Stomatal Function and Cuticular Conductance in Whole Tissue-cultured Apple Shoots

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Additional index words. water transport, water stress, gas exchange, Malus pumila, rootstock M.26

Abstract. The relative contribution of stomatal and cuticular conductance to transpiration from whole tissue-cultured apple shoots of Malus pumila Mill. M.26 was determined with a modified steady state porometer. When shoots were exposed to 90% RH and high boundary layer conductance, large (73%) and, in some cases, rapid (2 to 3 hours) reductions in leaf conductance occurred, indicating functional stomata. Stomatal closure was also observed microscopically. A maximum estimate for the cuticular conductance of these apple leaves was 18 to 40 mmol·m$^{-2}$·s$^{-1}$, which is lower than previous estimates and close to the upper limit of naturally occurring leaf cuticular conductances. Hence, both stomatal and cuticular restrictions of water loss appear to be of importance in determining the water balance of tissue-cultured apple shoots. The pathway of water transport in relation to water stress of tissue-cultured shoots is also discussed.

Lack of stomatal closure in tissue-cultured shoots has been implicated as a major cause of the desiccation that occurs when shoots are removed from in vitro conditions and placed in an unprotected environment (Brainerd and Fuchigami, 1982). Poor functioning of stomata has been attributed to abnormal microfibril development of guard cells (Wardle and Short, 1983) and to poor selectivity in the accumulation of Na, K, and Mg in guard cells while in vitro (Wardle et al., 1981).

The relative contribution of stomatal and cuticular water loss and the capacity of the stomata for normal functioning in cultured shoots remain controversial. Most of the data supporting the hypothesis that stomata in cultured shoots are nonfunctional is based on experiments with excised leaves (Brainerd et al., 1981; Fuchigami et al., 1981; Sutter and Langhans, 1982). Such experimental systems may not accurately portray conditions in vivo. Sutter (1988), using intact shoots, reported results in conflict with those discussed above. She found that stomatal conductance of intact tissue-cultured apple, cherry, and sweetgum shoots decreased when shoots were removed from in vitro conditions, but also reported rapid (within 1 to 2 min) decreases in cuticular conductance. In some cases, the magnitude of the changes in stomatal and cuticular conductance were similar; hence, the separation of stomatal from cuticular function remains unclear. Another difficulty with Sutter's study was that conductance was determined at laboratory relative humidity (37% to 45%), which was presumably much lower than the ambient humidity under tissue culture conditions.

The objectives of this study were to determine whether stomata of intact tissue-cultured apple shoots could close in response to a moderate increase in evaporative demand in situ and to estimate the relative contribution of stomatal and cuticular conductances to the regulation of water loss from these shoots. This paper also reports the development of a porometer-based gas exchange system that is suitable for the study of stomatal functioning in tissue-cultured shoots.

Materials and Methods

Gas exchange system. Measurement of the gas exchange of tissue-cultured shoots was made with a modified steady state porometer (model 1600; LI-COR, Lincoln, Neb.) in a controlled-temperature room (24 ± 1C, continuous fluorescent light of 30 to 35 µmol·s$^{-1}$·m$^{-2}$). The porometer was modified by replacing the leaf clamp and aperture cap with a glass adapter and by extending the leaf thermocouple wires (Fig. 1). A laptop computer (model PC-8201A; NEC, Tokyo) was programmed to operate the porometer and to collect, average, and store mea-
measurements for user selectable periods of time. Accuracy of the humidity sensor was verified at about weekly intervals by equilibration with a saturated NaCl solution of 75.1% RH, as recommended by the manufacturer, and was always within 3% RH. A micropipette gauge (model 1600MP; LI-COR) was used to verify the accuracy of the transpiration measurements at about biweekly intervals. The gauge was substituted for the culture vessel (Fig. 1) and petrolatum was used to seal the gauge and the glass adapter. A culture vessel filled with water was used to equilibrate the system at 90% RH for 1 to 2 hr before taking measurements. After this equilibration period, good agreement was found between the water loss rate, as measured by the porometer, and that determined directly by the rate of water movement in the gauge (Fig. 2). The data for Fig. 2 were obtained immediately after installing the gauge in the system. By increasing the quantity of water in the gauge before installation, the transpiration values could also be tested after longer time intervals (up to 5 hr) and were always accurate to within 6%. Boundary layer conductance within the glass adapter was similar to that used by the porometer for the calculation of stomatal conductance, so the conductance recorded by the porometer was only corrected for the shoot leaf area determined at the end of each experiment. Relatively small oscillations in room temperature with a 17-min period had a substantial influence on the measured values of transpiration (Fig. 2); for all experiments, the computer was programmed to collect data at 0.5-min intervals but to store only mean values for each 17-min period.

