
The Effect of Crop Load on Tree Water Consumption of ‘Golden Delicious’ Apples in Relation to Fruit Size: An Operative Model

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ABSTRACT. Water availability and crop load (number of fruit per tree) affect the fruit size of apple (Malus xdomestica Borkh.), but their interaction in relation to fruit size is not well understood. The objective of the present study was to explore the effect of crop load on water consumption of ‘Golden Delicious’ apple in relation to fruit size. A wide range of irrigation rates and crop loads was applied to mature, field-grown ‘Golden Delicious’ apple trees for two consecutive years, 1995 and 1996. The number of fruit, crop yield, and average fruit diameter were determined for each tree. A model was proposed to describe the combined effect of crop yield and irrigation rate on fruit size. In the model, irrigation waters were divided between two uses: vegetative water use ($U_v$), which enables the tree to produce a steady, long-term yield; and reproductive water use ($U_r$), which supports the production of the dry mass of commercial-size fruit. Potential fruit diameters were 77.1 and 72.2 mm for 1995 and 1996, respectively. Calculated vegetative use values were 300.2 and 323.4 mm for 1995 and 1996, respectively. The response of fruit diameter to reproductive water use per ton fresh weight (specific reproductive water use; $U_{fr}$) was fitted by a hyperbolic model in which the fruit diameter increases with increasing specific reproductive water use and approaches the yearly potential diameter at 60 to 70 m²·t⁻¹, irrespective of the potential fruit diameter. In both years, fruit diameter showed a closer correlation with the specific reproductive water use than with either crop load or irrigation rate. In conclusion, the crop yield and the potential fruit size determine the irrigation rate required to achieve a certain average fruit diameter. The year’s potential fruit diameter does not affect the total tree water use or its components. The proposed model can be used by growers for supporting decisions on irrigation and thinning strategies in commercial orchards.

Fruit size is a major aspect of apple quality. Since fruit size is affected by water deficit, and since water is a limiting factor to fruit production in some apple-producing regions, both economically and physically, optimizing water application is increasingly important.

Fruit size is negatively related to crop load (number of fruit/tree) in apple (Assaf et al., 1982; Erf and Proctor, 1987; Naor et al., 1997a, 1997b) and in other deciduous fruit trees (DeJong and Grossman, 1995; Naor et al., 1999, 2000, 2004). Limited availability of assimilates has been suggested as the reason for obtaining smaller fruit at high crop loads in peach (Prunus persica L.; DeJong and Grossman, 1995) and in apple (Wibbe et al., 1993). The number of cells in the pericarp may limit potential apple fruit size, and this number depends on the temperature during the cell-division phase (Austin et al., 1998; Bergh, 1990; Flaisalman and Naor, unpublished; Warrington et al., 1999). Fruit size also depends on crop loads in the current and the previous seasons (Bergh, 1985).

Apple yield and fruit-size distribution are affected by the irrigation rate (Assaf et al., 1982; Erf and Proctor, 1987; Naor et al., 1997a); fruit was smaller for a high crop load than for a lower crop load when subjected to “deficit irrigation” (Mpelasoka et al., 2001). Stomatal conductance increases (Erf and Proctor, 1987; Hansen, 1971) and midday stem water potential decreases (Naor et al., 1997b) as the crop load of stressed apples increases. These responses provide us with qualitative indications of the effect of crop load on tree water consumption. Compared with nonfruiting apple trees, fruiting trees had higher water consumption in containerized (Lenz, 1986) and lysimeter-grown trees (Mpelasoka et al., 2001). Lenz (1986) directly demonstrated that the tree water consumption of fruiting trees was 25% to 50% greater than that of nonfruiting ones in a controlled environment. Nevertheless, there is a lack of quantitative information on the effect of crop load on tree water consumption in field-grown apple.

Optimal irrigation is expected to maintain a well-balanced tree structure (roots, limbs, shoots, leaves, and flower buds) that can provide the fruit with enough assimilates at an adequate rate for them to reach commercial fruit size, and to ensure enough fertile-flower bud formation to enable steady commercial crop yield production. A balanced tree structure can be achieved provided vegetative growth is adequate for the formation of a sufficient number of flower buds and for the postharvest accumulation of assimilates in the tree-carbohydrate pool. The fulfillment of both these conditions would enable persistent cropping. Ebel (1991) showed that apple trees grown in a semiarid zone could survive one unirrigated season, provided that there was adequate moisture storage in the soil profile. Nevertheless, the canopy size of the unirrigated trees decreased to 40% in the subsequent year, and the number of fruit/tree was reduced to 55% of the number on the well-irrigated trees. This demonstrates that the tree water consumption required to maintain a well-balanced tree structure is considerably greater than the survival water consumption. There is a need to quantitatively evaluate the amount of irrigation required to maintain a well-balanced tree structure, and the additional water consumption necessitated by the presence of a crop.

