

The light subunit of system b⁰⁺ is fully functional in the absence of the heavy subunit

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The heteromeric amino acid transporters are composed of a type II glycoprotein and a non-glycosylated polytopic membrane protein. System b⁰⁺ exchanges dibasic for neutral amino acids. It is composed of rBAT and b⁰⁺AT, the latter being the polytopic membrane subunit. Mutations in either of them cause malfunction of the system, leading to cystinuria. b⁰⁺AT-reconstituted systems from HeLa or MDCK cells catalysed transport of arginine that was totally dependent on the presence of one of the b⁰⁺ substrates inside the liposomes. rBAT was essential for the cell surface expression of b⁰⁺AT, but it was not required for reconstituted b⁰⁺AT transport activity. No system b⁰⁺ transport was detected in liposomes derived from cells expressing rBAT alone. The reconstituted b⁰⁺AT showed kinetic asymmetry. Expressing the cystinuria-specific mutant A354T of b⁰⁺AT in HeLa cells together with rBAT resulted in defective arginine uptake in whole cells, which was paralleled by the reconstituted b⁰⁺AT activity. Thus, subunit b⁰⁺AT by itself is sufficient to catalyse transmembrane amino acid exchange. The polytopic subunits may also be the catalytic part in other heteromeric transporters.

Keywords: b⁰⁺AT–rBAT exchanger/b⁰⁺AT reconstitution/cystinuria/transport asymmetry/reabsorption

Introduction

The heteromeric amino acid transporters (HATs) are the only known eukaryotic transporters of amino acids composed of two different subunits bound by a disulfide bridge: (i) a type II glycoprotein or heavy subunit (rBAT and 4F2hc), and (ii) a non-glycosylated light subunit with 12 putative transmembrane domains (LAT-1, LAT-2, asc1, y⁺LAT-1, y⁺LAT-2, xCT and b⁰⁺AT) (for reviews see Verrey *et al.*, 1999, 2000; Devés and Boyd, 2000; Chillarón *et al.*, 2001; Wagner *et al.*, 2001). The heavy subunit has an ectodomain that is homologous to bacterial glucosidases (reviewed in Chillarón *et al.*, 2001). The light

subunit is suspected to be the catalytic part of the holotransporter because it is considered to be a polytopic membrane protein and heterodimers of 4F2hc with different light subunits yield different amino acid transport systems: (i) with LAT-1 or LAT-2, variants of system L (Kanai *et al.*, 1998; Mastroberardino *et al.*, 1998; Pineda *et al.*, 1999; Rossier *et al.*, 1999; Segawa *et al.*, 1999), (ii) with asc1, system asc (Fukasawa *et al.*, 2000; Nakauchi *et al.*, 2000), (iii) with y⁺LAT-1 or y⁺LAT-2, variants of system y⁺L (Torrents *et al.*, 1998; Pfeiffer *et al.*, 1999a; Kanai *et al.*, 2000) and (iv) with xCT, system x_c⁻ (Sato *et al.*, 1999, 2000; Bassi *et al.*, 2001).

The rBAT protein is the heavy subunit of b⁰⁺AT in kidney brush border membranes (Fernández *et al.*, 2002), and this heterodimer forms the holotransporter system b⁰⁺ [i.e. exchanger of dibasic amino acids and cystine (influx) for neutral amino acids (efflux)] (Feliubadaló *et al.*, 1999; Chairoungdua *et al.*, 1999; Pfeiffer *et al.*, 1999b; Mizoguchi *et al.*, 2001). The expression of both proteins is needed to express system b⁰⁺ transport at the cell surface (for reviews see Verrey *et al.*, 2000; Chillarón *et al.*, 2001). These two proteins when mutated cause the inherited disease named cystinuria, characterized by hyperexcretion of cystine and dibasic amino acids in urine (for reviews see Palacín *et al.*, 2001a,b): mutations in rBAT cause type I cystinuria (a completely recessive form of the disease) (Calonge *et al.*, 1994), and mutations in b⁰⁺AT cause non-type I cystinuria (in which the heterozygotes show moderate amino acid hyperexcretion) (Feliubadaló *et al.*, 1999; Font *et al.*, 2001). The rBAT cystinuria mutations already studied seem to cause trafficking defects (Chillarón *et al.*, 1997; reviewed in Palacín *et al.*, 2001b), whereas there are no studies demonstrating whether b⁰⁺AT cystinuria mutations cause trafficking defects or inactivate the transporter.

In the present study we reconstituted system b⁰⁺ activity from extracts of cells transfected with b⁰⁺AT. The reconstituted activity is dependent on the expression of a functional b⁰⁺AT protein and the presence of system b⁰⁺ amino acid substrates on the *trans*-side. In contrast, the expression of rBAT is not necessary to reconstitute the transport activity. Kinetic analysis demonstrated asymmetrical interaction of L-leucine with the reconstituted b⁰⁺AT.

Results

Expression of system b⁰⁺ in rBAT/b⁰⁺AT-transfected cells

Transient transfection of human rBAT and b⁰⁺AT resulted in the expression of sodium-independent cystine and arginine uptake in HeLa cells with characteristics of system b⁰⁺ (i.e. almost complete inhibition by L-ornithine, L-leucine and L-alanine, only partial inhibition by D-ornithine,

and no inhibition by L-glutamate) (Figure 1A). This inhibition pattern of cystine transport is identical to that obtained in rBAT-injected oocytes (Bertran *et al.*, 1992). Kinetic analysis showed an apparent K_m of 132 ± 36 and $226 \pm 64 \mu\text{M}$ for cystine and arginine uptake induced in HeLa cells (data not shown). In contrast, transfection of rBAT or b⁰⁺AT alone did not result in the induction of system b⁰⁺ activity (Figure 1A). Similarly, permanent transfection of rBAT and b⁰⁺AT, but not b⁰⁺AT alone, resulted in the expression of sodium-independent cystine and arginine uptake in Madin–Darby canine kidney (MDCK) cells (Figure 1B). This is in full agreement with previous results (Chairoungdua *et al.*, 1999; Feliubadaló *et al.*, 1999; Pfeiffer *et al.*, 1999b; Font *et al.*, 2001) and stresses the need for both subunits to express functional b⁰⁺ transporters at the cell surface. Indeed, expression of rBAT and b⁰⁺AT at the surface of HeLa cells needs cotransfection of both subunits, otherwise each of the proteins remains in an intracellular location (data not shown). The need for b⁰⁺AT for the routing of rBAT to the cell plasma membrane has also been reported in transfected COS cells (Feliubadaló *et al.*, 1999).

