The plasma membrane H⁺-ATPase, a simple polypeptide with a long history

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Abstract
The plasma membrane H⁺-ATPase of fungi and plants is a single polypeptide of fewer than 1,000 residues that extrudes protons from the cell against a large electric and concentration gradient. The minimalist structure of this nanomachine is in stark contrast to that of the large multi-subunit F₀F₁ ATPase of mitochondria, which is also a proton pump, but under physiological conditions runs in the reverse direction to act as an ATP synthase. The plasma membrane H⁺-ATPase is a P-type ATPase, defined by having an obligatory phosphorylated reaction cycle intermediate, like cation pumps of animal membranes, and thus, this pump has a completely different mechanism to that of F₀F₁ ATPases, which operates by rotary catalysis. The work that led to these insights in plasma membrane H⁺-ATPases of fungi and plants has a long history, which is briefly summarized in this review.

KEYWORDS
Arabidopsis thaliana, F-type ATPase, Neurospora crassa, Nicotiana tabacum, proton pump, P-type ATPase, Saccharomyces cerevisiae, Schizosaccharomyces pombe

1  |  INTRODUCTION

It has been known since ancient times that yeast secretes acid during glucose fermentation without this phenomenon being connected with any essential process in yeast. This review recapitulates the discovery of the plasma membrane proton pump (H⁺-ATPase) and will highlight recent advances in our understanding of the function, regulation, and structure of this pump, so surprisingly different from other proton pumps that it caused eyebrows to rise (as the introductory quotes indicate). The history to be described is divided into several parts, the biophysical, the biochemical, and the molecular biology approaches. Here, only a brief account is given, which is focused and dedicated to the work and memory of Professor André Goffeau. For a more detailed account of the early history, the reader is referred to Goffeau and Slayman (1981). Subsequently, some mention will be made to the history of the corresponding enzyme in plants.

... This raises a fundamental question. Why are the two presumed proton pumps of the yeast cell, the mitochondrial and the plasma membrane ATPases, so different in their structure and mechanism? ... A valid answer to the apparent paradox raised by the existence of two distinct proton pumps in yeast requires definite elucidation of another question. Does the plasma membrane ATPase from yeast really pump protons and only protons? Amory, Foury, and Goffeau (1980)

From time to time, surprise has been expressed that the proton-motive F₀F₁ ATPases should be so different structurally from the cation-motive E₁E₂ ATPases, and that their (chemiosmotic) mechanisms should presumably be so different inasmuch as the E₁E₂ ATPases involve a phosphorylated intermediate while the F₀F₁ ATPases do not. Mitchell and Koppenol (1982)
In the late 1960s, the chemiosmotic hypothesis of Peter Mitchell won acceptance and, with the words of Efraim Racker (1975), “became a favorite tool in the design of experiments.” Oxidative phosphorylation of ADP to generate ATP had been shown to be carried out by the F$_{0}$F$_{1}$ ATPase, a large mitochondrial complex, which as an energy source utilizes a proton motive force that has two components: a proton gradient and a membrane potential. In the 1970s, it became increasingly clear that the plasma membrane of fungi and plants is equipped with a system running in the reverse direction, thus operating as an ATP-driven proton pump that generates a proton motive force.

2 | HISTORY OF THE PHYSIOLOGY OF FUNGAL PLASMA MEMBRANE H$^{+}$-ATPases

The father of the plasma membrane H$^{+}$-ATPase is Clifford L. Slayman who, using inserted microelectrodes, recognized that cells of the fungus Neurospora crassa have a substantial plasma membrane potential approximating a quarter of a volt. Inhibitors of ATP synthesis caused depolarization of the membrane, suggesting that formation of the membrane potential depended on ATP consumption (Slayman, 1965a, 1965b; Slayman, Long, & Lu, 1973; Slayman, Lu, & Shane, 1970). It was suggested that the membrane potential is sustained by ejection of protons coupled to the splitting of ATP. This membrane potential then serves as the main energy distributor for transport and—together with an inward chemical gradient for protons—drives the uptake of a variety of different substances (Slayman & Gradmann, 1975). Protons were hypothesized to be the transported ion because hydrogen ions, almost alone among the common inorganic cations and anions, had a steep depolarizing effect: 30–40 mV for each unit decrease of pH (Slayman & Gradmann, 1975). This hypothesis has now proven to be correct; however, at the time, it was not supported by any biochemical or molecular evidence. Prokaryotes have plasma membrane-localized F$_{0}$F$_{1}$ ATPases. Therefore, even if the presence of a proton pump in the plasma membrane could be demonstrated, it needed not to be unrelated to the F$_{0}$F$_{1}$ ATPase of mitochondria.

An important characteristic of the F$_{0}$F$_{1}$ ATPase is that it is sensitive to the antibiotic oligomycin (Lardy, Johnson, & McMurray, 1958); in fact, the F$_{0}$ in F$_{0}$F$_{1}$ received its name for being the oligomycin-sensitive factor (Racker, 1963). Already in 1967, it was reported that the plasma membrane of the yeast Saccharomyces cerevisiae contained an ATPase activity that was insensitive to oligomycin (Matile, Moor, & Mühlthaler, 1967). In situ freeze-etching and subsequent negative staining by phosphotungstate of yeast cells revealed that the outer surface of the plasma membrane was covered with globular particles of about 150 Å in diameter. To purify plasma membranes, a microsomal membrane preparation was centrifuged through a linear density gradient of urografin, and a distinctive white band was isolated in which the membranes were sculptures with particles of the same size. It was thus concluded that this membrane fraction represented the plasma membrane. Matile et al. (1967) found that the most remarkable constituent of the plasma membrane was a polysaccharide identified as mannan, and the activity of the single enzyme they could detect, an ATPase. The ATPase was dependent on Mg$^{2+}$ for activity but was completely insensitive to the mitochondrial ATPase inhibitor oligomycin. It was stated:

No direct evidence with regard to the functional meaning of the ATPase can be presented yet. It is likely that this enzyme is involved in the energy-dependent uptake of nutrients from the culture medium; that is, permeases such as amino acid absorbing enzymes which are present in yeast cells may be identical with the ATPase bound to the plasma membrane.

The hypothesis that nutrient uptake systems were directly energized by ATP was prevalent at that time.

It had been suggested before that H$^{+}$ secretion and/or ATP was required for uptake of nutrients into fungal cells. Conway and O’Malley (1946) observed that addition of K$^{+}$ to the medium of fermenting yeast cells increased secretion of H$^{+}$ and noted:

The rapid absorption of K is essentially an interchange with H ions, as shown by the practically quantitative relations of K absorbed and H extruded ... It would appear that the nature of the potassium absorption consists very largely in a direct exchange of K and H ions.

Peña, Cinco, Puyou, and Tueno (1969) studied the effect of K$^{+}$ on the respiration and glycolysis of yeast cells and noted:

Measurement of the levels of ADP, ATP and Pi revealed that K$^{+}$ stimulates an enzymatic activity that diminishes ATP and increases ADP... This might mean that the ion induces the breakdown of ATP by an indirect mechanism, with another phosphorylated compound being hydrolyzed directly ... The most obvious condition of this sort would be the pH increase created inside the cell as H$^{+}$ is exchanged for K$^{+}$.