The objective of the present study was to establish a model for describing the combined effect of crop yield and irrigation rate on the fruit size of field-grown ‘Golden Delicious’ apple.
Materials and Methods

**Model Description.** It is assumed that the response of crop yield to soil moisture availability at a given crop load (number of flowering buds) has a single sigmoidal pattern (Fig. 1). There is a minimum soil moisture availability (A) below which trees might dry out and die; A–B represents a section where the proportion of fruit set and the proportion of surviving fruitlets increase with increasing water use (Powell, 1974) as a result of either improved turgor potential or of improved assimilate availability to the fruit that may prevent fruit drop. When water use is in the A–B range, a reduction in the structure dry biomass may occur, mainly by defoliation and by a negative balance in root turnover (Ebel, 1991; Lakso et al., 1999; Lenz, 1986). Nevertheless, trees will survive; B–C represents a section where both assimilate availability for fruit expansive growth and potential fruit size limit crop yield. When water availability is in the B–C section, trees may retain their structure. Shoot growth would increase with increasing water use to retain cropping potential until a certain point (undefined) from which additional water use might induce excessive shading within the canopy. Tree water use in commercial orchards is usually in section B–C, and reduced crop load in field-grown apple was reported only under nonirrigation (Ebel, 1991) or deficit irrigation in the cell division phase (Naor et al., unpublished; Powell, 1974). Therefore, the following analysis is restricted to high water use values (B–C) (Fig. 1) where both assimilate availability and potential fruit size limit crop yield and fruit size.

In the following model, irrigation water is divided into two components. The first is the water use for vegetative growth in addition to available soil moisture at budbreak (U\textsubscript{b}), which is the water use below which fruit drop would significantly limit crop yield. The second component is the reproductive water use (U\textsubscript{r}), which represents the water use designated to support the production of the dry mass of commercial-size fruit.

The degree to which the potential fruit size is achieved depends on the specific reproductive water use (reproductive water use per 1 metric ton of fruit; U\textsubscript{r}), which affects the ability of the canopy to supply the crop with assimilates at an adequate rate (DeJong and Grossman, 1995).

\[ U_{rr} = \frac{(I_{tot} - U_{b})}{Y_{tot}} \quad \text{Eq. 1} \]

where \( U_{rr} \) = irrigation per ton of fruit (m\textsuperscript{3}·t\textsuperscript{-1}); \( I_{tot} \) = total annual irrigation (m\textsuperscript{3}·ha\textsuperscript{-1}); \( U_{b} \) = annual amount of water designated for vegetative growth in addition to the available soil moisture at budbreak (m\textsuperscript{3}·ha\textsuperscript{-1}); and \( Y_{tot} \) = total crop yield (t·ha\textsuperscript{-1}).

In an “efficient” apple orchard the irrigation efficiency is assumed to be 100%, and the availability of assimilates is not a limiting factor; i.e., the crop load is in the range where the yield response to crop load is linear (DeJong and Grossman, 1995) (Fig. 2). The response of fruit diameter to the specific reproductive water use in an “efficient” apple orchard follows a hyperbolic curve:

\[ D = \frac{D_{max}(I_{tot} - U_{r})Y_{tot}}{(I_{tot} - U_{b})Y_{tot} + b} \quad \text{Eq. 2} \]

where \( D_{max} \) = fruit diameter, potential fruit diameter (mm); and \( b = \) slope constant.