Reconstitution of arginine/leucine exchange from extracts of b⁰⁺AT-transfected HeLa cells

Cells transfected with rBAT, b⁰⁺AT or both together were solubilized with detergent and mixed with phospholipids in order to reconstitute amino acid transporters. Uptake experiments using $0.5 \mu\text{M}$ L-[³H]arginine were performed for up to 6 h in the absence or presence of 2 mM leucine in the interior of the reconstituted liposomes (Figure 2A). When leucine was present, the uptake of arginine in liposomes obtained from extracts of cells transfected with b⁰⁺AT alone or together with rBAT revealed an overshoot that peaked at 1 min uptake and decayed thereafter to equilibrium values (3.9 ± 0.1 pmol/mg protein; $n = 11$ preparations) at 6 h (Figure 2A). The uptake in b⁰⁺AT- and rBAT/b⁰⁺AT-reconstituted liposomes at 1 min was, respectively, 4.8 ± 0.5 and 5.4 ± 1.0 -fold over the equilibrium value (i.e. overshoot factor) ($n = 6$ and 3 preparations, respectively, run in triplicate). In contrast, uptake of arginine into liposomes containing leucine and

obtained from rBAT-transfected cells or into liposomes containing no leucine and obtained from cells transfected with rBAT and b⁰⁺AT, or b⁰⁺AT alone, were indistinguishable and reached the equilibrium plateau without showing any overshoot (Figure 2A; and data not shown). Similar results were obtained from extracts of cells transfected with the arginine transport-irrelevant GABA amino acid transporter GAT-1 (data not shown).

The transport rate of $0.5 \mu\text{M}$ arginine, measured in linear conditions, in b⁰⁺AT- and rBAT/b⁰⁺AT-reconstituted liposomes was 10 times higher when 2 mM leucine was present in the interior of the liposome than when no amino acid was present (Figure 2B). The transport rate of arginine in liposomes obtained from cells transfected with GAT-1 or rBAT alone, irrespective of the presence of leucine in the internal medium, was identical to the transport rate in b⁰⁺AT- or rBAT/b⁰⁺AT-reconstituted liposomes when no amino acid was present in the internal medium. These results demonstrate that transfection of b⁰⁺AT alone in HeLa cells is enough to reconstitute the exchange of arginine and leucine in liposomes. Moreover, reconstitution from b⁰⁺AT- or rBAT/b⁰⁺AT-transfected cells yielded the same b⁰⁺ transport rates (Figure 2B).

The expression of rBAT and b⁰⁺AT in HeLa cells and the reconstitution of these proteins into liposomes was followed by western blot analysis using SDS–PAGE (Figure 2C–D). rBAT or b⁰⁺AT was present in the corresponding cell extracts and liposomes only when transfected (Figure 2C). Transfection of rBAT alone resulted in the expression of the immature core-glycosylated form (i.e. endoglucosidase H-sensitive; data not shown) of the protein. This core-glycosylated form has higher electrophoretic mobility than the matured glycosylated form (i.e. endoglucosidase H-resistant; data not shown) that was detected in cells cotransfected with rBAT and b⁰⁺AT and in renal brush borders (Figure 2C). These rBAT bands are similar to those detected during biogenesis of the protein in oocytes (Chillarón *et al.*, 1997). Moreover, the level of expression of rBAT was higher when cotransfected with b⁰⁺AT ($299 \pm 53\%$ of that found when rBAT was transfected alone; $n = 4$), but the expression of b⁰⁺AT seemed to be independent of the expression of rBAT ($83 \pm 5\%$ of expressed b⁰⁺AT when

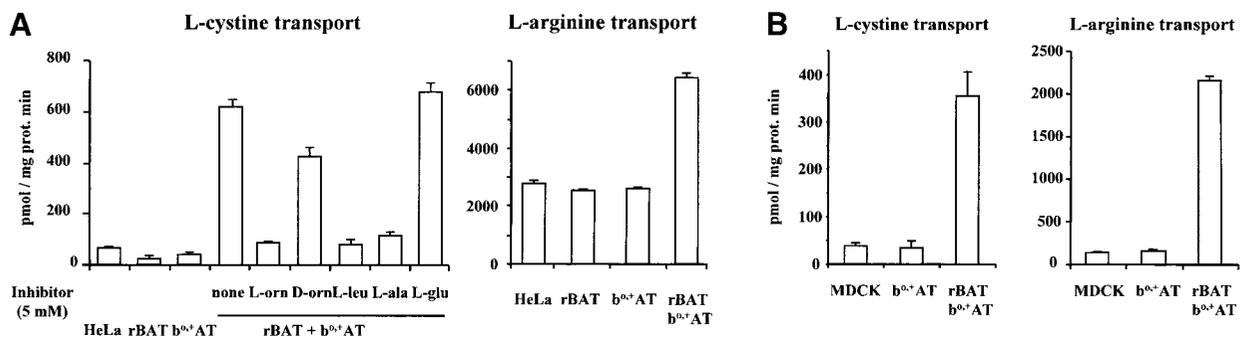


Fig. 1. Expression of amino acid transport in cells cotransfected with rBAT and b⁰⁺AT. **(A)** HeLa cells were transfected with rBAT or b⁰⁺AT alone or together, or were used untransfected (HeLa). Three days later transport of $20 \mu\text{M}$ L-[³⁵S]cystine or $50 \mu\text{M}$ L-[³H]arginine was measured in linear conditions. Only cells cotransfected with rBAT and b⁰⁺AT showed amino acid transport above background. For L-[³⁵S]cystine transport the inhibition pattern with the indicated amino acids at 5 mM was studied. **(B)** MDCK cells were permanently transfected with b⁰⁺AT alone or together with rBAT. The transport of $20 \mu\text{M}$ L-[³⁵S]cystine or $50 \mu\text{M}$ L-[³H]arginine was measured in linear conditions in non-transfected (MDCK) or transfected confluent cells. Only cells cotransfected with rBAT and b⁰⁺AT showed amino acid transport above background.

transfected alone; $n = 4$) (Figure 2C). These results suggest that $b^{o,+}AT$ helps the maturation and probably increases the stability of rBAT. In non-reducing conditions, as expected, bands corresponding to the electrophoretic mobility of the rBAT/ $b^{o,+}AT$ heterodimers were seen only in cell extracts and liposomes (although to a lesser extent) obtained from cotransfected cells, but most of

rBAT and $b^{o,+}AT$ migrated as the monomer (Figure 2D). In all, our results demonstrate that $b^{o,+}AT$ in the absence of rBAT mediates the exchange of arginine and leucine in the reconstituted liposomes.