Similarly, Eddy, Indge, Backen, and Nowacki (1970) noted that glycine uptake into yeast cells was markedly increased when pH of the medium was lowered from pH 7 to pH 4.5. This pH dependence was more marked when glucose was absent than when it was present. In contrast, high external K$^{+}$ inhibited glycine uptake. This led them to suggest:

The observations lead to the idea that, in certain circumstances, H$^{+}$ and K$^{+}$ may play part in the transport of amino acids by yeast that is analogous to the roles of Na$^{+}$ and K$^{+}$ in mammalian systems.

In these early studies, there was no suggestion that nutrient uptake could be a process indirectly fueled by metabolic energy, and there was no mentioning of an ATPase or a H$^{+}$ pump.

Foury and Goffeau (1975) observed a correlation between all three parameters: ATPase activity, extrusion of protons, and uptake of nutrients. They found that Dio-9, an F$_{0}$F$_{1}$ ATPase inhibitor (Schatz, Penefsky, & Racker, 1967), reduced intracellular ATP levels and instantaneously suppressed the cellular ejection of protons as well as the uptake of uridine and amino acids. However, Dio-9 also inhibited proton extrusion in the presence of antimycin A, an inhibitor of respiration. Furthermore, in the presence of antimycin A, addition of glucose to glucose-starved cells increased ATP levels and proton extrusion. This led Foury and Goffeau (1975) to conclude:
These results suggest that in these conditions, the target of Dio-9 is not the mitochondrial ATPase but a plasma membrane proton-translocating function generating an electrochemical gradient required for active transport.

3 HISTORY OF THE BIOCHEMISTRY OF FUNGAL PLASMA MEMBRANE H⁺-ATPases

The heroes who biochemically characterized the plasma membrane ATPase, demonstrated that it was a P-type ATPase and not an FₒFₒ ATPase, and finally proved it to be a proton pump were many, but prominent names were André Goffeau (working on Schizosaccharomyces pombe), Ramon Serrano (working on Saccharomyces cerevisiae), and Gene Scarborough (working on Neurospora crassa).

To characterize the elusive enzyme, plasma membranes first had to be purified with minimal contamination of mitochondrial membranes having FₒFₒ ATPase activity. This was not a trivial task as no marker for the plasma membrane was available at the time. Gene Scarborough solved this problem by employing a wall-less mutant of N. crassa that was labeled on the outside with concanavalin A, which binds to carbohydrates on the outer membrane surface. Cells were then disrupted, and a concanavalin A-labeled membrane fraction was obtained. The plasma membrane-enriched fraction contained a major membrane-embedded enzyme that hydrolyzed ATP and was stimulated by Na⁺ and K⁺ (Scarborough, 1975). Stimulation by these cations be purified with minimal contamination of mitochondrial membranes.

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The same characteristics were soon thereafter reported for the plasma membrane ATPase of the yeasts S. pombe (Delhez et al., 1977) and S. cerevisiae (Serrano, 1978). Delhez et al. (1977) separated microsomal membrane vesicles by centrifugation on a sucrose density gradient and obtained two peaks with ATP hydrolytic activity. Membrane vesicles in the fraction with the highest density were stained with osmium tetroxide, which is indicative of mannose and thus polysaccharides, a characteristic of the outer surface of the plasma membrane. The ATPase in the plasma membrane-enriched fraction had a pH optimum around pH 5.3, far more acidic than the pH optimum of 9 in the lighter fraction, which is characteristic of the FₒFₒ ATPase (Delhez et al., 1977). Serrano (1978) used a very similar purification strategy for isolating S. cerevisiae plasma membranes and tested their plasma membrane origin by postfixation in osmium tetroxide and staining with uranyl acetate and lead citrate. The plasma membrane fraction contained an ATPase activity with a pH optimum of 5.4 that was insensitive to oligomycin.

The work of Scarborough had established that there was an electrogenic ATPase at the plasma membrane of N. crassa, but the polypeptide(s) responsible for the process had not been identified. Characterization of the plasma membrane ATPase first required its purification in a catalytically active form. Using the mild detergent lysophosphatidylcholine, Dufour and Goffeau (1978) were able to solubilize the S. pombe plasma membrane fraction, and after subsequent centrifugation of the solubilized membrane proteins through a linear 10% to 30% (w/v) sucrose gradient, the ATPase activity became separated from the bulk of contaminating proteins, which were of lower sedimentation rate. The plasma membrane ATPase peak fraction of the sucrose gradient was analyzed by SDS-polyacrylamide gel electrophoresis followed by Coomassie blue staining. Only a single polypeptide with an apparent molecular weight of 105,000 could be observed. Thus, the purified plasma membrane ATPase of S. pombe exhibited a much simpler subunit composition than the mitochondrial, chloroplast, or bacterial ATPases. In contrast, it appeared very similar in size to the catalytic subunit of mammalian Na⁺/K⁺-stimulated plasma membrane and Ca²⁺-stimulated sarcoplasmic reticulum ATPases. These enzymes were at that time called E₁E₂ ATPases because during catalysis, they appeared to alternate between two major conformations, Enzyme 1 (E₁) and Enzyme 2 (E₂; Siegel & Albers, 1967). The plasma membrane ATPase of S. cerevisiae could not be solubilized in an active form with lysophosphatidylcholine, but could be solubilized with a synthetic zwitterionic detergent and subsequently purified by centrifugation through a linear glycerol gradient (Malpartida & Serrano, 1980). In the fraction with highest ATP hydrolytic activity, 85% of the protein corresponded to a 105,000-Mw polypeptide, thus confirming the single subunit composition of the plasma membrane ATPase. Subsequently, the N. crassa ATPase was purified and shown to have similar characteristics as the S. pombe and S. cerevisiae enzymes (Bowman, Blasco, & Slayman, 1981).

E₁E₂ ATPases are now commonly called P-type ATPases because phosphorylation of a conserved aspartate residue initiates pumping during catalysis. Before classification of the plasma membrane ATPase as a P-type ATPase, it was essential to demonstrate that a phosphorylated intermediate was formed. Willsky (1979) analyzed a crude microsomal fraction from S. cerevisiae and found that at pH 5.5 (optimum pH of the plasma membrane H⁺-ATPase), three polyproteptides at 210,000, 160,000, and 115,000 molecular weight, respectively, were labeled with [γ-³²P]ATP. All phosphoproteins behaved as phosphorylated intermediates of E₁E₂ ATPases. Thus, they formed rapidly in the presence of Mg²⁺, there was a rapid turnover of the bound phosphate when unlaabeled ATP was added, and dephosphorylation occurred after incubation with hydroxylamine, which is indicative that the phosphate linkage is a carboxylic acid anhydride linkage (Lipmann & Tuttle, 1945). In addition, vanadate, an inhibitor of E₁E₂ ATPase activity, blocked the phosphorylation of the 210,000- and 115,000- Da proteins. Serrano and Malpartida (1979) carried out similar experiments with a plasma membrane fraction from S. cerevisiae and found a 100,000-Mw band to be radiolabeled by ³²P-ATP. Using N. crassa plasma
membranes, Dame and Scarborough (1980) similarly found that, following phosphorylation by $^{32}$P-ATP, a radiolabeled protein migrated in gels with an apparent molecular weight of 100,000. The radiolabel disappeared if the membranes had been pretreated with trypsin but was protected if incubation was in the presence of MgATP. SDS polyacrylamide gel electrophoresis followed by Coomassie Blue staining showed that a polypeptide with an apparent molecular weight of 105,000 was cleaved by trypsin but was protected if MgATP was present during proteolysis. Strikingly, the ATP hydrolytic activity of the plasma membranes was sensitive to trypsin treatment, but not in the presence of MgATP. This served as a strong indication that the radiolabeled polypeptide indeed corresponded to a 100,000-Mw ATPase. Dame and Scarborough (1981) further characterized the chemical nature of the phosphoryl enzyme intermediate of the phosphorylated N. crassa plasma membrane enzyme and demonstrated that it was an aspartylphosphate, identical to that of the Na$^+$/K$^+$-ATPase and Ca$^{2+}$-ATPase of animal cells.

