Vacuolar Acidification in Citrus Fruit: Comparison between Acid Lime (*Citrus aurantifolia*) and Sweet Lime (*Citrus limmetioides*) Juice Cells

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ABSTRACT. Vacuolar acidification was investigated in ‘Palestine’ sweet (*Citrus limmetioides* Tanaka) and ‘Persian’ acid lime [(*Citrus aurantifolia* (Christm.) Swingle)] (vacuolar pHs of 5.0 and 2.1, respectively) using tonoplast vesicles isolated from juice cells. The ATPase activity of tonoplast-enriched vesicles from sweet limes was strongly inhibited by bafilomycin A₁ and NO₃⁻, but was unaffected by vanadate. In contrast, the ATPase activity in acid lime membranes was only slightly inhibited by bafilomycin A₁ and NO₃⁻ and was strongly inhibited by high concentrations of vanadate. The vacuolar origin of the acid lime vesicles was confirmed by immunoblotting. After solubilization and partial purification of the two enzymes by gel filtration, their inhibitor profiles were largely unchanged. Based on equal ATPase activities, vesicles from sweet and acid limes were able to generate similar pH gradients. However, in tonoplast vesicles from sweet limes, the maximum ΔpH was reached four times faster than in those from acid limes. Addition of ethylenediamine tetraacetic acid (EDTA) to chelate Mg⁺² after the maximal ΔpH was attained resulted in collapse of the pH gradient in vesicles from sweet limes, whereas no change in ΔpH was observed in vesicles from acid limes, indicating a less H⁺ permeable membrane. Vacuolar ATPases from both cultivars exhibited identical pH optima and showed similar Mg⁺² dependence, but only the acid lime ATPase activity was inhibited by Ca⁺². These data confirm that the vanadate-sensitive form of the V-ATPase found in lemon and acid limes is specific to hyperacidifying tissues rather than to citrus juice cells. Sweet lime vacuoles bear the classical V-ATPase also found in vegetative plant tissues.

The juice cells of most citrus fruit (*Citrus* L. sp.) accumulate high concentrations of citric acid and soluble sugars, and hyperacidify their vacuoles (Echeverria and Burns, 1989; Echeverria et al., 1992; Müller et al., 1996). In acid limes, for example, the vacuolar pH of juice cells has been determined to be as low as ≈2.0 by two separate and independent methods (Echeverria and Burns, 1989; Echeverria et al., 1992). Juice cells of other citrus fruit such as lemons [*Citrus limon* (L.) Burm.] (Müller et al., 1996) and sweet oranges [*Citrus sinensis* (L.) Osbeck] (Echeverria and Burns, 1989) are also highly acidic, with pH values of 2.2 and 2.8, respectively.

Vacuolar acidification in plant cells results from the H⁺ pumping activity of two tonoplast-bound proton pumps, the V-ATPase and the V-PPase (Rea and Sanders, 1987). In most mature citrus fruit, the trans-tonoplast pH gradient exceeds the thermodynamic maximum ΔpH attainable by the V-PPase, suggesting that the V-ATPase is the dominant pump (Davies, 1994; Schmidt and Briskin, 1993). In lemon juice cells, the large proton gradient between cytosol and vacuolar lumen (ΔpH) has been ascribed in

part to the existence of a novel vanadate-sensitive, chloride and bafilomycin-insensitive tonoplast H⁺-ATPase, characteristics which are contrary to V-ATPase from plant tissues (Müller et al., 1996, 1997, 1999). When compared to tonoplast vesicles from epicotyls of the same species, those from the acidic juice cells were able to generate a larger ΔpH in the presence of Mg-ATP. In addition, lemon fruit tonoplast vesicles maintained a preestablished H⁺ gradient even after cessation of H⁺-ATPase by addition of EDTA, thus demonstrating a high degree of H⁺ impermeability.