The cystinuria-specific A354T $b^{o,+}AT$ mutant revealed inactivation of transport in reconstituted systems

To examine whether the exchange of external arginine with internal leucine ($L\text{-arg}_o/L\text{-leu}_i$) is dependent on the reconstitution of a functional $b^{o,+}AT$, two cystinuria-specific $b^{o,+}AT$ mutants (A182T and A354T) were studied. Mutation A182T is one of the most common cystinuria mutations found in *SLC7A9*, whereas mutation A354T has been found in two cystinuria patients (Font *et al.*, 2001; and unpublished result from the International Cystinuria Consortium). We chose these mutations because previous functional analysis showed complete lack of transport activity of A354T, and ~50% residual transport activity of A182T when cotransfected with rBAT in HeLa cells (Font *et al.*, 2001). Identical results were obtained here (Figure 3A). Kinetic analysis showed an apparent K_m of $256 \pm 59 \mu M$ (mean \pm SEM) for arginine in rBAT/A182T $b^{o,+}AT$ -transfected cells (data not shown). This value is similar to that obtained in rBAT/wild-type $b^{o,+}AT$ -transfected cells (see first paragraph of Results). The reconstituted A354T $b^{o,+}AT$ mutant showed <10% of the wild-type $b^{o,+}AT$ transport of arginine *trans*-stimulated by leucine into liposomes (Figure 3B). In contrast, the reconstituted A182T $b^{o,+}AT$ mutant showed complete arginine transport activity like the wild-type $b^{o,+}AT$, which was totally dependent on the presence of leucine in the internal medium (Figure 3B). Western blot analysis revealed the presence of the wild-type and the mutated $b^{o,+}AT$ proteins in the cells and reconstituted systems in similar amounts (Figure 3C). Therefore, the defective transport activity of the A354T mutant could not be

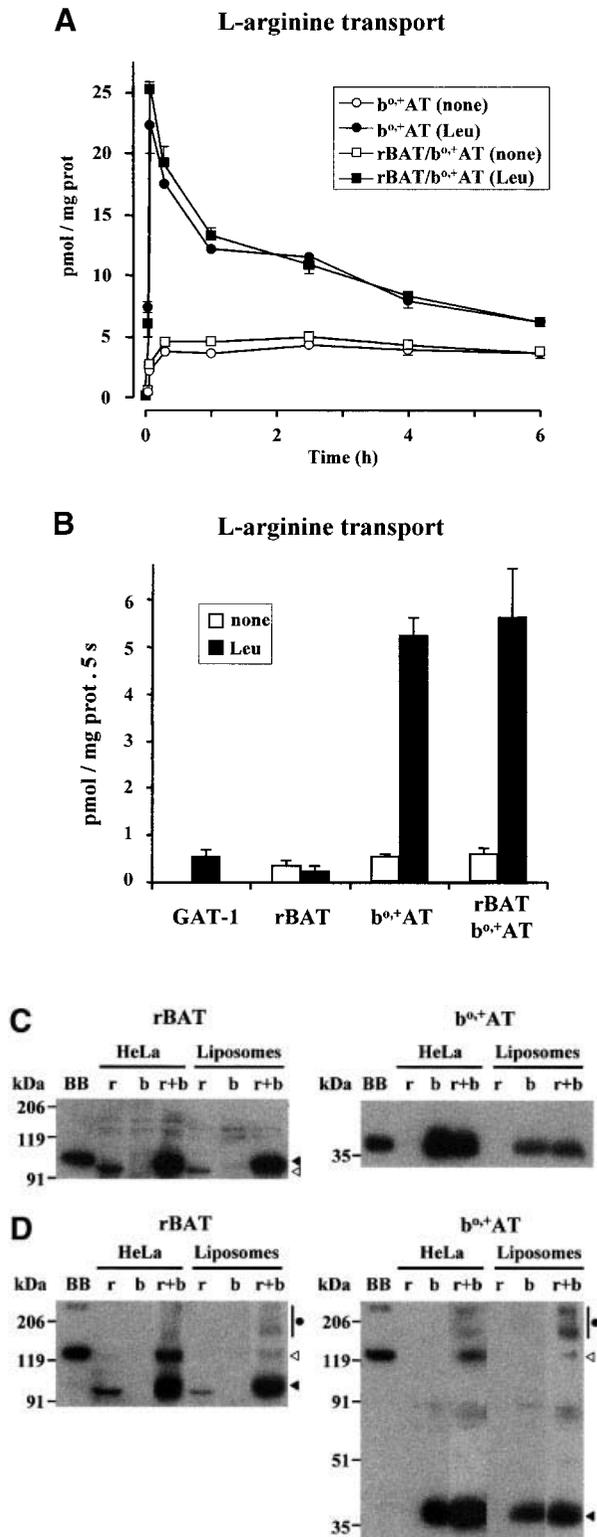


Fig. 2. Transport of arginine in reconstituted systems from HeLa cells. HeLa cells were transfected with rBAT or $b^{o,+}AT$ alone or together (rBAT/ $b^{o,+}AT$), or with the GABA transporter GAT-1. Three days later HeLa extracts were reconstituted into liposomes containing (Leu) or not containing (none) 2 mM leucine in the internal medium. (A) A time course of the transport of $0.5 \mu M$ $L\text{-}[^3H]$ arginine into the indicated liposomes. Transport was measured at 5 s, 1 and 15 min, and at 1, 2.5, 4 and 6 h. Data (mean \pm SEM) correspond to a representative experiment run in triplicate. When not visible, error bars are smaller than the symbol. (B) Initial transport rates of $0.5 \mu M$ $L\text{-}[^3H]$ arginine into the indicated liposomes. Data (mean \pm SEM) correspond to 12, five, five and one experiments run in triplicate for $b^{o,+}AT$, rBAT, rBAT/ $b^{o,+}AT$ and GAT-1 groups, respectively. (C and D) Western blot of rBAT (left panels) and $b^{o,+}AT$ (right panels) in reducing (C) or non-reducing (D) conditions. One hundred micrograms of protein of HeLa cells extracts (HeLa) or liposomes from the experiments shown in (A) and (B) were subjected to SDS-PAGE in the absence (D) or in the presence (C) of 100 mM DTT. Fifty micrograms of protein of human renal brush border membranes (BB) was loaded as control. Lanes in (C) and (D): rBAT-transfected (r), $b^{o,+}AT$ -transfected (b), and rBAT/ $b^{o,+}AT$ -cotransfected (r+b) cells (HeLa) or liposomes. (C) Black arrowhead, rBAT band with mature glycosylation (i.e. endoglucosidase H resistant; data not shown). White arrowhead, rBAT band with core glycosylation (i.e. endoglucosidase H sensitive; data not shown). (D) Black arrowhead, mature and core glycosylation bands of the rBAT monomer (left panel) and $b^{o,+}AT$ monomer (right panel). White arrowhead, rBAT/ $b^{o,+}AT$ heterodimer. Black circle, high molecular weight complexes. For (C) and (D) data correspond to two representative experiments. Two additional experiments gave similar results.