Still, the identity between the radiolabeled protein and the plasma membrane ATPase remained uncertain. Final proof for a phosphorylated reaction cycle intermediate of the plasma membrane ATPase came when the purified 100,000-Mw plasma membrane ATPase from S. pombe was shown to form a phosphorylated reaction cycle intermediate (Amory et al., 1980). The phosphorylated intermediate reached the steady-state level in less than 2 s and rapidly turned over. The phosphobond was cleaved by hydroxylamine and was relatively stable in acids but readily hydrolyzed in alkaline or in acid alcoholic media. These results demonstrated that the intermediate was an acylphosphate. Similar results were subsequently obtained with the purified S. cerevisiae enzyme (Malpartida & Serrano, 1981a).

Dame and Scarborough (1980), still using a plasma membrane preparation, used two independent methods to demonstrate that the ATPase of N. crassa plasma membranes is a proton pump. This was not a straightforward task as protons cannot be labeled and a demonstration of their transport across a membrane required that they could be measured on one side of a membrane or the other. First, isolated N. crassa plasma membrane vesicles were shown to catalyze concentrative uptake of the probe $^{14}$C imidazole, which is membrane permeant in its unprotonated form but remains trapped into vesicles when protonated. MgATP-dependent imidazole uptake and MgATP-dependent membrane potential generation (measured as SCN$^-$ uptake) displayed identical saturation kinetics with respect to the concentration of MgATP and were inhibited in parallel by the P-type ATPase inhibitor orthovanadate. Furthermore, the addition of ATP (in the presence of Mg$^{2+}$) to vesicles containing fluorescein-labeled dextran, which is a pH indicator in the pH range between 5 and 8, gave rise to time-dependent fluorescence quenching that was markedly stimulated by the addition of SCN$^-$ and was abolished by orthovanadate. Taken together, these experiments demonstrated that the N. crassa ATPase was an electrogenic enzyme catalyzing H$^+$ transport.

What remained to be shown was whether the purified single polypeptide plasma membrane ATPase was sufficient for proton pumping. In principle, many more polypeptides (subunits) or accessory proteins present in the plasma membrane preparation could be required for the process. This could be addressed by reconstituting the purified protein into artificial phospholipid bilayer vesicles and measuring ATP-dependent transport of protons into the vesicle lumen. Methods for reconstituting the F$_{O}$F$_{1}$ ATPase had been developed by Efraim Racker in the late 1970s (Racker, Violand, O’Neal, Alfonzo, & Telford, 1979). Ramon Serrano had learned the technique during a postdoctoral stay in Racker’s laboratory (Serrano, Kanner, & Racker, 1976) and reconstituted the purified plasma membrane ATPase of S. cerevisiae by a freeze-thaw sonication procedure (Malpartida & Serrano, 1981b). Proton pumping was subsequently measured by an indirect approach. The idea, which had been tested on the F$_{O}$F$_{1}$ ATPase (Serrano et al., 1976), was that, once a steep proton gradient had been established, the pump would operate in reverse and catalyze ATP synthesis. Conditions were established to measure $^{32}$Pi-ATP exchange, during which $^{32}$Pi is incorporated into cold ATP, and testing whether the reaction was sensitive to protonophores and other ionophores. Surprisingly, the reaction was only partly sensitive to the addition of the proton ionophore (uncoupler) carbonyl cyanide m-chlorophenyl hydrazine (CCCP), but was further inhibited by addition of the potassium ionophore valinomycin, which by itself was ineffective. This made it difficult to conclude whether the enzyme was an electrogenic H$^+$ pump or an electroneutral H$^+$/K$^+$ exchanger (Malpartida & Serrano, 1981b). In a subsequent publication (Malpartida & Serrano, 1981c), this view was modified in favor of transport of H$^+$ only, as another proton ionophore, 1799, was shown to completely inhibit the energy transfer reaction. Furthermore, ATP-dependent acidification of the vesicle lumen was documented by quenching of fluorescence of the acridine dye 9-amino-6-chloro-2-methoxyacridine (ACMA), which had previously been employed to monitor pH gradients (acidic inside) in submitochondrial particles (Azzi, Fabbro, Santato, & Gherardini, 1971).

Conclusive proof that the purified 105,000-Mw polypeptide operates on its own as an electrogenic H$^+$-ATPase came following reconstitution of the S. pombe enzyme into lipid vesicles (Villalobo, Boulty, & Goffeau, 1981). Proton movement in the external medium was directly monitored with a pH-electrode. Upon addition of MgATP, a fast proton uptake took place when countertransport of K$^+$ (for charge compensation) was facilitated by inclusion of the K$^+$ ionophore valinomycin. That the observed H$^+$ translocation only took place in the presence of valinomycin was evidence of the electrogenic nature of the H$^+$ pumping by the ATPase. Moreover, during steady-state ATP hydrolysis, a fast and transient proton uptake was also observed when the generated membrane potential was collapsed upon addition of valinomycin in the presence of K$^+$. The ATPase activity of the reconstituted enzyme was strongly stimulated by the H$^+$-conducting agent CCCP or by an association of the K$^+$/H$^+$ carrier nigericin plus valinomycin in the presence of K$^+$. An important discovery that demonstrated that the plasma membrane H$^+$-ATPase is under tight metabolic control came when it was shown that the enzyme is posttranslationally activated by glucose metabolism in a way that increases proton pumping tenfold (Serrano, 1983).
Ramon Serrano (working on S. cerevisiae), and Carolyn W. Slayman (working first on N. crassa and later on S. cerevisiae).

The plasma membrane H⁺-ATPase gene Plasma Membrane ATPase1 (PMA1) was first cloned from S. cerevisiae by Serrano, Kielland-Brandt, and Fink (1986), who could also show that the gene was essential for growth of yeast, and subsequently the corresponding gene was cloned from N. crassa (Addison, 1986; Hager et al., 1986) and S. pombe (Ghislain, Schlesser, & Goffeau, 1987). A second gene, PMA2, was identified in S. cerevisiae (Schlesser, Ulaszewski, Ghislain, & Goffeau, 1988) and S. pombe (Ghislain & Goffeau, 1991). Analysis of the sequences proved that the plasma membrane H⁺-ATPase is indeed related to mammalian P-type ATPases such as the Na⁺/K⁺- and the muscle Ca²⁺-ATPases. With the gene sequences in hand, the door was open for genetic work and for structure–function analysis by mutagenesis.

A significant finding was that the glucose activation of the S. cerevisiae enzyme involved a short stretch of 11 C-terminal residues that appeared to function as an autoinhibitor, which regulates the pump activity (Portillo, de Larrinoa, & Serrano, 1989). Later, it was found that glucose activation involved phosphorylation of two residues in this regulatory C-terminal stretch (Lecchi et al., 2007; Lecchi, Allen, Pardo, Mason, & Slayman, 2005).