In a few citrus cultivars, the high citric acid content characteristic of most citrus fruit is absent. In such cultivars, the lower citrate content is associated with a higher vacuolar pH. For example, the acidless sweet lime (*Citrus limmetioides*) accumulates only a fraction of the citric acid present in its close relatives Persian and Key limes (*Citrus aurantifolia*) and lemons, and is characterized by a vacuolar pH of ≈5.0 (Echeverria and Burns, 1989). The vacuolar H⁺-ATPase is therefore subjected to a different regulatory environment in sweet limes than in acidic limes and lemons.

In the present report, we investigated whether the vanadate sensitive V-ATPase identified previously in lemon juice cells represents a form genetically adapted for vacuolar hyperacidification, or whether it is characteristic of citrus juice cells and differentially regulated in the less acidic cultivars. We found that only the acid lime V-ATPase shares characteristics previously observed in lemon juice sacs and appears to be specific to

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hyperacidifying citrus vacuoles, whereas, the sweet lime juice sac V-ATPase has features similar to those of lemon epicotyls and other vegetative V-ATPases.

Materials and Methods

PLANT MATERIAL. 'Palestine' sweet limes and 'Persian' acid limes were collected August to September 1997 from the Citrus Arboretum at the Florida Department of Agriculture, Winter Haven. For some experiments, acid limes were collected from the lime orchard at the Citrus Research and Education Center, Lake Alfred, Fla.

TONOPLAST ISOLATION. For isolation of tonoplast vesicles from both sweet and acid limes, the procedure of Müller et al. (1996) was followed with a few modifications. About 150 mL of juice cell extract were squeezed directly into 100 mL of homogenization buffer under continuous stirring. In order to keep the pH of the homogenate at around neutrality, the strength of the buffer varied depending on acidity of the fruit. For sweet limes, the homogenization buffer contained 0.5 M MOPS/KOH (pH 8.5) and for acid limes the concentration was increased to 2 M. The homogenate was filtered through a 200 µm pore size nylon mesh and centrifuged at 12,000 g_n for 30 min. After this first centrifugation step, the upper soft part of the pellet was resuspended and combined with the supernatant prior to ultracentrifugation for 30 min at 132,000 g_n . The resulting pellet was resuspended in resuspension buffer (Müller et al., 1996), layered onto an 8%/17%/26%/34% sucrose gradient, and centrifuged for 60 min at 132,000 g_n . In contrast to the procedure of Müller et al. (1996), the sucrose gradient solutions used for these isolations did not contain bovine serum albumin (BSA), as it was found initially that BSA had no effect in recovery or stability of these membranes. Membrane vesicles at the interfaces were collected, rapidly diluted 10× with storage buffer of 10 mM Tris/Mes, 250 mM sorbitol, and 2 mM dithiothreitol (DTT) at pH 7.0, and centrifuged at 132,000 g_n for 30 min. After the final centrifugation, membrane samples were resuspended in 1 mL of storage buffer (at pH 7.0 or pH 5.5 depending on the experiments planned) and stored at -80 °C until needed. Before their use, all vesicles from sweet and acid lime were subjected to three freeze-thaw cycles to equilibrate the lumen with the same buffer solution and for experiments to be conducted under the same conditions.

SIMULTANEOUS MEASUREMENT OF ATPASE ACTIVITY AND PROTON PUMPING. Simultaneous measurements of ATPase activity and H⁺ pumping capacity were performed as described by Palmgren (1990). ATP hydrolysis was coupled to the oxidation of NADH through pyruvate kinase/lactate dehydrogenase, and oxidation of NADH was measured at 340 nm. Formation of a pH gradient was

measured at 496 nm following the absorbance quenching of acridine orange. Reactions were started by addition of MgCl₂ to a reaction mixture containing 50 µg tonoplast vesicle protein. All reactions were carried out at 30 °C in a Shimadzu UV-160 spectrophotometer (Shimadzu Corp., Kyoto, Japan). Data points were recorded and plotted using the Grafit computer program (Erithacus Software Ltd., Horley Surrey, U.K.). All experiments were conducted at least three times.