Shoot culture and general experimental procedures. Shoots of M.26 apple rootstock were grown in vitro on agar-solidified Linsmaier-Skoog nutrient medium (Linsmaier and Skoog, 1965) to, which 1 mg benzyladenine/liter and 162 mg phloroglucinol/liter were added. The cultures were incubated in a controlled-environment chamber at 25°C, under a 16-hr photoperiod and 75 μmol·s⁻¹·m⁻² provided by cool-white fluorescent lamps, and were transferred at 4-week intervals. Individual shoots, 1.5 to 2 cm tall, were excised from the proliferating mass of shoots and were placed in glass scintillation vials filled with 25 ml Linsmaier-Skoog medium. About 0.3 cm of the shoot was inserted into the medium. The vials were placed in a larger vessel and were allowed to continue to grow for an additional 4 days to 1 week. Water was placed in the larger vessel to just below the upper edge of the vial to increase ambient humidity and prevent drying of the shoot.

Tissue-cultured shoots were prepared for all gas exchange experiments in a humidified glove box; to prevent water evaporation from the nutrient medium, melted petrolatum was layered on the surface of the medium (Fig. 1) using a syringe and needle. Three protocols for shoot preparation were used. Unmodified shoots: no additional modifications were made; modified shoots: a syringe and needle were used to apply a layer of petrolatum to the abaxial surfaces of all leaves to prevent evaporation from the majority of stomata; isolated shoots: shoots were removed from the normal culture medium and inserted into a layer of petrolatum to isolate them from any supply of water. After shoot preparation, the gas exchange system was temporarily turned off, and the culture vessel was quickly substituted for the water-filled vessel that had been used to equilibrate the system at 90% RH. The system was then restarted, and the dry air flow control manually adjusted to maintain 90% ± 0.4% RH during the course of the experiment. Typically, this procedure required substantial adjustment only during the first 10 min and no additional adjustments after 1 to 2 hr.

At the completion of each experiment (typically 24 hr), the entire gas exchange system was transferred to a humidified glove box, shoots were removed, and leaf fresh weight was determined. Leaves were hydrated by placing them in a beaker of water and their leaf area was determined with a Delta T Area Meter (Decagon Instruments, Pullman, Wash.) while they were between sheets of clear plastic. Leaves were then dried in an oven for 2 days at 80°C for dry weight measurements. The dry weight and water content were also determined for a number of individual shoots taken directly from culture. Observations of abaxial and adaxial leaf surfaces on intact shoots were made at ×200 with a vertically illuminated microscope (BHMJ system; Olympus, New York) using a long working distance (11 mm) ×20 objective (1-LM546; Olympus). The shoot, vial, and microscope objective were enclosed in a loosely fitting, humidified plastic bag during the observations to prevent plant desiccation and allow flexibility in positioning the leaves.

