Chapter 3

Basic amino acid transport in plasma membrane vesicles of *Penicillium chrysogenum*


**SUMMARY**

The characteristics of the basic amino acid permease (system VI) of the filamentous fungus *Penicillium chrysogenum* were studied in plasma membranes fused with liposomes containing the beef heart mitochondrial cytochrome-c oxidase. In the presence of reduced cytochrome c, the hybrid membranes accumulated the basic amino acids arginine and lysine. Inhibition studies with analogs revealed a narrow substrate specificity. Within the external pH-range of 5.5-7.5, the transmembrane electrical potential ($\Delta\psi$) functions as the main driving force for uphill transport of arginine, although a low level of uptake was observed when only a transmembrane pH gradient was present. It is concluded that the basic amino acid permease is a H⁺-symporter. Quantitative analysis of the steady-state levels of arginine uptake in relation to the proton motive force suggest a H⁺-arginine symport stoichiometry of one-to-one. Efflux studies demonstrated that the basic amino acid permease functions in a reversible manner.
Amino acids are utilized by fungi as primary or secondary nitrogen sources or as building blocks for the synthesis of proteins and peptides. Systems involved in the translocation of amino acids across the plasma membrane have been studied in only a few filamentous fungi [119, 132, 307]. From these, *Neurospora crassa*, *Aspergillus nidulans*, and to a lesser extent *Penicillium chrysogenum* are genetically and biochemically the most extensively characterized species. Two distinct classes of plasma membrane located amino acid permeases are found in filamentous fungi: systems that catalyze the uptake of structurally related amino acids with a broad substrate specificity, like the general amino acid permeases of plant and animal cells [51, 172, 198], and systems with a narrow substrate specificity like bacterial amino acid transporters [229].

Fungi show a peculiar substrate specificity of their amino acid transport systems, and multiple transport mechanisms seem to exist for several amino acids. In *N. crassa*, five distinct transport systems have been identified, with specificity for aromatic and aliphatic amino acids (system I), aromatic, aliphatic and basic amino acids (system II), basic amino acids (system III), acidic amino acids (system IV), and L-methionine (system V) [119, 132, 218, 307]. Studies with mycelium of *P. chrysogenum* indicate that this fungus possesses at least six distinct amino acid transport systems [29, 119, 132, 135, 136, 304]. Most of these systems are specific for one amino acid and analogues except for system III which is a general amino acid permease, and system IV which transports acidic amino acids only [132, 135]. Amino acid transporters of fungi are assumed to possess some typical properties: i) they seem to function unidirectional; i.e., only uphill transport is observed while efflux or countertransport of the accumulated amino acids is not detected [132, 216]; ii) their activity is regulated by transinhibition, *i.e.*, a high internal concentrations of an amino acid appears to lower the activity of the transport system and thereby inhibits further uptake [132, 136, 218]. Transinhibition results in a decrease of the maximal transport rate \( V_{\text{max}} \) without affecting the affinity \( K_m \) of the system. Transinhibition is thought to occur through binding of the amino acid at a regulation site on the transport system that faces the cytosol. Binding would result in inactivation or a reduction of the transport activity. This phenomenon would allow a regulation of the intracellular amino acid pools and prevent the cell from reaching deleterious cytosolic amino acid concentrations.

Amino acid transport systems from filamentous fungi have been studied mainly in mycelial suspensions. Factors like growth phase, medium composition and stage of development strongly influence the expression, regulation and properties of these systems [29, 70, 132, 239, 304, 307]. In *P. chrysogenum* the transport systems for L-
arginine and L-lysine (system VI), and L-cysteine (system IX) are expressed constitutively. Transport systems specific for methionine (system I) and cystine (system II) are expressed under sulphur starvation, while carbon or nitrogen starvation results in the expression of the transport systems for neutral and basic amino acids (system III) and acidic amino acids (system IV) [29, 132].