IMMUNODETECTION OF V-PPASE. Sodium-dodecylsulfate gel electrophoresis (SDS-PAGE) was performed on slab gels containing 12.5% acrylamide and using the Laemmli buffer system (Laemmli et al., 1970). The gels were loaded with equal amount of membrane protein (10 µg) for all samples. For western blot analysis, proteins were transferred electrophoretically from acrylamide gels to cellulose nitrate membranes and subsequently immunostained. Polyclonal antibodies raised against purified mung bean [*Vigna radiatae* (L.) R. Wilcz.] V-PPase were kindly provided by M. Maeshima (Maeshima and Yoshida, 1989). Antibodies were detected with goat anti-rabbit IgG-conjugate alkaline phosphatase (AP) and visualized with bromochloro-indolylphosphate-nitroblue tetrazolium (BCIP/NBT).

PURIFICATION OF THE V-ATPASE. The V-ATPases from sweet and acid limes were solubilized and purified as described previously (Müller et al., 1997) with the following modifications. Fifteen milligrams of tonoplast-enriched sweet lime vesicles were solubilized with 2.5% octyl-β-glucoside in solubilization buffer (10 mM BTP-Mes, pH 7.6, 10% glycerol, 1 mM EDTA, 8 mM MgSO₄, 5 mM DTT, L-α-phosphatidylcholine liposomes at 200 µg·mL⁻¹, and 0.012% butylated hydroxytoluene) in a total volume of 5 mL. The solubilized proteins were centrifuged for 10 min at 412,000 g_n in a rotor (TLA-100.3; Beckman Instruments, Inc., Paloalto, Calif). The supernatant was applied to a 100 × 1-cm Sephacryl S-400HR (Amersham Pharmacia Biotech, Uppsala, Sweden) chromatography column equilibrated and eluted at 4.5 mL·h⁻¹ in running buffer (10 mM Tris-Mes, pH 7.0, 0.3% (w/w) Triton X-100, L-α-phosphatidylcholine at 100 µg·mL⁻¹, 10% glycerol, 1 mM EDTA, 4 mM MgCl₂, 5 mM DTT, and 50 µM PMSF). Fractions with ATPase activity were pooled and applied to a Mono-Q anion exchange chromatography column (Pharmacia) equilibrated in 5 mM Tris-HCl, pH 6.0, 0.3% (w/w) Triton X-100, L-α-phosphatidylcholine at 50 µg·mL⁻¹, 10% glycerol, 1 mM EDTA, 4 mM MgCl₂, 2 mM DTT, and 50 µM PMSF. The column was eluted with a linear 0 to 0.4 M KCl gradient, and the fractions with ATPase activity were pooled again, diluted to a final KCl concentration of 0.05 M, and applied to a second Mono-Q column. Elution from the second anion exchange column was performed with a 0.3 M KCl step gradient to recover all activity in one 0.5 mL fraction. The fraction containing ATPase activity was adjusted to

Table 1. ATPase activity in membrane fractions obtained from the interfaces of a discontinuous sucrose gradient. Original microsomal fractions were prepared from acid and sweet lime juice cells. Activity was determined in a coupled reaction with pyruvate kinase/lactate dehydrogenase and measured as the decrease in absorbance of NADH at 340 nm.

Treatment	Percent ATPase activity (Pi, nmol·min ⁻¹ ·mg ⁻¹ protein)					
	8%/17%		17%/26%		26%/34%	
	Acid lime	Sweet lime	Acid lime	Sweet lime	Acid lime	Sweet lime
Control (total activity)	803 (100%)	656 (100%)	513 (100%)	531 (100%)	345 (100%)	328 (100%)
+Bafilomycin A (10 nM)	96%	6.6%	96%	10%	94%	50%
+NO ₃ ⁻ (50 mM)	80%	15%	83%	25%	80%	70%
+Vanadate (100 µM)	31%	68%	34%	60%	75%	80%
+Azide (2 mM)	94%	80%	75%	72%	40%	38%

Fig. 1. Immunodetection of V-PPase in membrane samples from acid lime (lanes 1–3), sweet lime (lane 4), and red beet hypocotyl (lane 5) by antibodies against mung bean V-PPase. Lanes 1–3 correspond to the three interfaces of the sucrose gradient (8/17, 17/26, and 26/34 interfaces, respectively). All lanes were loaded with 10 µg total protein and u = unified atomic mass unit.