Results and Discussion

In all experiments, unmodified shoots exhibited rapid initial reductions in leaf conductance (g1) followed by more, gradual reductions, reaching relatively stable values of g1 by the end of the 24-hr test period (Fig. 3). The three curves for unmodified shoots represent the range of variability observed in the 11 unmodified shoots tested. They are typical in that shoots exhibiting high initial values of g1 (the highest recorded was 700 mmol·m⁻²·s⁻¹) also exhibited high final values and relatively slow rates of decline during much of the test period. The variability in shoot size was considerable, ranging from leaf areas of 3.08 to 9.81 cm², but, after correcting for leaf area, there was no apparent relation between shoot size or morphology and shoot gas exchange characteristics. The substantial reduction in g1, averaging 73%, indicated that the stomata of these shoots were functional. The relatively rapid reduction in g1 that was observed in some plants further supports this conclusion.

The apparent reduction in g, over time for unmodified shoots (Fig. 3) could have been due to water contamination in the gas exchange system itself. Evaporation of water derived from the medium or of superficial water on leaf surfaces, for example, would cause overestimation of conductance initially. The rate of water loss from vials containing uncoated media was substantial. No water loss was detected, however, when the me-

![Fig. 2. Comparison between calculated (porometer measurement) and observed (pipette measurement) rates of water loss from a paper evaporation surface in the gas exchange system of Fig. 1. Porometer measurements were collected by a computer at 0.5-min intervals, and pipette measurements were taken in increments of 1 μl water loss. Room air temperature over time (hours : minutes) is also shown.](image-url)
Fig. 3. Leaf conductance over time (hours : minutes) for representative shoots of three experimental treatments (unmodified, abaxial surfaces coated, and isolated from media) in the gas exchange system of Fig. 1. Values of conductance >300 mmol·m⁻²·s⁻¹ are omitted for clarity.

Medium was completely covered by petrolatum, indicating that the petrolatum seal was effective. High rates of water loss also occurred initially from wet filter paper placed on the petrolatum, but these dropped to zero after 1 hr. Hence, even a substantial quantity of superficial water on the shoot (none was apparent) would have a negligible effect on $g_c$ after 1 hr. The apparent reduction in stomatal conductance may also have been due to the loss of tissue-derived water and shoot desiccation. Presumably, such desiccation would be associated with visual symptoms of damage, especially if the observed 73% reduction in apparent $g_c$ largely had been a consequence of the desiccation. However, for unmodified shoots, there was no visual evidence of damaged leaves at the completion of each experiment.

Isolated shoots were used to evaluate the maximum impact that loss of tissue-derived water would have on $g_c$, since, for these shoots, all transpired water would be tissue derived. The $g_l$ of isolated shoots exhibited a rapid linear decline during the first 3 to 4 hr of gas exchange (Fig. 3), and, within 12 to 16 hr of the beginning of each experiment, all leaves were wilted, stems were black and brittle, and no water loss could be detected. These tests confirmed that desiccation could cause decreases in $g_c$, but that substantial desiccation would also be associated with visible symptoms of injury. Some desiccation by the end of 24 hr was evident by the lower leaf water content of both isolated and unmodified shoots, compared with that of shoots taken directly from culture (Table 1). However, unmodified shoots lost considerably more water per unit leaf dry weight (13.6 g) than their initial water content (control, 6.7 g), indicating that the major portion of water transpired by unmodified plants was not water lost by desiccation of tissues, but rather was water replaced by uptake from the medium.

In summary, the data presented indicate that the stomata of these tissue-cultured shoots were functional and closed in response to environmental conditions of increased evaporative demand. Evidence of stomatal closure was also found by microscopic examination (Fig. 4). On all leaves observed, stomata were open when shoots were taken directly from tissue culture conditions (Fig. 4, top). For unmodified shoots, stomata were closed after 24 hr in the gas exchange system (Fig. 4, bottom).