Factors like metabolism and compartmentalization interfere with the analysis of the plasma membrane transport processes when performed with intact mycelium. Internal amino acid pools are sequestered in at least two compartments, the cytosol with a high turnover rate, and the vacuole with a low turnover rate [119]. To study plasma membrane located transport systems in *P. chrysogenum*, we have developed a model system devoid of metabolic activities by fusing membrane vesicles with liposomes containing the mitochondrial beef heart cytochrome-c oxidase [82, 127]. We have used this hybrid system to analyze the characteristics of the constitutive basic amino acid transport system *in vitro* [135, 217]. Several properties of this transport system which were difficult to access in mycelial suspension could be well characterized in this hybrid system. Our data demonstrate that the basic amino acid permease is reversible proton symporter with a narrow substrate specificity.

**EXPERIMENTAL PROCEDURES**

**Organisms and culture conditions**

*P. chrysogenum* strains Wisconsin 54-1255 and Panlabs P2 (kindly supplied by Gist-brocades NV) were grown on production medium (pH 6.3) as described by Lara *et al.* (1982) supplemented with 10 mM glutamate and 10 % (mass/vol.) glucose. Cultures were incubated for approximately 70 h in a rotary shaker at 200 rpm and 25 °C. The Wisconsin 54-1255 strain was previously cultured for 24 h on production medium, with the omission of phenylacetic acid and lactose, containing 16% (mass/vol.) glucose. The P2 strain was previously cultured on YPG medium [1 % (mass/vol.) Yeast extract, 2 % (mass/vol.) Peptone and 2 % (mass/vol.) Glucose] for 72 h at pH 7.

**Plasma membrane isolation and fusion**

Bovine heart mitochondria were obtained according to the procedure described by King (1967). Cytochrome-c oxidase was isolated from these mitochondria as described by Yu *et al.* (1975), suspended in 50 mM sodium phosphate (pH 7.5) containing 1.5 % (mass/vol.) cholic acid and stored in liquid nitrogen. Cytochrome-c oxidase was reconstituted in liposomes, composed of 75 % (by mass) acetone/ether
washed *Escherichia coli* lipids and 25 % (by mass) Egg Yolk L-phosphatidylcholine, at a protein/lipid ratio of 0.16 nmol heme a/mg lipid [82]. *P. chrysogenum* plasma membranes were isolated according the procedure described by Hillenga *et al.* (1994). Cytochrome-c-oxidase-containing liposomes (10 mg lipid) and plasma membranes (1 mg protein) were mixed, rapidly frozen in liquid nitrogen and thawed slowly at 21 °C [83]. The freeze-thaw step was repeated once, and hybrid membranes were sized with a small-volume extrusion apparatus (Avestin Inc., Ottawa, Canada) [184] using polycarbonate filters (Avestin) with pore sizes of 400 nm and 200 nm. Fused membranes had a protein/lipid ratio of approximately 0.08-0.09 (mass/mass) (relative to phospholipid).

**Electrical and pH gradients across the membrane**

The transmembrane electrical potential (ΔΨ, interior negative) was calculated from the distribution of the tetraphenylphosphonium ion (TPP+), assuming concentration dependent binding to the membranes as described [178]. The external concentration of TPP+ was determined with a TPP+-selective electrode. Cytochrome-c oxidase vesicles (corresponding to 0.23 nmol cytochrome-c oxidase) were added to 50 mM potassium phosphate of the indicated pH, containing 5 mM MgSO₄ and 2 µM TPP+. A proton-motive force (Δp) was generated by the addition of ascorbate (10 mM, adjusted to the desired pH), N,N,N',N'-tetramethyl-p-phenylenediamine (Ph(NMe₂)₂; 200 µM) and horse heart cytochrome c (20 µM). When indicated, the ionophores nigericin and valinomycin were used at concentrations of 10 nM and 100 nM, respectively. The pH gradient across the membrane (ΔpH, interior alkaline) was determined from the fluorescence of pyranine (excitation, 450 nm; emission, 508 nm), measured with a Perkin Elmer LS50B luminescence spectrophotometer. Pyranine (100 µM) was entrapped in proteoliposomes by freeze-thaw-extrusion [82]. External pyranine was removed with a Sephadex G-25 column (coarse, 1/20 cm). Valinomycin was added to a final concentration of 50 nM. A Δp was generated by addition of the electrondonor system as described for ΔΨ-measurements. When indicated, nigericin was added to a final concentration of 1 µM.