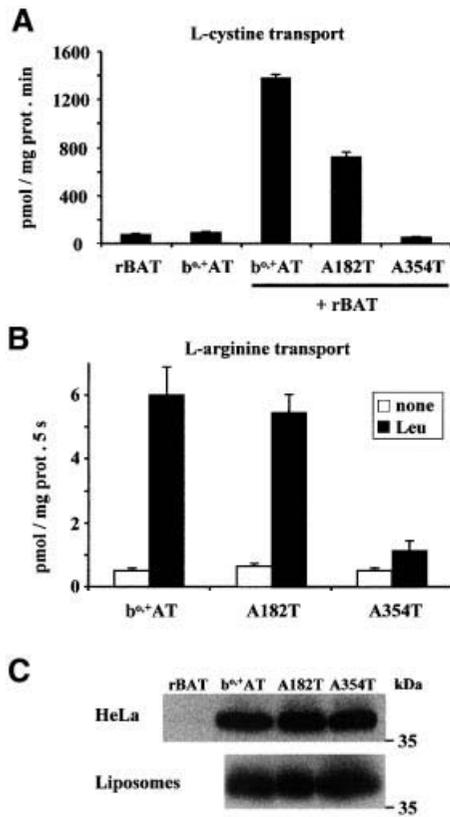


Fig. 3. Transport activity of cystinuria-specific b⁰⁺AT mutants in HeLa cells and reconstituted systems. HeLa cells were transfected with rBAT or b⁰⁺AT alone, or with rBAT plus wild-type b⁰⁺AT, A182T- or A354T-mutated b⁰⁺AT. (A) Three days later transport rates of 20 μ M L-[³⁵S]cystine into cells were measured. Data (mean \pm SEM) from a representative experiment. Three additional experiments gave similar results. (B) Transport of arginine in reconstituted systems. Cells were transfected with wild-type b⁰⁺AT, A182T- or A354T-mutated b⁰⁺AT. Three days later, cells were used for reconstitution into liposomes containing (Leu) or not containing (none) 2 mM leucine in the internal medium, and the transport rates of 0.5 μ M L-[³H]arginine were measured. Transport of arginine *trans*-stimulated by leucine was significant in the three groups. Data (mean \pm SEM) correspond to a representative experiment run in triplicate. A second experiment gave similar results. (C) Western blot of b⁰⁺AT in reconstituted systems. Cell extracts (HeLa) and liposomes (100 μ g protein) from the experiment shown in (B) were used for SDS-PAGE in the presence of 100 mM DTT. As a negative control, extracts from cells transfected with rBAT are shown.

attributed to lower expression of this protein in the liposomes. These results demonstrate that (i) L-arg_o/L-leu_i exchange activity is dependent on the reconstitution of a functional b⁰⁺AT protein, (ii) mutation A354T abolishes b⁰⁺AT transport activity almost completely, and (iii) mutation A182T retains full b⁰⁺AT transport activity.

Reconstituted b⁰⁺AT amino acid transport activity shows substrate specificity and electrogenic activity characteristic of system b⁰⁺

Next, we attempted the reconstitution of system b⁰⁺ transport activity from MDCK cells permanently transfected with b⁰⁺AT alone or together with rBAT. In these reconstituted systems similar results were obtained to those shown from HeLa transfected cells (Figure I in Supplementary data). Thus, b⁰⁺AT-reconstituted liposomes showed an overshoot of 0.5 μ M L-[³H]arginine transport

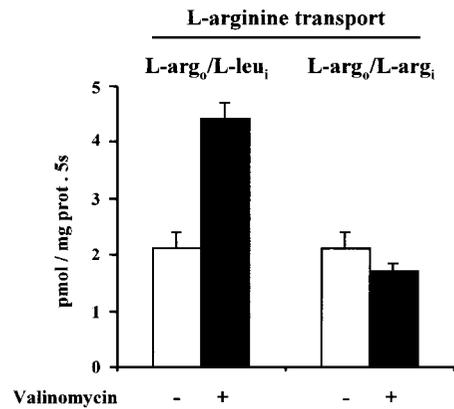


Fig. 4. Effect of membrane potential on the amino acid transport activity elicited by b⁰⁺AT in reconstituted systems. b⁰⁺AT-expressing MDCK cells were used for reconstitution in liposomes. Initial transport rates of 0.5 μ M L-[³H]arginine (L-arg_o) into liposomes in the exchange mode L-arg_o/L-leu_i or L-arg_o/L-arg_i were measured. Liposomes containing no amino acids or 2 mM leucine (L-leu_i) or arginine (L-arg_i) in the internal medium were used. Transport in exchange with the internal amino acid was calculated by subtracting transport in the absence of amino acid in the internal medium from transport in its presence. Valinomycin was added when indicated to liposomes containing 120 mM KP_i in the internal medium to impose a membrane potential (negative inside). Data (mean \pm SEM) correspond to a representative experiment run in triplicate.

generated by 2 mM leucine in the internal medium. The transport at the peak of the overshoot was 4.9 ± 0.7 times (i.e. overshoot factor) higher than the equilibrium values (3.4 ± 0.3 pmol/mg protein) ($n = 7$ preparations). Similar results were obtained from rBAT/b⁰⁺AT-reconstituted liposomes with an average overshoot factor of ~ 6 in two independent experiments. In contrast, reconstitution into liposomes of extracts from untransfected MDCK cells did not show any overshoot of arginine and uptake increased with time approaching equilibrium (data not shown). The transport rate of 0.5 μ M L-[³H]arginine in the presence of 2 mM leucine in the internal medium of b⁰⁺AT- and rBAT/b⁰⁺AT-reconstituted liposomes was 10 times higher than the background transport rates (i.e. transport in liposomes obtained from non-transfected MDCK cells or in b⁰⁺AT- and rBAT/b⁰⁺AT-reconstituted liposomes with no amino acids inside) (Figure I in Supplementary data). Similarly, no differences were observed between b⁰⁺AT- and rBAT/b⁰⁺AT-reconstituted liposomes when transport of L-[³H]arginine was measured at saturating concentrations (5 and 10 μ M) (data not shown).

Western blot analysis showed an identical situation (Figure I in Supplementary data) to that described in the HeLa experiments. (i) rBAT and b⁰⁺AT proteins were expressed only in the liposomes obtained from the corresponding transfected cells. (ii) In non-reducing conditions, heterodimers of rBAT and b⁰⁺AT were visible only in cells expressing rBAT and b⁰⁺AT. (iii) These heterodimers were not detected in cells transfected with b⁰⁺AT alone. These results demonstrate that b⁰⁺AT, which does not form heterodimers with rBAT, can also mediate the exchange of arginine and leucine in liposomes reconstituted from b⁰⁺AT-expressing MDCK cells.