**Feasibility Test.** Data from an old irrigation experiment were used for the feasibility test. The experimental settings have been reported in detail elsewhere (Naor et al., 1997a, 1997b). The experimental plot consisted of a 12-year-old drip-irrigated commercial orchard of ‘Golden Delicious’ apple on ‘Malling-Merton 106’ rootstock spaced at 4 m between rows × 2 m within row. The trees formed a full hedgerow, about 3 m in height with a relatively uniform canopy structure. The experimental plot was irrigated until mid-June according to commercial practice: irrigation started about 1 May at a Class A pan evaporation coefficient of 0.30 and was systematically increased, in coordination with the increase in canopy size and in fruit diameter, to reach 0.45 in mid-June. Differential irrigation treatments were started in mid-June, at the beginning of the main cell expansion phase in the fruit mesocarp. An additional 90 mm of water was applied to all treatments over 6 weeks from the end of harvest until the beginning of the rainy season in late October. The experimental design enabled the study of the interaction of irrigation and crop load during the main period of fruit growth. The experimental design was a split-plot factorial, with five irrigation rates as main plots and
crop loads as subplots. The treatments were replicated five times in a complete randomized block design. Each main plot consisted of four adjacent rows of four trees, and only the four inner trees of each plot were used for the four crop load treatments. The irrigation rates applied were Class A pan evaporation coefficients of 0.42, 0.58, 0.75, 0.90, and 1.06 in 1995, and 0.42, 0.58, 0.71, 0.87, and 1.03 in 1996. The fruit on each tree were counted at the beginning of June and were hand-thinned to four crop load levels: 0 to 150, 151 to 250, 251 to 350, and >350 fruit/tree. After harvest the fruit of each tree were weighed, and the diameter distribution was determined by means of a commercial grading machine (55 to 75 mm). The experiment was repeated with the same trees for two consecutive seasons, 1995 and 1996. Crop loads were switched between trees where necessary.

**Statistical Analysis.** Average fruit diameter, total crop yield for each tree, and total annual irrigation rates were used to evaluate the parameters of the model (Eq. 2) by means of nonlinear regression analysis (SigmaPlot, V 7.0; SPSS Inc., Chicago).

**Results**

The annual irrigation rates were 446, 536, 632, 723, and 817 mm in 1995, and 422, 524, 600, 702, and 804 mm in 1996. The crop loads ranged from 50 to 510 fruit/tree in 1995 and from 55 to 640 fruit/tree in 1996. Yields were linearly related to crop loads (Fig. 2) and ranged from 12 to 114 t·ha−1 in 1995 and from 10 to 118 t·ha−1 in 1996. Crop yields were higher in 1995 than in 1996 for each crop load (Fig. 2). The wide ranges of irrigation rates and crop loads allowed the investigation of wide ranges of specific reproductive use (Eq. 1): from 6 to 320 m³·t−1 in 1995 and from 4 to 240 m³·t−1 in 1996 (data not shown). Potential fruit diameters differed between years and were 77.1 and 72.2 mm in 1995 and 1996, respectively (Table 1; Fig. 3) Vegetative water use values were 300.2 and 323.4 mm for 1995 and 1996, respectively (Table 1). The relative fruit diameter (average diameter expressed as a fraction of the year’s potential fruit diameter) responded similarly to specific reproductive water use in both years, increasing with specific reproductive use and approaching 1 at 60 to 70 m³·t−1 (Fig. 4).

Fruit diameter was better correlated with crop load (number fruit/tree) than with annual irrigation rate in 1995 (Table 2) whereas the opposite was apparent in 1996. In both years fruit diameter had the highest correlation with specific reproductive use, a parameter that takes into account the effects of both the irrigation rate and the crop yield.

A simulation based on the combined data of 1995 and 1996 (Fig. 4; Table 1) shows that the reduction in fruit diameter in response to crop yield becomes more pronounced with decreasing annual irrigation rate (Fig. 5). A reduction of annual irrigation rate from 1000 to 700 mm would result in a decrease in fruit diameter of

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Table 1. The parameters of the model (Eq. 2) for 1995, 1996, and for the merged data with fruit diameter relative to its maximum value each year. The parameters were significant at α < 0.0001.

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<tr>
<td>Potential fruit diam (mm) ± se</td>
<td>77.1 ± 0.42</td>
<td>72.2 ± 0.55</td>
<td>100% ± 0.47</td>
</tr>
<tr>
<td>Vegetative water use (mm) ± se</td>
<td>300.2 ± 39.8</td>
<td>323.4 ± 30.2</td>
<td>320.5 ± 22.4</td>
</tr>
<tr>
<td>r²</td>
<td>0.58</td>
<td>0.56</td>
<td>0.58</td>
</tr>
<tr>
<td>n</td>
<td>79</td>
<td>92</td>
<td>171</td>
</tr>
</tbody>
</table>

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Table 2. Correlation coefficients (r²) between fruit diameter at harvest and total irrigation, number of fruit/tree, and specific reproductive water use (Eq. 1).

