material and growth conditions

Commercial ‘fresh’ strawberry plants \( (Fragaria \times ananassa \text{ Duch}) \) of three cultivars (‘Camarosa’, ‘Festival’, ‘Ventana’) were cultivated in a 567 m² plastic greenhouse (ARI, Cyprus, 34°94′N, 33°19′E) using coco-substrates from three different brands (BVB, Wonder Soil and Pelemix) wrapped in polyethylene slabs \((100 \text{ cm} \times 15 \text{ cm} \times 7 \text{ cm})\) and placed on Polygal troughs in a 2 levels-pyramid layout \((20.6 \text{ plants m}^{-2})\). The average temperature and humidity during the experimental period from October, 2010 to June, 2011, were 22°C and 75%. The mean global solar radiation was 17.5 MJ m\(^{-2}\).

2.2. Handling and harvesting

Plants were supplied with a known amount of nutrient solution, the drainage \((30\% \text{ run-off})\) was measured and the difference was the water consumption by the plants. The irrigation schedule was to keep the electrical conductivity in the root environment relatively constant. Water use efficiency (WUE) was defined as the amount of water consumed to produce one kg of fruit fresh weight. A nutrient solution consisting of the following concentrations of macronutrients \((\text{mmol/l}): 5.2 \text{ K, 3.5 Ca, 1.5 Mg, 12 N-NO}_3, 0.6 \text{ N-NH}_4, 1.25 \text{ P-H}_2\text{PO}_4, 1.15 \text{ S-SO}_4, \text{ and trace elements (\text{mmol/l}): 10 Mn, 20 Fe, 4 Zn, 0.75 Cu, 0.5 Mo and 20 B,} \) was applied to deliver essential nutrients.

Harvesting (twice a week) started in January and lasted until June. Harvested fruits were weighted and separated into ‘extra’ and ‘B’ categories. Fruits of ‘B’ category were characterised by malformation, various defects, or weight less than 15 g. Fruits for quality analysis were harvested at the optimum of fruit maturity at three harvesting periods (January, March and May) and results are the mean values, unless noted otherwise. The fruits of each replicate were weighted and flesh firmness was determined. Then calyces were removed, fruits were frozen, placed in polyethylene bags and stored at –30°C.

2.3. Photosynthetic parameters

The following measurements were performed on the youngest fully expanded leaf, 2-h after the sunrise in six plants per replication at monthly intervals and results are the mean values i) net assimilation rate \((A)\), stomatal conductance \((g_s)\), intercellular CO\(_2\) concentration \((C_i)\) and transpiration \((E)\) were measured using Li-6400 (Li-Cor, Lincoln, NE-USA) according to the equations derived by von Caemmerer and Farquhar [27].

2.4. Analyses

Soluble Solids concentration \((SS)\) was determined in the homogenized sample using a refractometer (Atago PR-1, Tokyo, Japan). The titratable acidity (TA) was measured by mixing 10 g of the pulp and 124 ml distilled water and titrating with 0.1 N NaOH \((\text{pH} 8.1)\), and expressed as % citric acid. The ratio of SS/TA was calculated. For the determination of dry matter \((DM)\), 10 g of berry pulp were dried at 60°C for 48 h.

Ascorbic acid content was determined by using Reflectoquant ascorbic acid test strips in an RQflex reflectometer (Merck, Darmstadt, Germany). Results were expressed as mg ascorbic acid \((\text{AA})\) per 100 g FW. Anthocyanin content was determined according to Cordemus et al. [5]. Anthocyanin content was estimated as mg pelargonidin-3-glucoside equivalents per 100 g FW using a molar absorptivity coefficient of 36,000. For total phenolics, flavonoids and ferric reducing antioxidant power (FRAP assay) samples \((5 \text{ g})\) were extracted with a mixture containing acetone, water and acetic acid \((70:29.5:0.5, v:v:v)\) [10]. The content of total phenolics was measured according to Scalbert et al. [24]
Table 1