Soon, the importance of other residues for functioning of the yeast pump was investigated by site-directed mutagenesis (Serrano & Portillo, 1990). However, studying in detail loss-of-function mutants was a problem, as the yeast plasma membrane H⁺-ATPase is an essential enzyme and the cells would stop growing if the pump was not active (Serrano et al., 1986). This problem was solved by designing expression systems in which the endogenous wild-type plasma membrane H⁺-ATPase was put under the control of an inducible promoter (Cid, Perona, & Serrano, 1987) or in which mutant H⁺-ATPases accumulated in secretory vesicles (Nakamoto, Rao, & Slayman, 1991). However, a problem that still remained was that mutations of many conserved residues gave rise to misfolded or mistargeted mutant proteins (DeWitt, Santos, Allen, & Slayman, 1998; Nakamoto et al., 1998), which complicated their analysis. For an in-depth review on the structure, function, and biogenesis of the yeast plasma membrane H⁺-ATPase, the reader is referred to Morsomme, Slayman, and Goffeau (2000).

5 | A FOLLOW UP ON THE PLANT PLASMA MEMBRANE H⁺-ATPase

It was suggested already in the beginning of the 20th century that ion uptake in plants was an active process distinct from diffusion. For reviews on the early history of the study of ion transport in plants, the reader is referred to Higinbotham (1973a, 1973b), Poole (1978), and Palmgren (1998).

At about the same time as the chemiosmotic hypothesis for ATP synthesis in mitochondria was proposed by Mitchell (Mitchell, 1961), Jack Dainty (1962) in a review discussed the presence of a plasma membrane potential in plants and their importance for ion transport and discussed whether ion pumps could be responsible for the electric potential. It is evident from the text that even the notion of a plasma membrane in plants was controversial at the time:

We may fairly ask the question: do the ion pumps make any direct contribution to the membrane potential? In the jargon of the specialists in this field: are any of the ion pumps electrogenic? Or are they neutral... I do not propose to enter here into the controversy about the existence of the external plasma membrane, which I shall call the plasmalemma despite objections to the universal application of this term to the, presumed, external permeability barrier. I personally consider the evidence in its favor... I think this article should have shown that the proper study of ion transport in plant cells is in its infancy. Any investigation which is concerned only with ion concentrations is practically certain to lead to incorrect deductions because electrical potential gradients are just as important as concentration gradients in determining passive ion movements...

A problem with plant material for electrophysiological studies was that the cells are small, the vacuole is big, the existence of a plasma membrane was not even established, so it was difficult to be certain what was actually measured and across which membranes following insertion of the microelectrodes. To solve this problem, Spanswick, Stolarek, and Williams (1967) measured membrane potentials in the giant internodal cells of the macroscopic green alga Nitella translucens where it could be excluded that potential differences between the vacuole and its surroundings contributed to the recordings. As is evident from the conclusion, no mention is made about the possibility of a H⁺ pump:

The possibility of a contribution to the plasmalemma potential from electrogenic pumps is briefly discussed...

This analysis shows that Na is actively transported from the cytoplasm into the medium as well as into the vacuole; K is pumped into the cytoplasm from the medium but appears to be close to electrochemical equilibrium across the tonoplast.

Inspired by the work of Slayman (1965a, 1965b), where the effect of external ions and metabolic inhibitors on the N. crassa membrane potential were studied, Kitasato (1968) used the metabolic inhibitor dinitrophenol (DNP) to study the origin of the membrane potential in N. translucens and became the first to obtain electrophysiological evidence pointing to a H⁺ extruding pump in plants. He concluded:

In artificial pond water containing DNP, the resting membrane potential decreased; this suggested that some energy-consuming mechanism maintains the membrane potential at the resting level. It is probable that there is a H⁺ extrusion mechanism in the Nitella cell, because the potential difference between the resting potential and the H⁺ equilibrium potential is always maintained notwithstanding a continuous H⁺ inward current which should result from the potential difference... Perhaps the effect of DNP is the most conclusive evidence for a metabolic component of the observed membrane potential of the Nitella cell. Such a
A plant plasma membrane ATPase was first characterized by Hodges, Leonard, Bracker, and Keenan (1972). As a marker for the plant plasma membrane, they used a stain consisting of periodic acid, chromic acid, and phosphotungstic acid and could separate them from other membranes by discontinuous sucrose-gradient centrifugation of homogenates from oat roots. An ATPase activity was found to copurify with the plasma membranes and was characterized (Hodges et al., 1972; Leonard & Hodges, 1973). They found the ATPase to be stimulated by multiple cations and suggested (Leonard & Hodges, 1973) that this single ATPase could be involved in the direct uptake of several ions:

In conclusion, purified plasma membranes of oat roots contain an ion-stimulated ATPase that has kinetic properties similar to the kinetics of monovalent cation transport. This, along with other results discussed, provides strong evidence for a role for this ATPase in ion absorption by plants. The kinetic results presented here are also consistent with the concept that only one transport system, consisting of binding sites possessing varying affinities for ions, is involved in the absorption of ions from very low (0.01 mM) to very high (100 mM) concentrations.

With the advent of techniques of molecular biology, sequences of plasma membrane H+-ATPase genes were cloned from Arabidopsis thaliana (Harper, Surowy, & Sussman, 1989; Harper, Manney, DeWitt, Yoo, & Sussman, 1990; Pardo & Serrano, 1989), Nicotiana plumaginifolia (Boutry, Michelet, & Goffeau, 1989), and Lycopersicum esculentum (Ewing, Wimmers, Meyer, Chetelat, & Bennett, 1990). This definitely showed the plant plasma membrane H+-ATPase to be evolutionarily related not only to its fungal counterparts but also to P-type ATPases of animals. Soon thereafter, it became apparent that in each plant investigated, a multigene family of around 10 plasma membrane H+-ATPase genes is present (reviewed in Arango, Gévaudant, Oufattole, & Boutry, 2003; Sondergaard, Schulz, & Palmgren, 2004). Different plasma membrane H+-ATPase genes vary in their spatiotemporal expression patterns and often have specialized physiological roles in the plant body. In A. thaliana, the two most highly expressed isoforms are Autoinhibited H+-ATPase1 and 2 (AHA1 and AHA2). Under optimal laboratory growth conditions, plants carrying deletions in either gene appear normal, but the homozygous double mutations cause embryo lethality (Haruta et al., 2010). This demonstrates that in plants, like in yeast, the plasma membrane H+-ATPase is essential for cell function.

The heterologous expression of the plant H+-ATPase in cells of S. cerevisiae with reduced expression or devoid of endogenous plasma membrane H+-ATPase proved to be a very convenient tool for producing recombinant plant plasma membrane H+-ATPases in large quantities (de Kerchove d’Exaerde et al., 1995; Villalba, Palmgren, Berberian, Ferguson, & Serrano, 1992). With these systems in hand, it was possible to carry out detailed structure–function analysis by mutagenesis of plant plasma membrane H+-ATPases. It has been used later on by the group of Goffeau to study thermophilic H+-ATPases from Archea (Morsomme et al., 2002).