System b⁰⁺ induced by rBAT in *Xenopus* oocytes has been shown to be electrogenic when a neutral and a dibasic

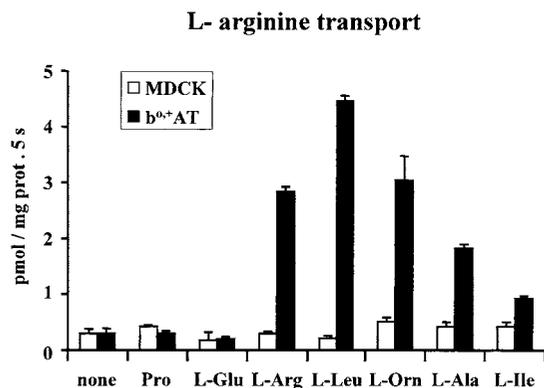


Fig. 5. Trans-stimulation of arginine transport by amino acids in b⁰⁺AT-reconstituted systems. b⁰⁺AT-expressing MDCK cells (filled bars) or non-transfected cells (open bars) were used for reconstitution in liposomes. Initial transport rates of 0.5 μM L-[³H]arginine into the liposomes were measured in the absence (none) or presence of the amino acids indicated at 2 mM concentration inside the liposomes. Data (mean ± SEM) from a representative experiment run in triplicate.

amino acid are exchanged through the transporter (Busch *et al.*, 1994; Coady *et al.*, 1994, 1996; Chillarón *et al.*, 1996). For this reason all the experiments in reconstituted systems in this study were performed with an imposed membrane potential (negative inside) generated by 120 mM K_P_i in the internal medium of the liposomes and the addition of valinomycin. The effect of the membrane potential on the exchange of external arginine for internal leucine (L-arg_o/L-leu_i) or external arginine for internal arginine (L-arg_o/L-arg_i) was tested in b⁰⁺AT-reconstituted liposomes from MDCK transfected cells. As shown in Figure 4, the imposed membrane potential (negative inside) doubled the transport rate of arginine in the L-arg_o/L-leu_i exchange mode, whereas it did not affect the transport rate of arginine in the L-arg_o/L-arg_i exchange mode. This demonstrates that the exchange L-arg_o/L-leu_i in the b⁰⁺AT-reconstituted system depends on the membrane potential, whereas the exchange L-arg_o/L-arg_i is independent, as expected for system b⁰⁺ transport activity.

In order to characterize further the amino acid transport activity of the reconstituted b⁰⁺AT, several amino acids were analysed for their capacity to *trans*-stimulate the transport of arginine. Substrates of the rBAT/b⁰⁺AT-induced system b⁰⁺, such as arginine, ornithine, alanine, leucine or isoleucine, produced *trans*-stimulation of arginine transport in b⁰⁺AT-reconstituted liposomes, whereas proline or glutamate, which are not substrates of system b⁰⁺, did not *trans*-stimulate arginine transport (Figure 5). The level of *trans*-stimulated arginine transport elicited by these system b⁰⁺ substrates parallels the transport of these amino acids induced in cells expressing rBAT and b⁰⁺AT (Chairoungdua *et al.*, 1999; Pfeiffer *et al.*, 1999b). No *trans*-stimulation was detected in liposomes from non-transfected cells (Figure 5).

Asymmetry of b⁰⁺AT

Kinetic analysis of arginine uptake into b⁰⁺AT-reconstituted liposomes revealed a high-affinity (apparent K_m ~0.5 μM) exchange with 2 mM leucine in the internal medium (Figure 6A). Next, we studied the effect of the variation of the leucine concentration in the internal

medium on the exchange with 0.5 μM L-[³H]arginine in the external medium. Internal leucine *trans*-stimulated transport of external arginine with saturation kinetics, and the concentration of internal leucine needed to *trans*-stimulate half-maximally the transport of arginine was estimated to be ~2.5 mM (Figure 6B). In order to check the interaction of leucine with b⁰⁺AT at the external face of the liposomes, the exchange of external 0.5 μM L-[³H]arginine with 2 mM internal leucine was measured at different leucine concentrations in the external medium. External leucine almost abolished the exchange at 50 μM; similarly, 100 μM external cystine also almost abolished the exchange (Figure 6C). To characterize the *cis*-inhibition elicited by leucine, the exchange L-arg_o/L-leu_i was measured at three different external arginine concentrations and five different external leucine concentrations. Dixon plots of these studies showed convergent lines crossing at negative values of the *x*-axis, demonstrating competitive inhibition by external leucine of the L-arg_o/L-leu_i exchange, with an apparent K_i of 1.6 μM. In summary, internal leucine *trans*-stimulated transport of arginine with a semi-maximal concentration in the mM range, whereas external leucine *cis*-inhibited the exchange L-arg_o/L-leu_i with a K_i in the μM range. These results demonstrate that the reconstituted b⁰⁺AT transporter interacts asymmetrically with leucine.

Discussion

In this study we show system b⁰⁺ transport activity in b⁰⁺AT-reconstituted systems from HeLa and MDCK b⁰⁺AT-transfected cells. This transport activity is dependent on (i) the expression of a functional b⁰⁺AT protein in the liposomes, (ii) the presence of system b⁰⁺ amino acid substrates on the *trans*-side and (iii) the membrane potential for electrogenic arginine/leucine exchange. System b⁰⁺ transport activity in b⁰⁺AT-reconstituted systems is independent of the expression of heterodimers of b⁰⁺AT and rBAT. Thus, the b⁰⁺AT subunit, in the absence of its heavy subunit rBAT, folds into an active form that shows full transport activity. In contrast, routing to the plasma membrane of rBAT and b⁰⁺AT (Feliubadaló *et al.*, 1999; this study) and expression of system b⁰⁺ transport in the plasma membrane require the expression of both the heavy and the corresponding light subunit, as demonstrated earlier for this and other HATs (for reviews see Verrey *et al.*, 2000; Chillarón *et al.*, 2001). Indeed, processing of rBAT from the ER to the Golgi (i.e. maturation of the *N*-glycosylation of rBAT) depends on the presence of the light subunit b⁰⁺AT. Therefore we have reconstituted functional b⁰⁺AT from an intracellular location. Moreover, b⁰⁺AT seems to protect rBAT against cellular degradation. In contrast, the level of expression of b⁰⁺AT seems to be independent of the expression of rBAT. Therefore, b⁰⁺AT might act as a specific chaperone for rBAT.