2% octyl- β -glucoside, loaded onto a 12% to 25% glycerol gradient in 10 mM BTP-Mes, pH 7.0, 0.15% Triton X-100, 1 mM EDTA, 4 mM MgSO₄, 10 mM KCl, L- α -phosphatidylcholine at 50 µmol·L⁻¹, 5 mM DTT, and 50 µM PMSF, and centrifuged for 18 h at 250,000 g_n in a Beckman SW-41 rotor. The purified V-ATPase was used for activity measurements and its subunits analyzed by SDS-PAGE. The purification method for the acid lime H⁺-ATPase was similar to that of sweet limes, except that the solubilization buffer contained only 1% octyl- β -glucoside, and after centrifugation of the solubilized membranes, the fluffy top layer of the pellet was resuspended and loaded together with the supernatant onto the Sephacryl S-400 HR column. In addition, before glycerol gradient centrifugation, the detergent concentration of the partially purified V-ATPase of acid limes was not adjusted to 2%. Concentration of octyl- β -glucoside were optimized for solubilization and for the maintenance of V-ATPases integrity. These optimal concentrations were established after conducting solubility-activity curves for both types of V-APases under increasing concentrations of detergent.

PROTEIN DETERMINATION. Protein was determined as described by Bradford (1976) using Commassie Plus Protein Assay Reagent (Pierce Co. Rockford, Ill.).

Results

IDENTIFICATION OF TONOPLAST FRACTIONS. Isopycnic separation of microsomal samples from sweet and acid lime juice cells in a discontinuous sucrose gradient resulted in three distinct membrane layers. ATPase activity in the lightest fraction from sweet limes exhibited the typical inhibitor profile of tonoplast H⁺-ATPases. The sweet lime enzyme was strongly inhibited by NO₃⁻ and bafilomycin A₁, and only slightly inhibited by vanadate or azide (Table 1). In contrast, the acid lime tonoplast ATPase activity was mostly insensitive to NO₃⁻ and bafilomycin A₁, but was inhibited by high concentrations of vanadate.

To verify the vacuolar origin of the low density fractions from acid limes, we used antibodies raised against the mung bean V-PPase (Maeshima and Yoshida, 1989). Highly pure tonoplast samples from red beet (*Beta vulgaris* L. ssp. *vulgaris*) hypocotyl (Echeverria and Gonzalez, 2000) and sweet lime (Echeverria et al., 1997) were included for comparative purposes (Fig. 1). Although some limited cross-reactivity was observed in all membrane fractions, the strongest antibody signal corresponding to V-PPase was increasingly evident in samples from the lower density layers (8%/17% sucrose), confirming that tonoplast-enriched vesicles were recovered from these fractions (Fig. 1). In lemon

juice cells, kinetic and biochemical characterization had already demonstrated the corresponding fractions to be of tonoplast origin (Müller et al., 1996, 1997, 1999). Secondary bands present in the sweet lime and red beet hypocotyl samples were attributed either to cross reactivity with aquaporin aggregates, or to proteolytic products of the V-PPase 73 ku (u = unified atomic mass unit) peptide (Maeshima, 1992). On a protein basis, the antibody cross-reactivity with acid lime tonoplast vesicles was significantly weaker than the signal obtained with sweet lime vesicles. This finding extends observations made in lemon, where, based on activity measurements, the V-PPase appeared to be only a minor constituent of the tonoplast membrane (Taiz, unpublished results). All subsequent experiments were performed with membranes collected from the lightest fractions.

The effects of increasing concentrations of bafilomycin A₁, NO₃⁻, and vanadate on the tonoplast-bound ATPase from sweet and acid limes are presented in Fig. 2. Bafilomycin A₁ had little

Fig. 2. Activity of acid and sweet lime juice cell V-ATPase in the presence of increasing concentrations of (A) bafilomycin-A, (B) nitrate, and (C) vanadate. Activities were measured by following the oxidation of NADH in the presence of pyruvate kinase-lactic dehydrogenase at 340 nm. Legend in A also applies to all figures and vertical bars = $\pm$ SD (n = 6).

effect on acid lime ATPase even at concentrations of 100 nM, whereas 10 nM inhibited corresponding activity from sweet limes by 90% (Fig. 2A). Acid lime tonoplast ATPase was partially inhibited by NO_3^- , but the degree of inhibition was smaller than for the sweet lime ATPase (Fig. 2B) and the V-ATPases from other sources (Brauer et al., 1992; Hosaka et al., 1994; Williams et al., 1990).