Using the technology of heterologous expression of plant plasma membrane H+-ATPase in yeast, it could be confirmed by mutagenesis that the C-terminal region of the protein serves as an autoinhibitory domain (Palmgren & Christensen, 1993). Thus, whereas the full-length A. thaliana AHA2 gene could not complement functionally the loss of endogenous yeast plasma membrane H+-ATPase PMA1, a mutated
AHA2 gene encoding a protein truncated by 92 C-terminal residues complemented a pma1 null mutant. Subsequent mapping by mutagenesis of the C-terminal domain revealed that it contained two autoinhibitory regions and thus was considerably larger and more complicated in structure than the short C-terminal autoinhibitory stretch in the yeast H⁺-ATPase (Axelsen, Venema, Jahn, Baunsgaard, & Palmgren, 1999; Regenberg, Villaiba, Lanfermeijer, & Palmgren, 1995).

Mutant screening further identified residues in plant plasma membrane H⁺-ATPases that were outside of the C-terminal domain but when they became mutated also gave rise to an activated enzyme (Morsomme et al., 1996; Morsomme, Dambly, Maudoux, & Boutry, 1998). These residues were scattered throughout the primary structure of the pump but were predicted in the 3D structure to form a possible intramolecular receptor for the C-terminal regulatory domain.

Another feature of the C-terminal domain of the plant pump that became apparent was that it interacted with activating 14-3-3 regulatory proteins (Jahn et al., 1997), that the interaction between the H⁺-ATPase and 14-3-3 protein produced a binding site for the fungal toxin fusicoccin (Baunsgaard et al., 1998; Piotrowski, Morsomme, Boutry, & Gecking, 1998), and that binding of 14-3-3 protein to the H⁺-ATPase required phosphorylation of the penultimate residue of the H⁺-ATPase (always a threonine) (Fuglsang et al., 1999; Maudoux et al., 2000; Svennelid et al., 1999). Later, it was realized that also the N-terminal domain is taking part in the regulation of the plant plasma membrane H⁺-ATPase (Ekberg, Palmgren, Veierskov, & Buch-Pedersen, 2010). For recent updates on the regulation of the plant plasma membrane H⁺-ATPase by phosphorylation and other factors, the reader is referred to Duby and Boutry (2009), Haruta, Gray, and Sussman (2015), and Falhof et al. (2016).

Site-directed mutagenesis of the plant plasma membrane H⁺-ATPase also resulted in the identification of residues in the transmembrane domain that influence proton pumping. A single proton acceptor-donor was found to be Asp-684 (AHA2 numbering) in transmembrane segment 6, whereas the positively charged Arg-655 in the neighboring transmembrane segment 5 appeared to regulate pumping albeit it was not essential (Buch-Pedersen et al., 2003a). Based on this work, a model was proposed in which a role for the conserved Arg-655 is to move in during pumping and provide positive charge to the proton binding site, possibly by polarizing the Asp-684 side chain and modulating its pKₐ, in this way promoting release of the bound H⁺ (Buch-Pedersen & Palmgren, 2003b). The yeast expression system allowed for production of amounts of AHA2 enzyme sufficient for crystallographic studies. The final crystal structure calculated at 3.6 Å resolution was a major step forward (Pedersen, Buch-Pedersen, Morth, Palmgren, & Nissen, 2007) and confirmed earlier predictions on the mechanism of H⁺ transport by the pump and proved the plant plasma membrane H⁺-ATPase to be very similar in structure to other P-type ATPases such as the Ca²⁺-ATPase and Na⁺/K⁺-ATPase (reviewed in Buch-Pedersen, Pedersen, Veierskov, Nissen, and Palmgren 2009). Unfortunately, neither the N- nor the C-terminal domains were ordered in the structures, so we still do not know how they interfere with pump activity, for which purpose a structure of the pump in its autoinhibited state is required. What is also lacking is high-resolution 3D structures of both plant and fungal plasma membrane H⁺-ATPases, which would allow for determining in more detail the H⁺ transport mechanism. These are important goals for the future.

As is evident from this work, many scientists have contributed to the history of the discovery of the plasma membrane H⁺-ATPase, and often the same discoveries were made by different laboratories at about the same time. We apologize for not having been able to cite all those who have been part of the journey and will end this historical treatise by a relevant quote of Efraim Racker:

> Perhaps scientists are too much concerned with giving and receiving credits. The research work in biochemistry in the twentieth century is probably like the building of cathedrals in the Middle Ages. It is the work of many, and the identity of those who participated in their creation is a matter of little consequence and will soon be forgotten. Racker (1976)

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**REFERENCES**

Addison, R. (1986). Primary structure of the Neurospora plasma membrane H⁺-ATPase deduced from the gene sequence. Homology to Na⁺/K⁺-, Ca²⁺-, and K⁺-ATPase. The Journal of Biological Chemistry, 261, 14896–14901.

Amory, A., Foury, F., & Goffeau, A. (1980). The purified plasma membrane ATPase of the yeast Schizosaccharomyces pombe forms a phosphorylated intermediate. The Journal of Biological Chemistry, 255, 9353–9357.

Arango, M., Gévaudant, F., Outefftote, M., & Boutry, M. (2003). The plasma membrane proton pump ATPase: The significance of gene subfamilies. Planta, 216, 355–365.

Axelsen, K. B., Venema, K., Jahn, T., Baunsgaard, L., & Palmgren, M. G. (1999). Molecular dissection of the C-terminal regulatory domain of the plant plasma membrane H⁺-ATPase AHA2: Mapping of residues that when altered give rise to an activated enzyme. Biochemistry, 38, 7227–7234. [https://doi.org/10.1021/bi982482l](https://doi.org/10.1021/bi982482l)

Azzl, A., Fabbro, A., Santato, M., & Gherardini, P. L. (1971). Energy transduction in mitochondriol fragments. Interaction of the membrane with acridine dyes. European Journal of Biochemistry, 21, 404–410. [https://doi.org/10.1111/j.1432-1033.1971.tb01484.x](https://doi.org/10.1111/j.1432-1033.1971.tb01484.x)

Baunsgaard, L., Fuglsang, A. T., Jahn, T., Korthout, H. A., de Boer, A. H., & Palmgren, M. G. (1998). The 14-3-3 proteins associate with the plant plasma membrane H⁺-ATPase to generate a fuscoiccin binding complex and a fuscoiccin responsive system. The Plant Journal, 13, 661–671. [https://doi.org/10.1046/j.1365-313X.1998.00083.x](https://doi.org/10.1046/j.1365-313X.1998.00083.x)

Boutry, M., Michelet, B., & Goffeau, A. (1989). Molecular cloning of a family of plant genes encoding a protein homologous to plasma membrane H⁺-translocating ATPases. Biochemical and Biophysical Research Communications, 162, 567–574. [https://doi.org/10.1016/0006-291X(89)92348-6](https://doi.org/10.1016/0006-291X(89)92348-6)

Bowman, B. J., Blasco, F., & Slayman, C. W. (1981). Purification and characterization of the plasma membrane ATPase of Neurospora crassa. The Journal of Biological Chemistry, 256, 12343–12349.