The biogenesis of the rBAT/b⁰⁺AT holotransporter remains unknown, but the present study suggests a working model. First, b⁰⁺AT folds into a functional conformation without the participation of rBAT. Then, b⁰⁺AT recognizes rBAT and favours its folding, and finally the rBAT/b⁰⁺AT heterodimers progress to the plasma membrane. Further research is needed to test this

hypothesis. In any case, this is at odds with the biogenesis of the K⁺-transporting P-type ATPases in eukaryotes (for a review see Geering, 2001). These transporters also have an oligomeric structure with a polytopic catalytic α subunit, and a type II glycoprotein β subunit, which is needed for the enzyme function. In these ATPases the β subunit acts as a chaperone for the corresponding α subunit by facilitating the correct membrane integration and packing, acquisition of functional properties and routing to the cell surface of the α subunit. It is becoming more apparent that some transporters, besides HATs and K⁺-transporting ATPases, require ancillary glycoproteins for their proper plasma membrane expression and current function in eukaryotes (Bruce *et al.*, 1994; Abumrad *et al.*, 1998; Kirk

et al., 2000). Thus, CD147 associates with the lactate transporter MCT1 or MCT4 for the expression of functional transporters at the cell surface (Kirk *et al.*, 2000; Wilson *et al.*, 2002). Reconstitution assays of MCT2 or MCT4 alone would indicate whether the ancillary subunit CD147 is necessary for the functional folding of these lactate transporters like the β subunit for the K⁺-transporting P-type ATPases, or whether it is only necessary for routing to the cell surface like the heavy subunit for HATs.

In addition to the chaperone function in K⁺-transporting P-type ATPases, β subunits also contribute to the intrinsic transport properties of the pump (i.e. they influence the apparent K⁺ affinity) (for a review see Geering, 2001). Interestingly, some cystinuria-specific rBAT mutations, in addition to having trafficking defects, also alter the apparent K_m for some amino acids of the system b⁰⁺ holotransporter (M.Pineda *et al.*, in preparation). This suggests that mutations in the heavy subunit rBAT transmit conformational changes to the catalytic light subunit b⁰⁺AT. Alternatively, rBAT *per se* might affect the kinetic properties of the holotransporter. Interestingly, the activity reconstituted in liposomes from b⁰⁺AT-transfected cells is lower than that expressed by rBAT and b⁰⁺AT at the cell surface. Moreover, arginine transport into HeLa cells expressing the holotransporter rBAT/b⁰⁺AT has an apparent K_m of ~200 μ M. This is at odds with the estimated apparent K_m of 0.5 μ M for arginine transport in b⁰⁺AT-reconstituted systems. This difference could be due to the presence of an unstirred layer surrounding cells not present in liposomes that increases the apparent K_m values. Indeed, this discrepancy in apparent K_m values in cells and liposomes has been reported for other transporters (e.g. GABA, glutamate transporters; Pines *et al.*, 1992). Alternatively, the holotransporter rBAT/b⁰⁺AT might have a lower apparent affinity and a higher transport turnover for amino acids like arginine than the catalytic subunit b⁰⁺AT alone.

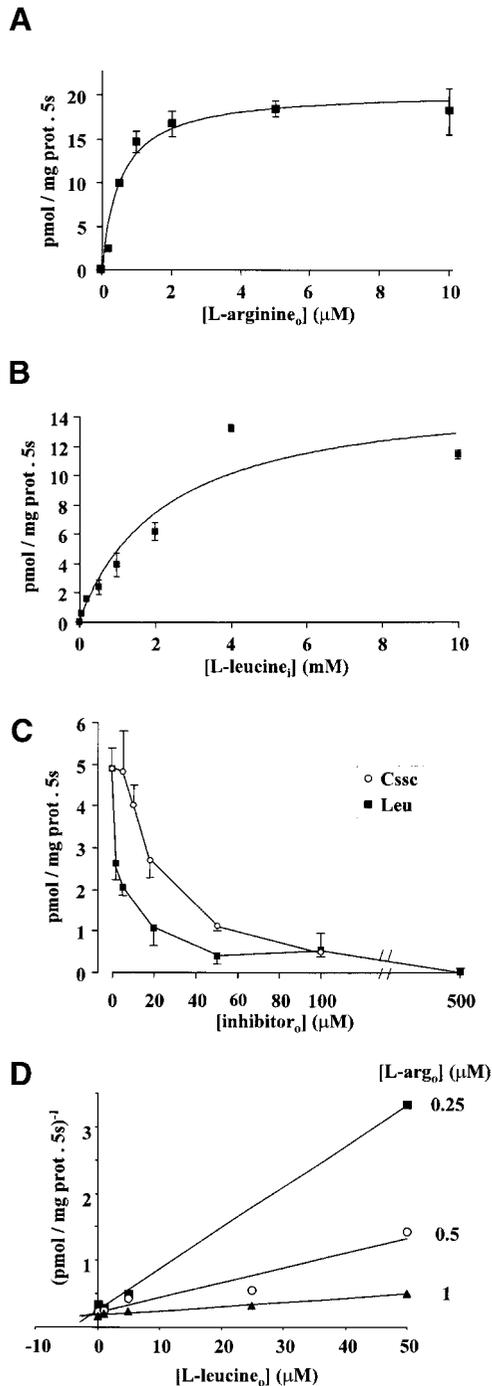


Fig. 6. Kinetic analysis of system b⁰⁺ amino acid transport in b⁰⁺AT-reconstituted liposomes. b⁰⁺AT-expressing MDCK cells were used for reconstitution in liposomes. **(A)** Kinetic analysis of the leucine-*trans*-stimulated L-[³H]arginine transport rates. The arginine transport *trans*-stimulated by leucine was calculated by subtracting transport into liposomes containing no amino acids from the transport into liposomes containing 2 mM leucine. The estimated apparent K_m for arginine outside ([L-arginine]_o) was $0.53 \pm 0.16 \mu\text{M}$ (GraphPad Prism program). Data (mean \pm SEM) correspond to a representative experiment run in triplicate. An additional experiment gave similar results. **(B)** Kinetic analysis of the leucine in the internal medium ([L-leucine]_i) *trans*-stimulating the transport rate of 0.5 μM L-[³H]arginine_o. The estimated apparent K_m for leucine_i was $2.5 \pm 0.6 \text{ mM}$ (GraphPad Prism program). Data (mean \pm SEM) correspond to a representative experiment run in triplicate. An additional experiment gave similar results. **(C)** Inhibition curve of the leucine-*trans*-stimulated arginine_o transport by the indicated concentrations of cystine (Cysc) and leucine (Leu) outside. The 0.5 μM L-[³H]arginine_o transport *trans*-stimulated by 2 mM leucine in the internal medium was calculated as in (A). Data (mean \pm SEM) correspond to a representative experiment run in triplicate. **(D)** Dixon plot of leucine-*trans*-stimulated arginine transport inhibited by different leucine concentrations outside ([L-leucine]_o). The transport of 0.25, 0.5 and 1 μM L-[³H]arginine_o *trans*-stimulated by 2 mM leucine in the internal medium was calculated as in (A). Data (mean \pm SEM) correspond to a representative experiment run in triplicate. In all panels, when not visible, error bars are smaller than the symbol.