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Extra fruits (g/plant)</th>
<th>Total yield (g/plant)</th>
<th>Berry mass (g/fruit)</th>
<th>WUE** (l/kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Cultivar</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Camarosa</td>
<td>224b</td>
<td>422</td>
<td>11.0c</td>
<td>48.5a</td>
</tr>
<tr>
<td>Festival</td>
<td>306a</td>
<td>447</td>
<td>13.6b</td>
<td>43.9ab</td>
</tr>
<tr>
<td>Ventana</td>
<td>323a</td>
<td>449</td>
<td>14.7a</td>
<td>41.4b</td>
</tr>
<tr>
<td><strong>Substrate</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BVB</td>
<td>290</td>
<td>450</td>
<td>12.9</td>
<td>41.4</td>
</tr>
<tr>
<td>Pelemix</td>
<td>289</td>
<td>440</td>
<td>13.4</td>
<td>45.7</td>
</tr>
<tr>
<td>WS</td>
<td>275</td>
<td>428</td>
<td>13.0</td>
<td>46.6</td>
</tr>
<tr>
<td>C × S ns</td>
<td>na</td>
<td>na</td>
<td>ns</td>
<td>ns</td>
</tr>
</tbody>
</table>

Means within a column followed by different letters are significantly different according to Duncan’s multiple range test at *P* < 0.05 level. *nonsignificant.

**amount of water consumed to produce one kg of fruit fresh weight.

using the Folin-Ciocalteu’s phenol reagent. Gallic acid was used as standard and the results expressed as milligrams of gallic acid equivalent (GAE) per g FW. For total flavonoids assay a protocol previously described by Kim et al. [14] was employed. Total flavonoid content was calculated from a calibration curve using catechin as the standard, and expressed as mg catechin equivalents per g FW. FRAP was determined using the method of Benzie and Strain [2]. Ascorbic acid (AA) was used as standard and the results were expressed as µmol AA per g FW.

2.5. Experimental design and statistical analysis

Experiment was arranged in a randomized block design with factorial arrangement of treatments and four replications. Each plot consisted of one 2 levels pyramid with a density of 20.6 plants m⁻². Data were analyzed using the means procedure of SAS/ASSIST.

3. Results and discussion

The total yield obtained from ‘Ventana’, ‘Festival’ and ‘Camarosa’ was similar (Table 1) and corresponded to 9.3, 9.2 and 8.8 kg m⁻² respectively. However, ‘Ventana’ and ‘Festival’ produced higher amount of ‘extra’ category fruits (Table 1) and increased berry mass compared to ‘Camarosa’ (Table 1). Total fruit yield, ‘extra’ fruits’ and berry mass were not affected by substrates and across cultivars averaged 439 g plant⁻¹, 284 g plant⁻¹ and 13.1 g fruit⁻¹, respectively. Fruit yield obtained in the current experiment when expressed per unit surface area, was within the range of earlier reports [17]. Cultivar and substrate selection for protected strawberry cultivation should be location specific and may depend on the desired period for peak production [1, 22]. In our study, harvesting started in January and lasted until June (Fig. 1). The peak of the production lasted approximately 30 days between 15 March and 15 April (Fig. 1) and this period coincides with high values in local market. Early yield (Jan-April), as a percentage of the total production, was higher in ‘Festival’ (82.9%) compared with ‘Camarosa’ (73.3%) and ‘Ventana’ was in between (77.5%)(Fig. 2). These results coincide with the notion that adapted strawberry cultivars must offer growers greater early fruit production when strawberries are not readily available in the market and carry a high price. No differences were observed on early yield results according to the substrates (data not shown).