Bowman, B. J., & Slayman, C. W. (1977). Characterization of plasma membrane adenosine triphosphatase of Neurospora crassa. The Journal of Biological Chemistry, 252, 3357–3363.
Briskin, D. P., & Leonard, R. T. (1982a). Phosphorylation of the adenosine triphosphatase in a deoxycholate-treated plasma membrane fraction from corn roots. *Plant Physiology, 70*, 1459–1464. https://doi.org/10.1104/pp.70.5.1459

Briskin, D. P., & Leonard, R. T. (1982b). Partial characterization of a phosphorylated intermediate associated with the plasma membrane ATPase of corn roots. *Proceedings of the National Academy of Sciences of the United States of America, 79*, 6922–6926. https://doi.org/10.1073/pnas.79.22.6922

Buch-Pedersen, M. J., & Palmgren, M. G. (2003a). Conserved Asp684 in transmembrane segment M6 of the plant plasma membrane P-type proton pump AHA2 is a molecular determinant of proton translocation. *The Journal of Biological Chemistry, 278*, 17845–17851.

Buch-Pedersen, M. J., & Palmgren, M. G. (2003b). Mechanism of proton transport by plant plasma membrane proton ATPases. *Journal of Plant Research, 116*, 507–515. https://doi.org/10.1007/s10265-003-0111-9

Buch-Pedersen, M. J., Pedersen, B. P., Veierskov, B., Nissen, P., & Palmgren, M. G. (2009). Protons and how they are transported by proton pumps. *Pflügers Archiv, 457*, 573–579. https://doi.org/10.1007/s00424-008-0503-8

Cid, A., Perona, R., & Serrano, R. (1987). Replacement of the promoter of *Saccharomyces carlsbergensis* by a galactose promoter gene from *Schizosaccharomyces pombe*. *Current Genetics, 19*, 2931–2937. https://doi.org/10.1007/BF00434664

Conway, E. J., & O'Malley, E. (1946). The nature of the cation exchanges during yeast fermentation, with formation of 0.02N-H ion. *The Biochemical Journal, 40*, 59–67. https://doi.org/10.1042/bj0400059

Dainty, J. (1962). Ion transport and electrical potentials in plant cells. *Annual Review of Plant Physiology, 13*, 379–402. https://doi.org/10.1146/annurev.pp.13.060162.022115

Dame, J. B., & Scarborough, G. A. (1980). Identification of the hydrolytic moiety of the Neurospora plasma membrane H+-ATPase and demonstration of a phosphoryl-enzyme intermediate in its catalytic mechanism. *Biochemistry, 19*, 2931–2937. https://doi.org/10.1021/bi00554a018

Dame, J. B., & Scarborough, G. A. (1981). Identification of the phosphorylated intermediate of the Neurospora plasma membrane H+-ATPase as beta-aspartyl phosphate. *The Journal of Biological Chemistry, 256*, 10724–10730.

de Kerchove d’Exaerde, A., Supply, P., Dufour, J. P., Bogaerts, P., Thinés, D., Goffau, A. T., Visconti, S., Drumm, K., Jahn, T., Stensballe, A., Mattei, B., ... Palmgren, M. G. (1990). Molecular cloning of tomato plasma membrane H+-ATPase. *The Plant Journal, 4*, 845–852. http://www.biochemj.org/content/120/4/845

Ekberg, K., Palmgren, M. G., Veierskov, B., & Buch-Pedersen, M. J. (2010). A novel mechanism of P-type ATPase autoinhibition involving both termini of the protein. *The Journal of Biological Chemistry, 285*, 7344–7350. https://doi.org/10.1074/jbc.M109.096123

Ewing, N. N., Wimmers, L. E., Meyer, D. J., Chetelat, R. T., & Bennett, A. B. (1990). Molecular cloning of tomato plasma membrane H-ATPase. *Plant Physiology, 94*, 1874–1881. https://doi.org/10.1104/pp.94.4.1874

Falhof, J., Pedersen, J. T., Fuglsang, A. T., & Palmgren, M. (2016). Plasma membrane H+-ATPase regulation in the center of plant physiology. *Molecular Plant, 9*, 323–337. https://doi.org/10.1016/j.molp.2015.11.002

Fouri, F., & Goffeau, A. (1975). Stimulation of active uptake of nucleosides and amino acids by cyclic adenosine 3’5’-monophosphate in the yeast *Schizosaccharomyces pombe*. *The Journal of Biological Chemistry, 250*, 2354–2362.

Fuglsang, A. T., Visconti, S., Drumm, K., Jahn, T., Stensballe, A., Mattei, B., ... Palmgren, M. G. (1999). Binding of 14-3-3 protein to the plasma membrane H+-ATPase AHA2 involves the three C-terminal residues Tyr946, Thr–Val and requires phosphorylation of Thr947. *The Journal of Biological Chemistry, 274*, 36774–36780. https://doi.org/10.1074/jbc.J045718200

Ghislain, M., & Goffeau, A. (1991). The pma1 and pma2 H+-ATPases from *Schizosaccharomyces pombe* are functionally interchangeable. *The Journal of Biological Chemistry, 266*, 18276–18279.

Ghislain, M., Schlessier, A., & Goffeau, A. (1987). Mutation of a conserved glycine residue modifies the vanadate sensitivity of the plasma membrane H+-ATPase from *Schizosaccharomyces pombe*. *The Journal of Biological Chemistry, 262*, 17549–17555.

Goffeau, A., & Slayman, C. W. (1981). The proton-translocating ATPase of the fungal plasma membrane. *Biochimica et Biophysica Acta, 639*, 197–223. https://doi.org/10.1016/0006-3898(81)90010-0

Hager, K. M., Mandalia, S. M., Davenport, J. W., Speicher, D. W., Benz, E. J. Jr., & Slayman, C. W. (1986). Amino acid sequence of the plasma membrane ATPase of *Neurospora crassa*: Deduction from genomic and cDNA sequences. *Proceedings of the National Academy of Sciences of the United States of America, 83*, 7693–7697. https://doi.org/10.1073/pnas.83.20.7693

Harper, J. F., Manney, L., DeWitt, N. D., Yoo, M. H., & Sussman, M. R. (1990). *The Arabidopsis thaliana* plasma membrane H+-ATPase multigene family. Genomic sequence and expression of a third isofrom. *The Journal of Biological Chemistry, 265*, 13601–13608.

Harper, J. F., Surowy, T. K., & Sussman, M. R. (1989). Molecular cloning and sequence of cDNA encoding the plasma membrane proton pump (H+-ATPase) of *Arabidopsis thaliana*. *Proceedings of the National Academy of Sciences of the United States of America, 86*, 1234–1238. https://doi.org/10.1073/pnas.86.4.1234

Haruta, M., Burch, H. L., Nelson, R. B., Barrett-Wilt, G., Kline, K. G., Mohsin, S. B., ... Sussman, M. R. (2010). Molecular characterization of mutant Arabidopsis plants with reduced plasma membrane proton pump activity. *The Journal of Biological Chemistry, 285*, 17918–17929. https://doi.org/10.1074/jbc.M110.101733

Haruta, M., Gray, W. M., & Sussman, M. R. (2015). Regulation of the plasma membrane proton pump (H+-ATPase) by phosphorylation. *Current Opinion in Plant Biology, 28*, 68–75. https://doi.org/10.1016/j.pbi.2015.09.005

Higinbotham, N. (1973a). The mineral absorption process in plants. *The Botanical Review, 39*, 15–69. https://doi.org/10.1007/BF02860069

Higinbotham, N. (1973b). Electropotentials of plant cells. *Annual Review of Plant Physiology, 24*, 25–46. https://doi.org/10.1146/annurev.ph.24.060173.000325

Hodges, T. K., Leonard, R. T., Bracker, C. E., & Keenan, T. W. (1972). Purification of an ion-stimulated adenosine triphosphatase from plant roots: Association with plasma membranes. *Proceedings of the National Academy of Sciences of the United States of America, 69*, 3307–3311. https://doi.org/10.1073/pnas.69.11.3307
Maudoux, O., Batoko, H., Oecking, C., Gevaert, K., Vandekerckhove, J., Matile, P., Moor, H., & Mühlethaler, K. (1967). Isolation and properties of Malpartida, F., & Serrano, R. (1981c). Reconstitution of the proton pump.