Similar to what was found in lemon (Müller et al., 1996, 1997), acid lime V-ATPase activity was strongly inhibited by 20 to 100 μM vanadate (Fig. 2C). Only a small reduction in sweet lime V-ATPase activity was observed under the same conditions. The profiles of inhibition of the acid lime V-ATPase by vanadate, NO_3^- , and bafilomycin A_1 are very similar to the profiles found in tonoplast-enriched vesicles of mature lemon juice cells, whereas the sweet lime juice cell V-ATPase resembles the V-ATPase of lemon epicotyls (Müller et al., 1996).

KINETIC PROPERTIES. V-ATPases from sweet and acid limes showed similar pH dependence with a pH optimum of 6.7 for both (data not presented). This value is slightly lower than the pH 7.0

optimum reported for lemon fruit and epicotyls (Müller et al., 1996), 7.5 for citrus leaves (Bañuls et al., 1993), and 8.0 in peanut (*Arachis hypogaea* L.) seedlings (Soma and Sharma, 1994). Sweet and acid limes were equally dependent on Mg^{2+} , and for both, the Michaelis-Menten curve was hyperbolic with an estimated K_m of ≈ 1.0 mM (Fig. 3A). There was little effect of Ca^{2+} on sweet lime activity up to 10 mM (Fig. 3B). By contrast, under similar conditions, the acid lime V-ATPase activity was inhibited over 80% by 4 mM Ca^{2+} .

FORMATION OF H^+ GRADIENT AND MEMBRANE PERMEABILITY.

The proton pumping activities of tonoplast enriched fractions from acid and sweet limes were compared by using acridine orange absorption quenching. Based on equivalent ATPase activity, both acid and sweet lime samples generated similar pH gradients (Fig. 4). However, sweet lime tonoplast membranes reached steady state faster than acid lime vesicles (6 min vs. 30 min in Fig. 4). After steady state was reached, addition of EDTA to chelate Mg^{2+} and stop the ATPase reaction resulted in collapse of the pH gradient within 10 min in tonoplast vesicles from sweet limes. In contrast, the pH gradient built up by acid lime vesicles was unaffected under similar conditions (Fig. 5). Addition of 10 μM gramicidin collapsed the remaining pH gradient in both cases. The impermeability of the acid lime tonoplast to protons was very comparable to that of the lemon tonoplast (Müller et al., 1996) and may be a common characteristic of highly acidic cells.

PURIFICATION OF THE V-ATPASES OF SWEET AND ACID LIMES.

After solubilization with octyl- β -glucoside and centrifugation, the soluble ATPase activities from sweet and acid limes were layered onto a Sephacryl S-400 HR column for initial purification by gel filtration. Elution profiles are shown in Fig. 6. Consistent

Fig. 3. Effect of (A) Mg^{2+} and (B) Ca^{2+} on the V-ATPase activity from acid and sweet lime juice cells. The reaction mixture contained 150 mM KCl before addition of MgCl_2 . Activities were measured by following the oxidation of NADH in the presence of pyruvate kinase-lactic dehydrogenase at 340 nm. Legend in A also applies to B and vertical bars = $\pm$ SD ($n = 6$).

Fig. 4. Simultaneous measurements of H^+ /ATPase activity ($\circ$, left scale) and H^+ gradient formation ($\bullet$, right scale) by 50 μg tonoplast vesicles from (A) sweet lime and (B) acid lime juice cells. Reactions were started by addition of MgCl_2 to a final concentration of 4 mM. ATPase activity and pH formation were measured simultaneously as described in Materials and Methods.