Reconstitution of the holotransporter is needed to solve this question.

In this study we were unable to reconstitute the holotransporter rBAT/b⁰⁺AT, for two reasons. (i) Total integrity of the disulfide bound rBAT/b⁰⁺AT heterodimers needs the presence of *N*-ethylmaleimide (NEM) in the homogenization buffer to prevent disulfide bond shuffling (Wang and Tate, 1995). Unfortunately, NEM dramatically inhibits human rBAT/b⁰⁺AT-elicited system b⁰⁺ transport activity, and reconstitution into liposomes of rBAT and b⁰⁺AT in the absence of NEM diminishes the rBAT/b⁰⁺AT heterodimers. (ii) Even in the presence of NEM, there is an excess of b⁰⁺AT expression over that of rBAT in both transient and permanent transfections (data not shown), most probably due to the higher stability of b⁰⁺AT in heterologous expression systems. One way to overcome this situation would be the expression of a b⁰⁺AT–rBAT concatamer, which expresses system b⁰⁺ activity in *Xenopus* oocytes (Pfeiffer *et al.*, 1999b). Reconstitution of this b⁰⁺AT–rBAT concatamer from HeLa transfected cells shows system b⁰⁺ transport activity but the reconstituted activity is very low (i.e. 2-fold over background conditions) (data not shown). Purification before reconstitution will be needed to study the transport characteristics of the concatamer. This may allow the study of the role of rBAT in the holotransporter function.

The reconstituted system b⁰⁺ shows dramatic asymmetry: high-affinity (μ M range) interaction of arginine and leucine from the outside and low-affinity (mM range) interaction of leucine from the inside. Very recently, similar asymmetry of substrate apparent affinities (extracellular apparent K_m in the micromolar range and intracellular apparent K_m in the millimolar range) has been reported for other heteromeric amino acid transporters (LAT-1 and LAT-2) expressed in oocytes (Meier *et al.*, 2002). We do not know the orientation of b⁰⁺AT protein in our reconstituted systems. Detection of the asymmetry for substrate affinity was possible due to the large difference in the substrate apparent affinity on either side of the transporter: since b⁰⁺AT is an exchanger, at very low substrate concentration outside (μ M), only the transporters oriented with the high-affinity site outside would be active and show low-affinity inside. Substrate affinity asymmetry was reported in system b⁰⁺-like exchanger activity from intestinal chicken brush border vesicles (Torrás-Llort *et al.*, 2001): the estimated apparent K_m for arginine was $\sim 8 \mu$ M outside and $\sim 180 \mu$ M inside. This suggests that the human b⁰⁺AT may be oriented with the high-affinity side towards the renal tubule lumen and the low-affinity side inside the cell. This orientation has a physiological significance. System b⁰⁺ is an exchanger for the influx of cystine and dibasic amino acids and the efflux of neutral amino acids (for a review see Chillarón *et al.*, 2001; Palacín *et al.*, 2001b). Immunoprecipitation studies in renal brush border preparations and clinical data on cystinuria strongly suggest that the rBAT/b⁰⁺AT heterodimer is the main, if not the only, transporter responsible for the apical reabsorption of cystine along the renal proximal tubule (Fernández *et al.*, 2002). More than 99% of the cystine and arginine filtered in the glomerulus is reabsorbed in the proximal tubule (Frimpter *et al.*, 1962; Silbernagl, 1988). Then, a high-affinity system would ensure efficient reabsorption of the substrates even at the end of the

proximal tubule. Moreover, low-affinity at the intracellular site of the b⁰⁺ transporter would ensure efficient exchange with the high intracellular concentration of neutral amino acids in the epithelial cells of the proximal tubule.

Mutation in either subunit of the rBAT/b⁰⁺AT holotransporter produces loss of renal reabsorption of cystine and dibasic amino acid that results in cystinuria, and mutations in y⁺LAT-1 (4F2hc/y⁺LAT-1 holotransporter) impair renal reabsorption of dibasic amino acid in lysinuric protein intolerance (LPI) (for a review see Palacín *et al.*, 2001b). Eighteen and seven missense mutations have been described for b⁰⁺AT and y⁺LAT-1 in cystinuria and LPI, respectively (Font *et al.*, 2001; Palacín *et al.*, 2001b). These mutations may cause transport dysfunction by affecting trafficking to the plasma membrane or intrinsic transport activity (Mykkanen *et al.*, 2000). Here we show that reconstitution of the A354T-mutated b⁰⁺AT yields an almost inactive transporter, whereas A182T-mutated b⁰⁺AT retains full reconstituted activity. This demonstrates that, in addition to possible trafficking defects not characterized here, mutation A354T inactivates the transporter, whereas mutation A182T may alter the routing of the holotransporter to the plasma membrane. The degree of the defect caused by these two mutations correlates with the levels of urine cystine and dibasic amino acids in their heterozygotes (i.e. higher hyperexcretion in A354T carriers than in A182T carriers) (Font *et al.*, 2001). Residue A354 is located in putative transmembrane IX and corresponds to a residue with a short side chain (i.e. glycine, alanine or serine) in all known catalytic subunits of HATs. In contrast, residue A182, located in putative transmembrane V, is not conserved among the catalytic subunits of HATs (Font *et al.*, 2001). Further studies are needed to characterize the defects that cause transport inactivation of A354T b⁰⁺AT mutated proteins and the trafficking defect suspected for A182T b⁰⁺AT. Reconstitution of mutated b⁰⁺AT or y⁺LAT-1 proteins may help to dissect the defects associated with cystinuria and LPI mutations.