Malpartida, F., & Serrano, R. (1980). Purification of the yeast plasma mem-

Lecchi, S., Nelson, C. J., Allen, K. E., Swaney, D. L., Thompson, K. L., Coon, J.

Mitchell, P., & Koppenol, W. H. (1982). Chemiosmotic ATPase mechanisms.

Malpartida, F., & Serrano, R. (1981c). Reconstitution of the proton translocating adenosine triphosphatase of yeast plasma membranes.

Leonard, R. T., & Hodges, T. K. (1973). Characterization of plasma membrane-associated adenosine triphosphate activity of oat roots. Plant Physiology, 52, 6–12. https://doi.org/10.1101/pp.52.1.6

Lipmann, F., & Tuttle, L. C. (1945). The detection of activated carboxyl groups with hydroxylamine as interceptor. The Journal of Biological Chemistry, 161, 415–416.

Malpartida, F., & Serrano, R. (1980). Purification of the yeast plasma mem-

Lecchi, S., Allen, K. E., Pardo, J. P., Mason, A. B., & Slayman, C. W. (2005). Tandem phosphorylation of Ser111–Thr52–Thr116–Thr116–7940 in the carboxy tail.

The Journal of Biological Chemistry 1990–1993, 265, 13423–13426.

Malpartida, F., & Serrano, R. (1980). Purification of the yeast plasma mem-

Malpartida, F., & Serrano, R. (1981a). Phosphorylated intermediate of the ATPase from the plasma membrane of yeast. European Journal of Biochemistry, 116, 413–417. https://doi.org/10.1111/j.1432-1033.1981.tb05350.x

Malpartida, F., & Serrano, R. (1981b). Proton translocation catalyzed by the purified yeast plasma membrane ATPase reconstituted in liposomes. FEBS Letters, 131, 351–354. https://doi.org/10.1016/0014-5793(81)80401-2

Malpartida, F., & Serrano, R. (1981c). Reconstitution of the proton-translocating adenosine triphosphatase of yeast plasma membranes. The Journal of Biological Chemistry, 256, 4175–4177.

Matile, P., Moor, H., & Mühlthaler, K. (1967). Isolation and properties of the plasmalemma in yeast. Archiv für Mikrobiologie, 58, 201–211. https://doi.org/10.1007/BF00408804

Maudoux, O., Batoko, H., Oecking, C., Gavaert, K., Vankerckhove, J., Boutry, M., & Morsomme, P. (2000). A plant plasma membrane H+-ATPase expressed in yeast is activated by phosphorylation at its penultimate residue and binding of 14-3-3 regulatory proteins in the absence of fusicoccin. The Journal of Biological Chemistry, 275, 17762–17770. https://doi.org/10.1074/jbc.M909690199

Mitchell, P. (1961). Coupling of phosphorylation to electron and hydrogen transfer by a chemi-osmotic type of mechanism. Nature, 191, 144–148. https://doi.org/10.1038/191144a0

Mitchell, P., & Koppel, W., H. (1982). Chemiosmotic ATPase mechanisms. Annals of the New York Academy of Sciences, 402, 584–601. https://doi.org/10.1111/j.1749-6632.1982.tb25785.x

Morsomme, P., Chami, M., Marco, S., Nader, J., Ketchum, K., Goffeaule, A., & Rigaud, J. L. (2002). Characterisation of an hyperthermophilic P-type ATPase from Methanococcus jannaschii expressed in yeast. The Journal of Biological Chemistry, 277, 29608–29616. https://doi.org/10.1074/jbc.M203871200

Morsomme, P., Dambly, S., Maudoux, O., & Boutry, M. (1998). Single point mutations distributed in 10 soluble and membrane regions of the Nicotiana plumbaginifolia plasma membrane PMA2 H+-ATPase activate the enzyme and modify the structure of the C-terminal region. The Journal of Biological Chemistry, 273, 34837–34842. https://doi.org/10.1074/jbc.273.52.34837

Morsomme, P., de Kerchove d’Exaerde, A., De Meester, S., Thélin, D., Goffeau, A., & Boutry, M. (1996). Single point mutations in various domains of a plant plasma membrane H+-ATPase expressed in Saccha-

Morsomme, P., Slayman, C. W., & Goffeau, A. (2000). Mutagenic study of the structure, function and biogenesis of the yeast plasma membrane H+-ATPase. Biochimica et Biophysica Acta, 1469, 133–157. https://doi.org/10.1016/S0304-4157(00)00015-0

Nakamoto, R. K., Rao, R., & Slayman, C. W. (1991). Expression of the yeast plasma membrane [H+]ATPase in secretory vesicles. A new strategy for directed mutagenesis. The Journal of Biological Chemistry, 266, 7940–7949.

Nakamoto, R. K., Verjovski-Almeida, S., Allen, K. E., Ambesi, R., Rao, R., & Slayman, C. W. (1998). Substitutions of aspartate 378 in the phosphorylation domain of the yeast PMA1 H+-ATPase disrupt protein folding and biogenesis. The Journal of Biological Chemistry, 273, 7338–7344. https://doi.org/10.1074/jbc.273.13.7338

Palmgren, M. G. (1998). Proton gradients and plant growth: Role of the plasma membrane H+-ATPase. Advances in Botanical Research, 28, 1–70. https://doi.org/10.1016/S0065-2296(08)60293-1

Palmgren, M. G., & Christensen, G. (1993). Complementation in situ of the yeast plasma membrane H+-ATPase gene pma1 by an H+-ATPase gene from a heterologous species. FEBS Letters, 317, 216–222. https://febs.onlinelibrary.wiley.com/doi/abs/10.1016/0014-5793(93)92893-9

Palmgren, M. G., Larsson, C., & Sommarin, M. (1990). Proteolytic activation of the plant plasma membrane H+-ATPase by removal of a terminal segment. The Journal of Biological Chemistry 1990, 265, 13423–13426.

Palmgren, M. G., Sommarin, M., Serrano, R., & Larsson, C. (1991). Identification of an autoinhibitory domain in the C-terminal region of the plant plasma membrane H+-ATPase. The Journal of Biological Chemistry, 266, 20470–20475.

Pardo, J. M., & Serrano, R. (1989). Structure of a plasma membrane H+-ATPase gene from the plant Arabidopsis thaliana. The Journal of Biological Chemistry, 264, 8557–8562.