Fig. 5. Effect of 20 mM EDTA (to stop ATPase activity) and 10 μ M gramicidin on the established H^+ gradient in vesicles from sweet (○) and acid (■) lime juice cells.

with what was found previously for the V-ATPases from lemon juice sacs and epicotyls, the ATPase activities from sweet and acid limes eluted in a single peak. The partially purified sweet lime juice cell ATPase was found to be sensitive to nitrate but insensitive to inhibition by vanadate. In contrast, the partially purified acid lime ATPase was inhibited by both, nitrate and high concentrations of vanadate, and thus resembles the lemon fruit V-ATPase. The V-ATPases of sweet and acid limes were further purified by glycerol gradient centrifugation and anion exchange chromatography. The purified enzymes were then analyzed by SDS-PAGE. Results for the sweet lime V-ATPase are shown in Fig. 7. Subunits of ≈ 97 , 66, 55, 52, 42/43, 31, 17, and 13 ku are visible. A 33 ku-band, like that found to comigrate with the ATPase activity of lemon fruit, was absent in the sweet lime V-ATPase and the pronounced 16 ku-band, previously described for lemon fruit, was barely visible in the sweet lime. Although a similar analysis of the acid lime V-ATPase indicated that it was enriched in the usual V-ATPase subunits, the acid lime enzyme was less stable than the sweet lime V-ATPase and appeared to disintegrate during the medium pressure Mono-Q anion exchange chromatography (data not presented).

Discussion

Acidification of the vacuolar compartment in plant cells is essential for maintenance of the electrochemical potential gradient ($\Delta\mu H^+$) that serves to energize the secondary active transport of sugars (Getz, 1991; Keller, 1992), amino acids (Homeyer and Schultz, 1988), ions (Barkla and Pantoja, 1996; Schumaker and Sze, 1986), and a large number of secondary metabolites (Blom et al., 1991; Deus-Neumann and Zenk, 1986). The transport of organic anions into the vacuole is thought to be driven by the H^+ concentration difference across the tonoplast mem-

brane (Brune et al., 1998; Oleski et al., 1987; Rentsch and Martinoia, 1991), generated by two electrogenic proton pumps, a V-ATPase and an inorganic V-PPase, with the V-ATPase being dominant in most tissues (Cheffings et al., 1997; Müller et al., 1996).

Most citrus fruit highly are acidic, a property associated with a high concentration of citric acid. In the exceedingly acidic Persian and Key limes (*Citrus aurantifolia*) and Bearss lemon, for example, the concentration of citric acid reaches beyond 325 mM, whereas in the acidless sweet lime, citric acid does not exceed 5 mM (Echeverria and Ismail, 1987). Differential intracellular fractionation (Echeverria and Valich, 1988) and direct pH measurements (Echeverria et al., 1992) have established that the low pH of cell extracts and the intracellular location of citric acid can be ascribed to the large central vacuole.

Citrate accumulation in the vacuole of plant cells is determined primarily by the vacuolar $\Delta\mu H^+$ (Brune et al., 1998; Oleski et al., 1987; Rentsch and Martinoia, 1991). In citrus juice cells, although a portion of the accumulated citrate is transported by a $\Delta\mu H^+$ -independent, ATP-dependent uptake mechanism (Brune et al., 1998; Canel et al., 1995), the majority of the accumulation is driven by the ΔpH . Therefore, the final concentration of citric acid in the vacuole is dependent on the ability of the tonoplast-bound ATPase to pump H^+ into the lumen, and on the permeability of the tonoplast to retain the H^+ . The V-PPase is unlikely to be

Fig. 6. Partial purification of the V-ATPases from (A) sweet and (B) acid limes. Membrane proteins from tonoplast-enriched vesicles from sweet limes and acid limes were solubilized with octyl- β -glucoside and separated on a 1×100 -cm Sephacryl S-400 HR chromatography column. Profiles of the ATPase activity sensitive to 50 mM nitrate (○) or 100 μ M vanadate (●) are shown. Legends in A applies to B.

involved in hyperacidification for three reasons: 1) PPase activity in lemon fruit juice cells is negligible (Müller et al., 1996); 2) thermodynamic constraints prevent the V-PPase from generating such steep pH gradients (Schmidt and Briskin, 1993); and 3) V-PPase activity in acidic citrus fruit has been demonstrated to function in the PPi synthesis direction rather than in the hydrolytic mode (Marsh et al., 2000)