<table>
<thead>
<tr>
<th></th>
<th>1995</th>
<th>1996</th>
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<tbody>
<tr>
<td>Total annual irrigation</td>
<td>0.262</td>
<td>0.431</td>
</tr>
<tr>
<td>Crop load (fruit/tree)</td>
<td>0.354</td>
<td>0.268</td>
</tr>
<tr>
<td>Specific reproductive water use</td>
<td>0.539</td>
<td>0.570</td>
</tr>
<tr>
<td>n</td>
<td>79</td>
<td>92</td>
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size in 1995. The ful
crop load in 1995 than in 1996, indicating a larger potential fruit
diameter between years are large at low crop yield and
0 to 2 mm, while a reduction of irrigation rate from 700 to 400
mm would result in a dramatic decline in fruit diameter, namely
7 mm at 400 fruit/tree and 10 mm at 700 fruit/tree. The effects
crop yield on fruit diameter at three crop loads, 100, 400, and
700 fruit/tree, are presented in solid lines (Fig. 5). Differences
in fruit diameter between years are large at low crop yield and
high annual irrigation rate, whereas they decrease at high crop
yield and low annual irrigation rate.

Discussion

In the present study, higher crop yields were apparent for every
crop load in 1995 than in 1996, indicating a larger potential fruit
size in 1995. The fulfillment of the differences in potential fruit
size occurred at high irrigation rates and low crop yields (Fig.
5) when availability of assimilates did not limit fruit growth.
On the other hand, at low irrigation rates and high crop yields,
where assimilate availability limits fruit expansive growth, the
difference in fruit size between years decreased.

The cost in terms of water use for the production of 1 t (fresh
weight) of apple fruit was estimated as follows. A typical water
use efficiency ratio of C3 plants on a mass basis is 1:205 (Taiz
and Zeiger, 1991); typical dry matter content for ‘Golden Deli-
cious’ apple fruit is 16% (Lakso et al., 1999). There would be
an additional small cost to account for maintenance and growth
respiration of the fruit (Proctor et al., 1976) as well as a more
substantial metabolic cost for processes involved in translocation
of assimilates from the leaves into the fruit (Wibble et al., 1993).
On the assumption of a respiratory cost equivalent to 30% of
the total carbon fixed by the canopy (Walton and Fowke, 1995),
the overall cost in terms of water consumption for the production of
1 t (fresh weight) of apples would be ~47 m³. It should be noted
that this value is only a rough estimate and may change from one
region to another because of climatic differences, which affect both
respiration and water use efficiency on a leaf basis (WUEleaf).

Applying specific reproductive water use amounts lower than
47 m³·t⁻¹ might cause a shift in assimilate translocation in favor
of the fruit, which would harm tree structure in the long term,
because of attenuation of shoot growth (Ebel, 1991). Applying
specific reproductive use levels higher than 47 m³·t⁻¹ would
increase the proportion of assimilates transported to the tree
structure, which could be manifested in excessive shoot growth.
Consistent application of specific reproductive use levels either
higher or lower than 47 m³·t⁻¹ for a few years is expected to result
in the adjustment of the tree structure so as to change vegetative
water use until the actual specific reproductive use approaches
47 m³·t⁻¹ in a new steady-state condition. The similar response
of fruit diameter to specific reproductive use in both years (Fig.
5) suggests that the damage to the structure in 1995 at specific
reproductive use lower than 47 m³·t⁻¹ was nonsignificant over a
2-year period.

The vegetative water use depends on the demand for mainte-
nance respiration of the leaves, roots, and stem (Grossman and
De Jong, 1994), and on the evaporative demand, which determines
transpiration rates throughout the growing season. Since respira-
tion exhibits an exponential response to temperature (Grossman
and De Jong, 1995; Proctor et al., 1976), weather conditions
during fruit growth may have an impact both on dark respiration
and on net photosynthesis during the day, so that the weather is a
major factor in determining the water consumption of orchards.
Vegetative use is positively related to the canopy biomass per unit
area of orchard (i.e., hedgerow height) as it dictates the amount
of light intercepted by the canopy (Lakso et al., 1999) and the
total maintenance respiration of the tree (Grossman and De Jong,
1994). Canopy size affects the yield capacity of the tree and the
crop yield at which assimilate availability becomes a limiting
factor; therefore, a high vegetative use does not necessarily imply
that the trees are inefficient.

The low response of fruit diameter at high specific reproduc-
tive water use levels (Fig. 4) demonstrates a carbon partitioning
pattern that favors fruit over nonreproductive tissue, which is
consistent with previous findings (e.g., De Jong and Grossman,
1995). Therefore, if the irrigation supply falls short of meeting the
added water requirements of the canopy and of the crop, it will
necessarily result both in a smaller crop yield in the current year
through a reduction in fruit size, and in inhibited canopy growth,
which might be accompanied by a decrease in the reserves of
assimilate stored in the stem (Lakso et al., 1999) and thus limit
canopy development and fruit set early in the following year
(Ebel, 1991; Grossman and De Jong, 1995). On the other hand, if
irrigation rates were above the requirements of the fruit and the
balanced tree structure, it would result in extensive shoot growth,
which may require additional labor for summer pruning.