It is not possible to distinguish stomatal ($g_s$) from cuticular ($g_c$) conductance in entire leaves (Ball, 1987), but if all stomata were fully closed after 24 hr in the gas exchange system, then the minimum $g_c$ would represent twice $g_s$, since cuticular water loss occurs from both leaf surfaces. Consequently, for the 11 unmodified shoots tested, the range of apparent $g_c$ was 18 to 64 mmol·m⁻²·s⁻¹. If any water loss occurred from surfaces other than the leaves (i.e., stems) or if stomatal closure on the leaves was not complete, then these values of $g_c$ would be overestimates. On hypostomatous leaves, a more reliable method of estimating $g_c$ (Holmgren et al., 1965) involves prevention of stomatal transpiration by coating abaxial leaf surfaces with a layer of petrolatum, as was done to modified shoots. Microscopically, stomata were largely confined to abaxial leaf surfaces, as had been reported by Sutter (1988). The gas exchange behavior (Fig. 3) for the four modified shoots indicated a range of apparent $g_c$ of 19 to 40 mmol·m⁻²·s⁻¹. The lower limit of this range was somewhat less than that observed for unmodified shoots, indicating that some incomplete stomatal closure may have occurred in unmodified shoots. However, the changes in $g_l$ that were exhibited by some of the modified shoots over 24 hr (Fig. 3) also indicated that stomata in other locations, i.e., the margins of the adaxial leaf surface, as observed by Sutter (1988), may have influenced the values of apparent $g_c$ in modified shoots. Since any stomatal water loss would cause an overestimate of $g_c$, the range of 18 to 40 mmol·m⁻²·s⁻¹ may be regarded as a maximum estimate for the $g_c$ of leaves on these shoots. This entire range is less than the value of $g_c$ reported earlier for similar plants, i.e., 83 mmol·m⁻²·s⁻¹ after conversion from the original units (Sutter, 1988), but may be more reliable since it is based on a larger sample of leaf area and more rigorously calibrated methods. The lower portion of this range is also close to the upper bound of the range of $g_c$ commonly occurring in fully expanded leaves, i.e., 2 to 13 mmol·m⁻²·s⁻¹ after conversion from the original units (Nobel, 1974); hence, we conclude that tissue culture conditions do not necessarily result in abnormal cuticular properties, such as uniformly high values of $g_c$ in apple leaves.

Stresses that are associated with excessive water loss in plants result from an interaction of two independent processes-regulation of water loss and efficient water uptake. In this study, substantial variability between shoots was observed in the rapidity of stomatal closure and in the final value of $g_l$ achieved when shoots were exposed to an increase in evaporative demand. Hence, when exposed to an environmental stress, such

| Treatment                        | N | Leaf | Shoot | Water loss | Water content |
|----------------------------------|---|------|-------|------------|---------------|
| Control                          | 10| 3.56 ± 0.97 | 6.7 ± 2.2 | ---         | 6.7 ± 2.2     |
| Unmodified (stem base in agar)    | 11| 2.01 ± 0.56 | ---       | 13.6 ± 2.0  | ---           |
| Isolated (stem base in petrolatum)| 3 | 1.28 ± 0.32 | ---       | 5.6 ± 2.4   | ---           |

Table 1. Leaf and shoot water content (grams of water per gram of leaf dry weight) for control shoots (directly from tissue culture) and experimental shoots (after 24 hr in the gas exchange system), and the calculated amount of water lost by experimental shoots during gas exchange measurements.

1Number of individual shoots sampled.
2Mean ±1 SD.
3Not applicable.
as removal from culture and transfer to greenhouse conditions, the ability of any individual shoot to restrict its water loss and avoid desiccation may be variable. Some shoots required >24 hr to achieve minimal $g_1$ after being transferred to new environmental conditions. For such shoots, a gradual transition from tissue culture to greenhouse conditions may be necessary. This study has also demonstrated that, despite stomatal closure, unmodified shoots lost an average of twice their water content over 24 hr (Table 1). Presumably, this required water uptake from the medium through the epidermal and cortical cells of the stem. Since these stem tissues may not be specifically differentiated for the radial transport of water, as are the comparable tissues in roots, it is possible that a substantial gradient in water potential was associated with this water movement (Boyer, 1985). Consequently, even low rates of transpiration may have been associated with substantial water stress in the leaves. The occurrence of water stress under gas exchange conditions may account for both the stomatal closure (Fig. 3) and the reduced leaf water content (Table 1) observed in the unmodified shoots of this study. Since the survival of tissue-cultured shoots after removal from culture may depend largely on the balance between water loss and water uptake, factors that enhance the ability of plants to transport water, such as increased length of stem in contact with the media or the presence of roots, may
be as important to survival as stomatal or cuticular regulation of water loss.