**Transport studies**

Uptake of arginine and lysine was studied at 25 °C and pH 6.5, unless stated otherwise. Mycelium was suspended in 50 mM potassium phosphate at final densities of 10 mg/ml (P2) or 6 mg/ml (Wisconsin-54-1255), and stored on ice until further use. L-[U-¹⁴C]-arginine (Amersham, 38 Ci/mol) or L-[U-¹⁴C]-lysine (Amersham, 43 Ci/mol), previously 10-fold diluted with non-labeled substrate, were added to the mycelial suspension to 30 µM unless indicated otherwise. At given time intervals,
samples of 0.5 ml were taken, added to 2 ml of ice cold 0.1 M LiCl, and filtered immediately on paper filters (296 PE, type 0860; Schleicher & Schuell). Filters were washed once with 2 ml ice cold 0.1 M LiCl, and the amount of radioactivity was determined with a liquid scintillation counter (Packard Tri-Carb 460 CD; Packard Instruments). Cells were de-energized by preincubation with the protonophore carbonyl-cyanide-m-chlorophenylhydrazone (CF$_3$O$_2$Ph$_2$C(CN)$_2$, 10 µM) for 5 min at 25 °C. For uptake studies with hybrid membranes, vesicles were suspended to a final concentration of approximately 1.2 mg protein/ml in 50 mM potassium phosphate (pH 6.5, unless indicated otherwise) containing 5 mM MgSO$_4$. After 1-min incubation in the presence of the electron donor system ascorbate (30 mM), Ph(NMe)$_2$$_2$ (150 µM) and horse heart cytochrome c (7.5 µM), L-[U-$^{14}$C]-amino acids were added to 30 µM unless indicated otherwise. Samples of 20 µl were taken at given time intervals and processed as described above. Samples were filtered on 0.45-µm pore-size diameter cellulose-nitrate filters (Schleicher and Schuell). For efflux studies hybrid membrane vesicles were washed twice with a 20-fold volume of 50 mM potassium phosphate, pH 6.5. Concentrated suspensions of 25-30 mg protein/ml in 50 mM potassium phosphate (pH 6.5) containing 5 mM MgSO$_4$, CF$_3$O$_2$Ph$_2$C(CN)$_2$ (10 µM) and arginine at the indicated concentration, were supplemented with a tracer of L-[U-$^{3}$H]-arginine (Amersham, 4 Ci/mmol) and incubated for 3 h at 25 °C. Samples of 8 µl were rapidly diluted into 400 µl 50 mM potassium phosphate (pH 6.5) containing 5 mM MgSO$_4$ and CF$_3$O$_2$Ph$_2$C(CN)$_2$ (10 µM), filtered on 0.45-µm pore-size diameter cellulose-nitrate filters (Schleicher and Schuell), washed once with 2 ml ice cold 0.1 M LiCl and processed as described above. Kinetic data were analyzed with the GraFit program (Erithacus Software Ltd.).

Other methods

Protein concentrations were determined in the presence of 0.5 % (wt/vol) SDS using a modified Lowry assay [289]. Bovine serum albumin was used as a standard.

RESULTS

Kinetics of arginine and lysine transport

The constitutive basic amino acid permease was studied in plasma membranes obtained from the *P. chrysogenum* strains Wisconsin 54-1255 and P2. Plasma membranes were fused with cytochrome-c oxidase containing liposomes by a freeze/thaw-extrusion technique. To establish whether membrane isolation or fusion affected the basic amino acid permease, the kinetics of arginine and lysine uptake were
determined both in hybrid membranes and mycelial suspensions. Compared to mycelium, the specific activity of arginine and lysine uptake was clearly increased in hybrid membranes, \( i.e. \), a \( V_{\text{max}} \) of 0.63 and 4 nmol/min·mg of protein, respectively (Fig. 3.1). Arginine was accumulated to a higher extent than lysine by the hybrid membranes. An increase in apparent \( K_m \) value of arginine uptake was noted in the hybrid membranes as compared to intact mycelium, \( i.e. \), 120 and 25 µM, respectively. Similar results were obtained with mycelia and hybrid membranes of strain P2, yielding \( V_{\text{max}} \) values of 0.45 and 2.2 nmol/min·mg of protein, and \( K_m \) values of 30 and 170 µM, respectively (data not shown). These studies show that the basic amino acid transport system is active in the hybrid membranes. The proton motive force generated by the beef heart cytochrome-c oxidase functions as a driving force for the accumulation of arginine and lysine.