Water use efficiency at the orchard level (WUEorchard; fresh
weight production/applied irrigation) is the reciprocal of the
specific reproductive water use, indicating that WUEorchard in-
creases with decreasing specific reproductive use. A decrease in
specific reproductive use can be obtained by decreasing annual
irrigation rate or increasing crop yield (Eq. 1), and the tree may
respond differently to each factor. When specific reproductive
use decreases in response to an increase in crop yield (higher
crop load) the whole-tree assimilation rate would be increased
because of the increased stomatal aperture (Erf and Proctor, 1987;
Hansen, 1971). However, the assimilation rate can increase by
no more than ~15% because of increased stomatal conductance
(De Jong 1986), whereas in the current study there was a 10-fold

Fig. 5. A simulation of the effect of crop yield on fruit diameter at various annual
irrigation rates in 1995 (dashed lines) and 1996 (dotted lines) and the effect of
crop yield on fruit diameter at 100, 400, and 700 fruit/tree (solid line). Underlined
numbers are annual irrigation rates (mm). The simulation was restricted to
irrigation rates and crop loads within the actual experimental ranges.

difference in crop yields between the lowest and highest crop loads. If specific reproductive use decreased because of a reduction in annual irrigation rate, it would result in a deterioration of tree water status, which would be manifested in a reduction in stomatal conductance and assimilation rate. This reduction, in turn, would lead to an increase in WUE\textsubscript{c} (assimilation rate/transpiration rate), because the assimilation resistance is always higher than the transpiration resistance due to additional nonstomatal resistance (Jones, 1992); nevertheless, the tree assimilation rate would decrease. Both annual irrigation rate and crop yield affect the rate of the transport of assimilates into the fruit by either changing the supply of assimilates or changing sink capacity. However, severe water stress may attenuate expansive growth by a low turgor potential and a decrease in the rate transport of assimilates in the phloem (Nobel, 1991). It seems, therefore, that the water status in the current study was not low enough to limit expansive growth by a low turgor potential and a decrease in the rate transport of assimilates in the phloem.

The link found in the present study between crop yield and tree annual irrigation rate could potentially be used for planning irrigation and thinning regimes in commercial orchards, in order to achieve the optimum fruit-size distribution. The annual irrigation rate and fruit thinning level can be optimized during the growing season once sufficient good estimates of crop yield and potential fruit size are available. For example, different specific reproductive water use values would be applied in 1995 and in 1996, assuming that the commercially optimal fruit diameter is 71 mm; For 1995, when the potential fruit diameter reached 77.1 mm, a 20 m\textsuperscript{3}t\textsuperscript{–1} specific reproductive use would be sufficient to achieve the desired fruit diameter, whereas in 1996, when the potential fruit diameter was 72.2 mm, the specific reproductive use required to reach the same mean diameter would be substantially higher, at 58 m\textsuperscript{3}t\textsuperscript{–1}.

The availability of irrigation water would dictate the optimal crop load for each potential fruit size. For example: should water supply be restricted to 550 mm during the 2 years of the present study, the total reproductive water use for both years would be 230 mm. For an optimal fruit diameter of 71 mm, in 1995 this quantity of water would suffice for a crop yield of more than 110 t ha\textsuperscript{–1} commercially optimal-size fruit, whereas in 1996 thinning should probably be applied, to reduce the crop yield to some 40 t ha\textsuperscript{–1} of similarly sized fruit. It should be noted, however, that reliable technologies for the early-stage assessments of crop yield and of potential fruit size are yet to be developed to enable the benefits from the utilization of the model in commercial orchards to be maximized.

After harvest, when the actual crop yield and fruit size distribution data for each commercial plot are available, the following analysis can be performed. If the data for a specific commercial plot lie on the regression line (Fig. 4), it suggests that the trees were efficient and that whether the irrigation was suboptimal, optional, or excessive depended on the optimal fruit diameter. If the data for a specific plot lie below the regression line, the trees are considered to be inefficient, which could be attributed to low irrigation efficiency, to an excessive crop load, or to limitation of availability of assimilates attributed to the activities of pests and/or diseases.

**Literature Cited**