Pederen, B. P., Buch-Pedersen, M. J., Morth, J. P., Palmgren, M. G., & Nissen, P. (2007). Crystal structure of the plasma membrane proton pump. Nature, 450, 1111–1114. https://doi.org/10.1038/nature06417

Peña, A., Cinco, G., Puyou, A. G., & Tuena, M. (1969). Studies on the mechanism of the stimulation of glycolysis and respiration by K+ in Saccharomyces cerevisiae. Biochimica et Biophysica Acta, 180, 1–8. https://doi.org/10.1016/0006-2789(69)90187-X

Piotrowski, M., Morsomme, P., Boutry, M., & Oecking, C. (1998). Complementation of the Saccharomyces cerevisiae plasma membrane H+-ATPase by a plant H+-ATPase generates a highly abundant fusicoccin binding site. The Journal of Biological Chemistry, 273, 30018–30023. https://doi.org/10.1074/jbc.273.45.30018

Poole, R. J. (1978). Energy coupling for membrane transport. Annual Review of Plant Physiology, 29, 437–460. https://doi.org/10.1146/annurev.pp.29.060178.002253

Portillo, F., de Larinoa, I. F., & Serrano, R. (1989). Deletion analysis of yeast plasma membrane H+-ATPase and identification of a regulatory domain at the carboxyl-terminus. FEBS Letters, 247, 381–385. https://doi.org/10.1016/0014-5793(89)81375-4

Racker, E. (1963). A mitochondrial factor conferring oligomycin sensitivity on soluble mitochondrial ATPase. Biochemical and Biophysical Research Communications, 10, 435–439. https://doi.org/10.1016/0006-291X(63)90375-9

Racker, E. (1975). Reconstitution, mechanism of action and control of ion pumps. Biochemical Society Transactions, 3, 785–802. http://www.biochemsoctrans.org/content/3/6/785
Racker, E. (1976). A new look at mechanisms in bioenergetics, Academic Press. eBook ISBN: 9780323160117

Racker, E., Violand, B., O’Neal, S., Alfonso, M., & Telford, J. (1979). Reconstitution, a way of biochemical research; some new approaches to membrane-bound enzymes. Archives of Biochemistry and Biophysics, 198, 470–477. https://doi.org/10.1016/0003-9861(79)90521-6

Regenberg, B., Villalba, J. M., Lanfermeijer, F. C., & Palmgren, M. G. (1995). C-terminal deletion analysis of plant plasma membrane H+-ATPase: Yeast as a model system for solute transport across the plant plasma membrane. Plant Cell, 7, 1655–1666.

Scalla, R., Amory, A., Rigaud, J., & Goffeau, A. (1983). Phosphorylated intermembrane bound enzymes. Proceedings of the National Academy of Sciences of the United States of America, 73, 1485–1488. https://doi.org/10.1073/pnas.73.5.1485

Schatz, G., Penefsky, H. S., & Racker, E. (1967). Partial resolution of the enzymes catalyzing oxidative phosphorylation. XIV. The Journal of Biological Chemistry, 242, 2525–2560.

Schlessner, A., Ulaszewski, S., Ghislain, M., & Goffeau, A. (1998). A second transport ATPase gene in Saccharomyces cerevisiae. The Journal of Biological Chemistry, 263, 19480–19487.

Serrano, R. (1978). Characterization of the plasma membrane ATPase of Neurospora crassa plasma membranes. The Journal of Biological Chemistry, 250, 1106–1111.

Serrano, G. A. (1976). The neurospora plasma membrane ATPase is an electrophoretic pump. Proceedings of the National Academy of Sciences of the United States of America, 73, 1485–1488. https://doi.org/10.1073/pnas.73.5.1485

Scalia, R., Amory, A., Rigaud, J., & Goffeau, A. (1983). Phosphorylated intermediate of a transport ATPase and activity of protein kinase in membranes from corn roots. European Journal of Biochemistry, 132, 525–530. https://doi.org/10.1111/j.1432-1033.1983.tb07393.x

Scarborough, G. A. (1975). Isolation and characterization of Neurospora crassa plasma membranes. The Journal of Biological Chemistry, 250, 1106–1111.

Serrano, R. (1990). Catalytic and regulatory sites of yeast plasma membrane H+-ATPase studied by directed mutagenesis. Biochimica et Biophysica Acta, 1018, 195–199. https://doi.org/10.1016/0005-2728(90)90247-2

Siegel, G. J., & Albers, R. W. (1967). Sodium-potassium-activated adenosine triphosphatase of Electrophorus electric organ. IV. Modification of responses to sodium and potassium by arsenite plus 2,3-dimercaptopropanol. The Journal of Biological Chemistry, 242, 4972–4979.

Slayman, C. L. (1965a). Electrical properties of Neurospora crassa. Effects of external cations on the intracellular potential. The Journal of General Physiology, 49, 69–92. https://doi.org/10.1085/jgp.49.1.69

Slayman, C. L. (1965b). Electrical properties of Neurospora crassa. Respiration and the intracellular potential. J. Gen. Physiol., 49, 93.

Slayman, C. L., & Gradmann, D. (1975). Electrogenic proton transport in the plasma membrane of Neurospora. Biophysical Journal, 15, 968–971. https://doi.org/10.1016/S0006-3495(75)85877-2

Slayman, C. L., Long, W. S., & Lu, C. Y. (1973). The relationship between ATP and an electroneutral pump in the plasma membrane of Neurospora crassa. The Journal of Membrane Biology, 14, 305–338. https://doi.org/10.1007/BF01868083

Slayman, C. L., Lu, C. Y., & Shane, L. (1970). Correlated changes in membrane potential and ATP concentrations in Neurospora. Nature, 226, 274–276. https://doi.org/10.1038/226274a0

Sondergaard, T. E., Schulz, A., & Palmgren, M. G. (2004). Energization of transport processes in plants. Roles of the plasma membrane H+-ATPase. Plant Physiology, 136, 2475–2482. https://doi.org/10.1104/pp.104.048231

Spanswick, R. M., Stolarek, J., & Williams, E. J. (1967). The membrane potential of Nitella translucens. Journal of Experimental Botany, 18, 1–16. https://doi.org/10.1093/jxb/18.1.1

Svennelid, F., Olsson, A., Piotrowski, M., Rosenquist, M., Ottman, C., Larsson, C., ... Sommarin, M. (1999). Phosphorylation of Thr-948 at the C terminus of the plasma membrane H+-ATPase creates a binding site for the regulatory 14-3-3 protein. Plant Cell, 11, 2379–2391.

Vara, F., & Serrano, R. (1982). Partial purification and properties of the proton-translocating ATPase of plant plasma membranes. The Journal of Biological Chemistry, 257, 12826–12830.

Vara, F., & Serrano, R. (1983). Phosphorylated intermediate of the ATPase of plant plasma membranes. The Journal of Biological Chemistry, 258, 5334–5336.

Villalba, J. M., Palmgren, M. G., Berberian, G. E., Ferguson, C., & Serrano, R. (1992). Functional expression of plant plasma membrane H+-ATPase in yeast endoplasmic reticulum. The Journal of Biological Chemistry, 267, 12341–12349.

Villalobo, A., Boulter, M., & Goffeau, A. (1981). Electrogenic proton translocation coupled to ATP hydrolysis by the plasma membrane Mg2+-dependent ATPase of yeast in reconstituted proteoliposomes. The Journal of Biological Chemistry, 256, 12081–12087.

Willsky, G. R. (1979). Characterization of the plasma membrane Mg2+-ATPase from the yeast, Saccharomyces cerevisiae. The Journal of Biological Chemistry, 254, 3326–3332.

How to cite this article: Palmgren M, Morsomme P. The plasma membrane H+-ATPase, a simple polypeptide with a long history. Yeast. 2019;36:201–210. https://doi.org/10.1002/yea.3365