**Fig. 3.1 Kinetics of L-arginine and L-lysine uptake in mycelium and hybrid membranes.**

(A) Uptake of L-arginine (○,●) and L-lysine (▲,▼) in mycelium (open symbols) and hybrid membranes (closed symbols) from the Wisconsin 54-1255 strain; (B) Kinetics of arginine uptake in mycelium (○) and hybrid membranes (●) from the Wisconsin 54-1255 strain. The error bars indicate the standard error of the mean of three independent experiments.

**Specificity of the arginine/lysine permease**

The specificity of the basic amino acid permease was analyzed by competition experiments. Addition of a large excess of lysine completely inhibited arginine uptake and vice versa in hybrid plasma membranes from the Wis 54-1255 strain. Based on the observed initial velocity of arginine uptake at three L-[U-\( ^{14} \)C]-arginine concentrations (30, 100 and 300 µM) and various concentrations of unlabeled lysine (0-3 mM; Fig. 3.2), a \( K_i \) of 190 µM was calculated for the inhibition of arginine transport by lysine.
Fig. 3.2 Competitive inhibition of L-arginine uptake by L-lysine in hybrid membranes. Inhibition of L-arginine uptake by L-lysine was determined in hybrid membranes of Wisconsin 54-1255 at L-[U-¹⁴C]-arginine concentrations of 30 ( ), 100 ( ) and 300 ( ) µM. The error bars indicate the standard error of the mean of three independent experiments.

Table 3.1. Inhibition of L-arginine transport in hybrid Wisconsin 54-1255 membranes by different L-amino acids.

<table>
<thead>
<tr>
<th>Inhibitor</th>
<th>% Inhibition of L-arginine uptakea</th>
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<tr>
<td>Arginine</td>
<td>100</td>
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<tr>
<td>Lysine</td>
<td>72</td>
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<tr>
<td>Canavanine</td>
<td>89</td>
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<tr>
<td>Ornithine</td>
<td>87</td>
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<tr>
<td>Homoarginine</td>
<td>82</td>
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<tr>
<td>Citrulline</td>
<td>17</td>
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<tr>
<td>Histidine</td>
<td>24</td>
</tr>
<tr>
<td>2,4-Diaminobutyric acid</td>
<td>55</td>
</tr>
<tr>
<td>2-Amino-3-guanidinopropionic acid</td>
<td>29</td>
</tr>
<tr>
<td>Glutamate</td>
<td>3</td>
</tr>
<tr>
<td>Glutamine</td>
<td>42</td>
</tr>
<tr>
<td>Cysteine</td>
<td>8</td>
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<tr>
<td>Serine</td>
<td>12</td>
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<td>Valine</td>
<td>6</td>
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<tr>
<td>Leucine</td>
<td>14</td>
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*aInhibitors were added at a 33-fold higher concentration than [¹⁴C]arginine that was used at a concentration used was 30 µM. Inhibition by arginine was set at 100%.
The observation that the $K_i$ for lysine is in the same range as the $K_m$ value for arginine uptake suggests that these amino acids are equivalent substrates for the basic amino acid permease.

The specificity of the basic amino acid permease was further assessed from the extent of inhibition of arginine uptake by addition of a 33-fold excess of analogs and structurally non related amino acids (Table 3.1). The initial rate of $\Delta p$ driven arginine uptake was strongly inhibited by lysine, canavanine, ornithine and homoarginine. Of the other amino acids only 2,4-diaminobutyric acid and glutamine inhibited the uptake of arginine slightly, while typical substrates for the general amino acid permease, like leucine and valine, were without effect. The observed inhibition pattern indicates that the specificity of this permease is mainly determined by the presence and structural configuration of the $\omega$-amine-group. These observations are consistent with the narrow substrate specificity observed in mycelial suspensions for system VI, the arginine/lysine permease [135, 217].