Water consumption of ‘Ventana’, ‘Festival’ and ‘Camarosa’ strawberry plants corresponded to 366, 385 and 405 l m⁻², respectively (data not shown). Accordingly, WUE was lower in ‘Ventana’ followed by ‘Festival’ and ‘Camarosa’ (Table 1). Substrates did not affect WUE (Table 1), although plant water consumption showed a decline with BVB substrate (369 l m⁻²) compared with Pelemix and Wonder Soil, that gave similar values (avg. 395 l m⁻²).
Statistical analysis determined that there were no effects due to cultivar or substrate on photosynthetic parameters. Net assimilation rate (A) averaged 16.1 μmol CO₂ m⁻² s⁻¹; stomatal conductance (gs), averaged 0.332 mol H₂O m⁻² s⁻¹; intercellular CO₂ concentration (Ci) averaged 279 μmol CO₂ mol⁻¹ and transpiration (E) averaged 3.39 mmol
Table 2

The effect of strawberry cultivar and substrate on dry matter (DM), pH, soluble solids (SS, °Brix), titratable acidity (TA, % citric acid), ascorbic acid (mg AA (100 g FW)−1), total anthocyanins (mg pelargonidin-3-glucoside (100 g FW)−1), total flavonoids (mg catechin (g FW)−1), total phenolics (mg Gallic Acid (g FW)−1) and antioxidant capacity (FRAP values [µ mol AA (g FW)−1]) of strawberry fruits.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>DM</th>
<th>pH</th>
<th>SS</th>
<th>TA</th>
<th>SS/TA</th>
<th>Asc. acid</th>
<th>Total anth.</th>
<th>Total flavon.</th>
<th>Total phen.</th>
<th>FRAP values</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Cultivar</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Camarosa</td>
<td>7.4</td>
<td>3.6</td>
<td>6.9a</td>
<td>0.92a</td>
<td>7.7b</td>
<td>35a</td>
<td>30a</td>
<td>0.69a</td>
<td>2.5a</td>
<td>20a</td>
</tr>
<tr>
<td>Festival</td>
<td>7.3</td>
<td>3.7</td>
<td>6.8a</td>
<td>0.72b</td>
<td>9.5a</td>
<td>36a</td>
<td>23b</td>
<td>0.47c</td>
<td>2.2b</td>
<td>16b</td>
</tr>
<tr>
<td>Ventana</td>
<td>7.1</td>
<td>3.7</td>
<td>6.3b</td>
<td>0.69b</td>
<td>9.2a</td>
<td>31b</td>
<td>24b</td>
<td>0.56b</td>
<td>2.1b</td>
<td>16b</td>
</tr>
<tr>
<td><strong>Substrate</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BVB</td>
<td>7.1</td>
<td>3.7</td>
<td>6.7</td>
<td>0.77</td>
<td>8.9</td>
<td>35</td>
<td>26</td>
<td>0.56</td>
<td>2.2</td>
<td>17</td>
</tr>
<tr>
<td>Pellemix</td>
<td>7.3</td>
<td>3.6</td>
<td>6.7</td>
<td>0.78</td>
<td>8.9</td>
<td>33</td>
<td>26</td>
<td>0.59</td>
<td>2.3</td>
<td>18</td>
</tr>
<tr>
<td>WS</td>
<td>7.2</td>
<td>3.6</td>
<td>6.7</td>
<td>0.78</td>
<td>8.7</td>
<td>34</td>
<td>25</td>
<td>0.56</td>
<td>2.2</td>
<td>17</td>
</tr>
<tr>
<td>C × S ns*</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
<td>ns</td>
</tr>
</tbody>
</table>

Means within a column followed by different letters are significantly different according to Duncan’s multiple range test at P<0.05 level. *nonsignificant.

Table 3

The overall effect of the harvesting period on dry matter (DM), pH, soluble solids (SS, °Brix), titratable acidity (TA, % citric acid), ascorbic acid (mg AA (100 g FW)−1), total anthocyanins (mg pelargonidin-3-glucoside (100 g FW)−1), total flavonoids (mg catechin (g FW)−1), total phenolics (mg Gallic Acid (g FW)−1) and antioxidant capacity (FRAP values [µ mol AA (g FW)−1]) of strawberry fruits.