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**Inhibition by NaCl of Net CO₂ Fixation and Yield of Cucumber**

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**Abstract.** Cucumber (*Cucumis sativus* L. cv. Fidelio) grown in sand culture in the greenhouse was trickle-irrigated with nutrient solution containing 0, 10, or 50 mM NaCl. Gas exchange of individual leaves was measured by a portable infrared gas analyzer at saturating photosynthetic photon flux. Salt at 10 mM had no detectable effect on plant performance, but exposure to 50 mM NaCl caused net CO₂ fixation to decline by 33% and 48% in the eighth and ninth oldest leaves, respectively. Stomatal conductance and transpiration rate were also reduced (*≈ 50%*) in these leaves. These differences, as well as lower leaf water potentials, were associated with a 60% reduction in fruit fresh weight. The relationship between net CO₂ fixation and intercellular (substomatal) CO₂ concentrations was determined for individual, attached leaves of plants with roots exposed to various concentrations of NaCl in hydroponics. With 50 and 100 mM NaCl, a nonstomatal contribution to the inhibition of photosynthesis at the chloroplast level was Indicated by strong inhibition of CO₂ fixation at a saturating CO₂ concentration. Salt-induced inhibition of CO₂ fixation was associated with accumulation of Na⁺ and Cl⁻, and lower K⁺ in the individual leaves examined.

High productivity of cucumbers in the greenhouse results from optimization of the aerial and rooting environments, with yields between 223 to 446 t·ha⁻¹, compared with 10.7 t·ha⁻¹ under field conditions (Ware and McCollum, 1975). Such heavy yields, combined with higher market prices obtainable out of season, offer advantages to producers located in the southern and southwestern United States, where solar radiation is rarely limiting. However, the quality of water supply in the region is potentially limiting to the productivity of greenhouse cucumbers, a moderately salt-sensitive crop (Gornat et al., 1973; Maas and Hoffman, 1977; Sonnveld and Vogt, 1978; Drews, 1979). The threshold electrical conductivity of the rooting medium of cucumbers, above which appreciable declines in growth and yield can be expected, is between 2.5 and 3.0 dS·m⁻¹ (reviewed by Maas and Hoffman, 1977; see also Kazim and Khaliel, 1983; Maas, 1985), and can be as low as 0.8 to 1.6 dS·m⁻¹ (Jones et al., 1989).

Although mechanisms of salt tolerance and salt injury have been examined in numerous crop species (Maas and Nieman, 1978; Wyn Jones, 1981), there is relatively little information available on cucumber. In general, the more-negative osmotic potential of saline irrigation water contributes adversely to plant water relations, and the excess accumulation of specific ions (especially Na⁺ or Cl⁻) within the leaf mesophyll is a major factor associated with depression of plant growth (Jennings, 1976; Greenway and Munns, 1980).

Many salt-sensitive species tolerate moderate exposure to salt by the roots' ability to exclude ions from the xylem sap and, hence, the leaves through energy-dependent processes (Läuchli, 1984; Drew and Läuchli, 1985). At higher concentrations of salt, the processes of exclusion are inadequate, and excess Na⁺ and/or Cl⁻ are transported via the xylem to the shoot (Drew and Dikumwin, 1985; Drew and Läuchli, 1985). Abnormal concentrations of these ions in the cytoplasm have been associated with...