**Effect of ionophores on L-arginine uptake.** Uptake of L-arginine in P2 hybrid membranes under energized conditions (●), and after previous incubation with $\text{CF}_3\text{OPh}_2\text{C}(\text{CN})_2$ (○), nigericin(Δ, panel A), valinomycin (■, panel B) or nigericin + valinomycin (□, panel C). Congruent closed symbols depict the effect of ionophores added after 5 min of arginine accumulation.

**Effect of ionophores on arginine uptake**

The energetic mechanism of arginine uptake in hybrid membrane vesicles from the P2 strain was studied in detail at pH 6.5 by addition of the ionophores nigericin and valinomycin (Fig. 3.3). The ionophore nigericin collapses the $\Delta p\text{H}$ by mediating electroneutral $K^+\text{-H}^+$ exchange. Upon a collapse of the $\Delta p\text{H}$ by nigericin a
compensatory increase of the $\Delta \psi$ occurs. Addition of nigericin enhanced the uptake of arginine (Fig. 3.3A). The ionophore valinomycin mediates electrogenic diffusion of $K^+$, thereby collapsing the $\Delta \psi$. Omitting the $\Delta \psi$ results in a slight increase in the $\Delta p\text{H}$. Previous incubation with valinomycin strongly, but not completely, reduced arginine uptake (Fig. 3.3B). Addition of valinomycin after 5 min of arginine accumulation resulted in a partial efflux of accumulated arginine. No uptake of arginine was observed after collapsing the $\Delta p$ with nigericin and valinomycin. Addition of these ionophores after 5 min of arginine accumulation caused a fast and almost complete efflux of the accumulated arginine (Fig. 3.3C). These results indicate that the $\Delta \psi$ is the major driving force for arginine uptake. The observation that arginine is still accumulated when only a $\Delta p\text{H}$ is present suggests that the basic amino acid permease is a proton symporter. Similar results were obtained in hybrid membranes of the Wisconsin 54-1255 strain.

**Fig. 3.4 Effect of the external pH on L-arginine uptake.** L-Arginine uptake in hybrid membranes of Wisconsin 54-1255 at pH 5.5 (A), 6.5 (B) and 7.5 (C). Uptake of arginine was determined under energized conditions (▪), and after previous incubation with CF$_3$OPh$_2$C(CN)$_2$ ( ○), valinomycin ( △) and nigericin ( ▽).

**pH dependency of the arginine/$H^+$ symport stoichiometry**

The role of the external pH in arginine uptake was studied in hybrid membranes from the Wisconsin 54-1255 strain (Fig. 3.4). Steady state uptake levels of arginine were similar at pH 5.5 and 6.5, while at pH 7.5 the level of arginine accumulation
Fig. 3.5. Relation between the external pH and the H⁺/Arginine symport stoichiometry. 
(A) Effect of the external pH on the internal pH (Δ) and magnitude and composition of the generated Δp in hybrid membranes of Wisconsin 54-1255; Δψ (●), -ZΔpH (○) and the calculated total Δp (△). (B) Effect of the external pH on the steady state accumulation levels of L-arginine (▼) and the H⁺/Arginine symport stoichiometry (▲). The error bars indicate the standard error of the mean of three independent experiments.

decreased about 2-fold. At all pH values studied, addition of nigericin enhanced arginine uptake while valinomycin strongly reduced steady state levels of arginine uptake.

To determine the H⁺/arginine stoichiometry, the magnitude and composition of the generated Δp was determined at different pH values (Fig. 3.5A). The Δψ and ΔpH were determined in the absence of ionophores. Δψ was highest between pH 6.0-7.0, with maximum values of about -100 mV. ΔpH was maximal at pH 5.5 and slowly decreased at pH values above 6.0. The Δp, calculated from the Δψ and ΔpH values, showed an optimum at around pH 6.0 (about -150 mV). Within the pH-range studied, the internal pH increased with the external pH (Fig. 3.5A), but remained within a range of pH 6.0 to 7.2. Steady state levels of arginine accumulation were determined under identical conditions as Δψ and ΔpH values (Fig. 3.5B). From the data depicted in Fig. 3.5A and the steady state levels of arginine accumulation the apparent H⁺/arginine stoichiometry (n_app) was calculated. The n_app varied between 0.7 and 0.9 within the examined pH range when it is assumed that arginine is accumulated as a positively charged amino acid (Fig. 3.5B). This slight variance indicates that there is no strong dependency of the n_app on the external or internal pH. Arginine uptake in hybrid membranes mainly relates to the magnitude and composition of the Δp. These data show that the basic amino acid permease takes up amino acids in symport with one
proton and effectively translocates two positive charges. This is consistent with the observation that the $\Delta\psi$ is the main driving force in amino acid accumulation via the basic amino acid permease.