<table>
<thead>
<tr>
<th>Harvest period</th>
<th>DM</th>
<th>pH</th>
<th>SS</th>
<th>TA</th>
<th>SS/TA</th>
<th>Asc. acid</th>
<th>Total anth.</th>
<th>Total flavon.</th>
<th>Total phen.</th>
<th>FRAP values</th>
</tr>
</thead>
<tbody>
<tr>
<td>January</td>
<td>7.4a</td>
<td>3.7</td>
<td>7.4a</td>
<td>0.71b</td>
<td>10.5a</td>
<td>34a</td>
<td>21c</td>
<td>0.47b</td>
<td>1.7c</td>
<td>14c</td>
</tr>
<tr>
<td>March</td>
<td>6.6b</td>
<td>3.6</td>
<td>6.1c</td>
<td>0.74b</td>
<td>8.3b</td>
<td>32b</td>
<td>25b</td>
<td>0.51b</td>
<td>1.3b</td>
<td>17b</td>
</tr>
<tr>
<td>May</td>
<td>7.5a</td>
<td>3.6</td>
<td>6.5b</td>
<td>0.87a</td>
<td>7.7b</td>
<td>35a</td>
<td>31a</td>
<td>0.74a</td>
<td>2.9a</td>
<td>22c</td>
</tr>
</tbody>
</table>

H₂O m⁻² s⁻¹. Net assimilation rate corresponded well to earlier reports [4] and it was consistent with the respective yield results.

Regarding fruit quality, ‘Festival’ and ‘Ventana’ were interesting for higher SS/TA ratio compared to ‘Camarosa’. This may suggest better taste since taste quality usually depends on the ratio of soluble solids to titratable acidity. However, all cultivars showed SS/TA ratio adequate to achieve best quality [5, 23] (Table 2). The phenolic compounds usually found in strawberry are phenolic acids, flavonoids and anthocyanins [15]. Dissimilar to SS/TA ratio, ‘Camarosa’ fruits performed higher values of anthocyanins, flavonoids and phenolics which resulted to a higher antioxidant capacity as measured by FRAP assay (Table 2). Moreover, higher anthocyanin content may indicate a more attractive color given that anthocyanins account for the color in strawberry fruits [8]. These values corresponded to earlier reports [12, 23] and any minor discrepancies may be attributed to various factors such as cultivar effect, climatic conditions and experimental set up. No differences in fruit quality were observed according to the type of coco-substrate (Table 2). These results revealed that differences in the antioxidant capacity exist in strawberry fruit, which depend on genotype [13], and may provide both growers and consumers with health benefit information.

Correlations coefficients for different antioxidant compounds and antioxidant capacity were calculated. A significant linear correlation (Pearson) was observed between antioxidant capacity (FRAP) and anthocyanins, flavonoids and total phenolics (r = 0.71***, r = 0.89***, r = 0.86***, respectively). In line with our results, a considerable body of data suggests that a higher content of total phenolics, flavonoids and anthocyanins in strawberry enhances their antioxidant activity [11, 15, 28]. Inversely, ascorbic acid was not correlated with antioxidant capacity. Torrønen and Määtä [26] mentioned only a small contribution of ascorbic acid to total antioxidant capacity compared to total phenolics and flavonoids. To summarize, quality was mainly related to genotype and not to the type of coco-substrate. This confirms the importance of having soilless-adapted varieties for greenhouse production with improved and stable fruit quality [6].
Finally the impact of the harvesting period on fruit quality parameters was studied. Harvesting from January to May improved studied bioactive compounds and antioxidant activity of fruits, but taste quality related SS/TA ratio was negatively affected (Table 3). These variations were probably influenced by temperature, irradiance and leaf:fruit ratio, which may affect the formation and accumulation of carbon-based metabolites, thus, the quality parameters of strawberry fruits [6, 16]. Seasonal variation in antioxidants compounds observed may influence consumers’ preference in case of a diet rich in antioxidants.

4. Conclusions

‘Festival’ and ‘Ventana’ appear to be interesting alternative cultivars. These cultivars attained early fruit production with large fruits of good taste quality and good efficiency in water management. However, ‘Camarosa’ produced fruits with higher antioxidant compounds and a more attractive color. The type of coco-substrate was not a limiting factor. The use of an optimum time for planting, the cultivars and plant support media used along the management of the nutrient solution, would allow lengthening of the fruiting season and particularly greater yields in December and January of high quality fruits and has to be investigated.

References