Müller et al. (1996) suggested that vacuolar hyperacidification in lemon juice cells was facilitated by a novel tonoplast-bound V-ATPase that was sensitive to high concentrations of vanadate. This enzyme was absent from lemon epicotyl cells which exhibited a higher vacuolar pH. In addition, the tonoplast of juice cells was more impermeable to protons than that of epicotyls. The results of the present investigation confirm that hyperacidification in juice cells of acidic citrus fruit is achieved via a vanadate-sensitive V-ATPase and a tonoplast that is impermeable to H⁺ diffusion. This conclusion is based on: 1) data demonstrating that a vanadate and Ca²⁺ sensitive, NO₃⁻ and bafilomycin insensitive V-ATPase (Figs. 2 and 3) similar to that of lemon, is present at the tonoplast of acid limes juice cells, but absent from the tonoplast of a high pH, low citrate variety; 2) an apparent lower H⁺/ATP coupling ratio for the acid lime V-ATPase (Fig. 4), which may be necessary to reach the trans tonoplast pH difference observed in vivo; and 3) notable differences in the ability of the tonoplasts to maintain a pH gradient after inactivation of the V-ATPase (Fig. 5).

Compared to the V-ATPases from sweet limes and those from sugarcane (*Saccharum officinarum* L.) (Williams et al., 1990), pear (*Pyrus communis* L.) fruit (Hosaka et al., 1994) and maize (*Zea mays* L.) roots (Brauer et al., 1992), tonoplast-bound ATPases from lemon and acid lime juice cells are unusual in their lack of sensitivity to bafilomycin A₁, NO₃⁻, and by their sensitivity to vanadate. This unusual inhibitor profile was maintained during purification of the enzyme by gel filtration and anion exchange chromatography, and thus appears to represent a hallmark of hyperacidifying proton pumps in *Citrus*. The conventional inhibitor profile of the sweet lime V-ATPase proves that juice cells with less acidic vacuoles do not possess the vanadate-sensitive pump.

When normalized to the initial rates of H⁺ pumping, Müller et al. (1996) noted that the lemon juice cell V-ATPase was able to generate a significantly larger ΔpH than the corresponding enzyme from epicotyls. However, when based on identical ATPase activity rates, the lemon fruit and epicotyl enzymes reached similar pH gradients, with the fruit enzyme taking twice as long to reach equilibrium as the epicotyl V-ATPase (Müller et al., 1999). This was also the case for acid and sweet lime juice cell samples when compared on an equal V-AT-

Fig. 7. SDS-polyacrylamide gel of the purified V-ATPase from sweet lime. The solubilized V-ATPase from sweet lime was purified by gel filtration, anion exchange chromatography, and glycerol gradient centrifugation, and separated by gel electrophoresis. The molecular weight of standard proteins (Std) is indicated on the left of the gel. V-ATPase subunits are identified on the right.

Pase activity basis (Fig. 4). Although both V-ATPases were able to generate comparable ΔpHs, the H⁺ pumping rate of the sweet lime ATPase was virtually double that of acid limes, suggesting a difference in their H⁺/ATP coupling ratio.

In agreement with the previous characterization of lemon fruit V-ATPase (Müller et al., 1996, 1997, 1999), formation and maintenance of the large ΔpH across the tonoplast of highly acidic citrus juice cells is made possible by the presence of a specialized V-ATPase and a tonoplast highly impermeable to H⁺. The low vacuolar pH of ≈2.0, and a trans-tonoplast ΔpH of about five units are sufficient to drive the uptake of 325 mM citric acid from a cytosolic citrate concentration of <10 mM (Brune et al., 1998). This capacity for vacuolar hyperacidification is absent from citrus fruit and other vegetative cells with a higher vacuolar pH. Therefore, the novel vanadate-sensitive V-ATPase described for lemons and acid limes appears to be a characteristic of hyperacidifying tissues and not a property of citrus juice cells in general.

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