**Fig. 3.6 L-Arginine efflux in hybrid membranes.** (A) Uptake of L-arginine in hybrid membranes of Wisconsin 54-1255 under energized conditions (○) or after previous incubation with CF$_3$OPh$_2$C(CN)$_2$ (□). U-[$^{14}$C]-L-arginine efflux or exchange was induced after 5 min of arginine accumulation by the addition of CF$_3$OPh$_2$C(CN)$_2$ (△) or a 100-fold excess (3 mM) of nonlabeled L-arginine (Δ). (B) Efflux of L-arginine in the presence of CF$_3$OPh$_2$C(CN)$_2$. Arginine was used at an internal concentrations of 0.25 (○), 0.5 (●), 1 (▲), 5 (▲) and 10 mM (▼) mixed with a fixed amount of the tracer [$^{14}$C]-arginine. The data is plotted as the amount of [$^{14}$C]-arginine (in cpm) present in the vesicle lumen as a function of time after 50-fold dilution of the membranes. (C) Kinetics of L-arginine efflux: $K_m$ 2-6 mM, $V_{max}$ 1-4 nmol arginine/mg of protein•min. The error bars indicate the standard error of the mean of two independent experiments.

**Arginine efflux and exchange**

Counterflow experiments were conducted to assess whether the basic amino acid permease mediates homologous exchange. Hybrid membranes were equilibrated with 0.5-10 mM L-arginine and diluted 50-fold into a buffer containing [$^{14}$C]arginine (33 µM). Under these conditions no significant uptake of arginine was detected (data not shown). In another experiment, hybrid membrane vesicles were first allowed to accumulate [$^{14}$C]-labeled arginine. Subsequent addition of the protonophore CF$_3$OPh$_2$C(CN)$_2$ caused a fast and almost complete efflux of arginine (Fig. 3.6A), while the addition of an excess of unlabeled arginine resulted in a slow and only partial release of the internal [$^{14}$C]-labeled arginine. Extensive studies showed that the efflux induced by the addition of CF$_3$OPh$_2$C(CN)$_2$ was not enhanced when an excess of unlabeled arginine was added simultaneously (data not shown).

Efflux of arginine occurred when the $\Delta\rho$ or $\Delta$pH was dissipated by ionophores or protonophores (Fig. 3.1 and 3.6A). To further establish that the basic amino acid
permease mediates efflux, this process was studied in more detail. In these experiments the protonophore CF₂OPh₂C(CN)₂ was added to prevent the build up of a Δp during efflux. Hybrid membranes were equilibrated with 0.05-10 mM arginine supplemented with a fixed amount of tracer [³²H]arginine, and diluted 50-fold to initiate efflux. The rate by which the radiolabel is released decreased with increasing internal arginine concentrations (Fig. 3.6B), and when converted to true initial rates, the efflux followed Michaelis Menten kinetics (Fig. 3.6C). The $K_m$ of the efflux process showed to be 2-6 mM and the maximum rate of efflux was determined at 2-4 nmol/min.mg protein. Thus the $K_m$ values of arginine efflux and Δp-driven arginine uptake differ substantially.

**DISCUSSION**

The filamentous fungus *P. chrysogenum* is used for the commercial production of penicillins. To study transport processes that play a role in penicillin biosynthesis, a procedure was developed for the isolation of plasma membranes from *P. chrysogenum* [127]. Hybrid membrane vesicles were obtained by fusing plasma membranes with cytochrome-c oxidase vesicles. These vesicles are endowed with a low ion permeability and can easily be energized by the external addition of reduced cytochrome c. This system was used to study in detail the kinetic and energetic properties of the constitutive arginine/lysine permease, (system VI).

Based on several observations, it was concluded that the transport activity described in this paper did not result from the general amino acid permease (system III, GAP). First, *P. chrysogenum* was grown in the presence of excess lactose and ammonium, the main carbon- and nitrogen sources, respectively. These compounds were not exhausted at the time the cells were harvested, and based on the presence of substantial amounts of intracellular glycogen and trehalose, cells were not carbon-deprived [Hillenga et al., 1995 unpublished results]. These are conditions at which GAP is not expressed [136, 217]. Second, both cells and hybrid membranes hardly showed any uptake of leucine and methionine, both typical substrates for the GAP [Hillenga et al., 1995 unpublished results]. Third, the narrow substrate specificity of the system described in this paper (Table 1) agrees with the specificity described for the arginine/lysine permease (system VI) [135].

In analogy to *Saccharomyces cerevisiae*, a symport stoichiometry of one H⁺ per amino acid was determined for the basic amino acid permease [119, 216]. Uptake was stimulated nor inhibited by Na⁺-ions (up to 50 mM) [Hillenga et al., 1995 unpublished results], excluding the possibility that Na⁺ is a co-transported ion. Within the pH-range studied, the $n_{up}$ varied only slightly indicating that the stoichiometry is not dramatically
affected by the internal nor the external pH. In hybrid membranes the $K_m$ for arginine and lysine uptake was significantly higher than in mycelium. This alteration might have resulted from the applied isolation or fusion procedures. Plasma membranes form a structural integrated part of the cell and isolation inevitably causes the disruption of this organelle and the attached cytoskeleton. It is commonly assumed that these events do not effect properties of transport systems unless important constituents (e.g. binding proteins) or co-factors are lost during the isolation. Some earlier reports implied amino acid-binding proteins in the uptake of amino acids by filamentous fungi [119]. However, no evidence to support these assumptions (e.g. cloned genes) has been presented since, whereas these studies clearly demonstrate that insofar as such binding proteins are involved, they cannot be soluble constituents. Several studies have shown that alterations in membrane composition strongly affected transport systems. For instance, changes in lipid headgroup, acyl chain carbon number and sterol concentration had a pronounced effect on the velocity of L-leucine uptake in *Lactococcus lactis* [140]. However, in those studies no alteration of the $K_m$ was noted. Since the kinetic parameters in intact mycelium are less well defined than in a vesicle system, the discrepancy may arise from the fact that mycelium, in particular that of the P2 strain, is inhomogeneous, while the presence of an anionic cell wall, with a variable thickness, may interfere with the bulk diffusion of cationic solutes like arginine.

Efflux studies of the basic amino acids in membrane vesicles clearly demonstrated that this symport system functions in a reversible manner. An apparent asymmetry is seen in the half-saturation parameters $K_m$ and not in the $V_{max}$ terms. The $K_m$ value for arginine efflux is almost 50-fold higher than the $K_m$ for $\Delta p$ driven arginine uptake. This, however, does not imply that the permease is functionally asymmetric, but rather suggests that the $K_m$ is influenced by the $\Delta p$. The observed linear correlation between the parameters $1/V$ and $1/[S]$ for arginine efflux indicates that in the concentration range studied “transinhibition” by arginine does not occur. Internal concentrations up to 10 mM did not inhibit the efflux process, but at these high internal concentrations no counterflow was observed in mycelium. Since this “transinhibition” phenomenon appears to be specific for intact mycelium, it could be related to metabolism or compartmentalization in the cell, rather than a specific kinetic effect on the permease. Alternatively, the “transinhibition” effect could arise from a regulatory phenomenon which is not operational in isolated plasma membranes.

In conclusion, these studies demonstrated that hybrid membrane vesicles are a powerful tool in investigating characteristics of transport systems from filamentous fungi. Important advantages of hybrid membrane vesicles over mycelial suspension are: i) transport systems can be investigated without the interference of compartmentalization or metabolic activities; ii) the magnitude of driving forces can
be accurately determined; iii) mechanical properties of transport systems can be assessed more adequately; iv) the specificity of a system can be resolved with more certainty.

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