Materials and methods

Cell culture

MDCK and HeLa cells were grown in Dulbecco's modified Eagle's medium (Gibco) supplemented with 10% fetal calf serum (heat-inactivated), 100 U/ml penicillin (Gibco) and 0.1 mg/ml streptomycin (Gibco) (D10) at 37°C in a humidified atmosphere containing 5% CO₂. Stably transfected MDCK-derived cell lines were grown in D10 supplemented with the appropriate selection agent (400 μ g/ml geneticin and/or 100 μ g/ml hygromycin B).

cDNAs and transfections in HeLa and MDCK cells

Transient transfections were performed in HeLa cells by standard calcium phosphate precipitation in 15 cm diameter plates with a mixture of DNA containing 4 μ g of pEGFP (green fluorescence protein; Clontech), 18 μ g of pCDNA3-rBAT and 18 μ g of pCDNA3-b⁰⁺AT as described (Font *et al.*, 2001). When rBAT, b⁰⁺AT or GAT-1 was transfected alone, the DNA transfection mixture contained 4 μ g of pEGFP and 36 μ g of pCDNA3-rBAT, pCDNA3-b⁰⁺AT or pCDNA3-GAT-1 (wild-type rat GAT-1 cDNA cloned into the pCDNA3 vector (Invitrogen) between the *Hind*III and *Xba*I sites). The GFP-encoding plasmid was included to monitor transfection efficiencies, which ranged from 60 to 90% as assessed by FACS analysis. To obtain a b⁰⁺AT MDCK cell line, parental MDCK-J cells were transfected with pCDNA3-b⁰⁺AT following the standard protocol (Font *et al.*, 2001). Forty-eight hours later, cells were plated at limiting dilution and grown in the presence of geneticin (400 μ g/ml) for 2–3 weeks. Individual clones were isolated, expanded and

analysed for expression of the b⁰+AT protein by western blot. One clone, number 5, was selected for further experiments based on the high level and stable expression of the protein. Similarly, an rBAT/b⁰+AT MDCK cell line was engineered starting with the selected b⁰+AT-expressing cell line. Briefly, b⁰+AT MDCK cells were transfected with pCDNA3Hygro-rBAT [rBAT cDNA cloned into the pCDNA3Hygro3.1 vector (Invitrogen) between the *Hind*III and *Xba*I sites] and individual colonies resistant to both geneticin (400 µg/ml) and hygromycin B (100 µg/ml) were isolated. Clones were screened by western blot analysis. In addition, a functional analysis was performed on preselected clones to identify clone 31B, which expresses a high L-cystine transport activity absent in parental MDCK cells.

Reconstitution

For liposome preparation, soybean phospholipids (Asolectin; Sigma) were partially purified as described (Kagawa and Racker, 1971) and crude pig brain lipids were extracted as described previously (Folch *et al.*, 1957). As detergent, cholic acid (Sigma) was used, recrystallized (Kagawa and Racker, 1971) and neutralized with NaOH to pH 7.4. Reconstitution of the transporter into liposomes was performed as described by Radian and Kanner (1985) with minor changes. Briefly, cells from a 15 cm diameter plate were washed twice in phosphate-buffered saline (PBS) and then harvested in 100 µl of PBS. A 25 µl aliquot of the cell suspension was mixed with liposomes and cholate up to a final volume of 200 µl. The mixture was passed through a Sephadex G-50 (Sigma) column pre-equilibrated with the reconstitution buffer (120 mM KPi, pH 7.4, 5 mM Tris-SO₄, 0.5 mM EDTA, 1 mM MgSO₄, 1% glycerol and, where indicated, with the desired concentration of amino acids). Liposomes were passed through a second spin column of identical composition but lacking the amino acid. Control liposomes (without entrapped amino acid) were also passed through a second spin column. The protein concentration in the cell suspension and in the liposomes was determined using a protein assay kit (Bio-Rad).

Uptake measurements

L-[³⁵S]cystine (Amersham) uptake measurements (20 µM) on whole cells were made as described (Font *et al.*, 2001). L-[³H]arginine (American Radiolabelled Chemicals) uptake measurements were performed following the same protocol, using 50 µM L-arginine (40 Ci/mmol, 0.5 µCi/well) and without L-glutamate in the uptake solution described (Font *et al.*, 2001). For L-[³H]arginine transport studies in the reconstituted system, 10 µl of liposomes was mixed with 180 µl of uptake solution [150 mM choline chloride, 10 mM Tris-HEPES, 1 mM MgCl₂, 1 mM CaCl₂, 0.5 µCi L-[³H]arginine, 2.8 µM valinomycin (Sigma) and L-arginine to the final concentration] equilibrated at 37°C, and incubated at 37°C for different periods. Reactions were terminated by the addition of 1 ml of ice-cold stop solution (150 mM choline chloride, 10 mM Tris-HEPES and 5 mM L-arginine) and filtration through membrane filters (Sartorius, 0.45 µm pore size). Filters were then washed three times in 2 ml of stop solution and dried, and the trapped radioactivity was counted. All experimental values were corrected by subtracting zero-time values obtained by adding the stop solution before the liposomes. Unless otherwise indicated only L-isomers of amino acids were used in this study.

Western blot analysis

Western blot analyses using anti-human b⁰+AT antibody (P6-870 Ab) and anti-human rBAT antibody (hrBAT-251 Ab) were carried out as described (Font *et al.*, 2001; Fernández *et al.*, 2002). For both total cell extracts and liposomes, 100 µg of protein in Laemmli sample buffer containing or not 100 mM dithiothreitol (DTT) was loaded in each lane for SDS-PAGE (10% polyacrylamide). As a control, brush border membrane preparations from human kidney were prepared by the Ca²⁺ precipitation method (Malathi *et al.*, 1979). For brush border membrane preparations, NEM at 5 mM was present in all buffers used (except the resuspension buffer) following Tate and colleagues (Wang and Tate, 1995), in order to prevent artifactual reduction/shuffling of disulfides. The reconstituted systems used in this study were prepared from cell extracts in the absence of NEM because it inactivates system b⁰+ in transfected HeLa cells (data not shown). Cell extracts, liposomes and membranes were stored at -80°C until used.

Supplementary data

Supplementary data are available at *The EMBO Journal* Online.

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References

- Abumrad, N., Harmon, C. and Ibrahimi, A. (1998) Membrane transport of long-chain fatty acids: evidence for a facilitated process. *J. Lipid Res.*, **39**, 2309–2318.
- Bassi, M.T., Gasol, E., Manzoni, M., Pineda, M., Riboni, M., Martín, R., Zorzano, A., Borsani, G. and Palacín, M. (2001) Identification and characterisation of human xCT that coexpresses, with 4F2hc heavy chain, the amino acid transport system x_c⁻. *Pflügers Arch.*, **442**, 286–296.
- Bertran, J., Werner, A., Moore, M.L., Stange, G., Markovich, D., Biber, J., Testar, X., Zorzano, A., Palacín, M. and Murer, H. (1992) Expression-cloning of a cDNA from rabbit kidney cortex that induces a single transport system for cystine, dibasic and neutral amino acids. *Proc. Natl Acad. Sci. USA*, **89**, 5601–5605.
- Bruce, L.J., Groves, J.D., Okubo, Y., Thilaganathan, B. and Tanner, M.J. (1994) Altered band 3 structure and function in glyophorin A- and B-deficient (MkMk) red blood cells. *Blood*, **84**, 916–922.
- Busch, A.E., Herzer, T., Waldegger, S., Schmidt, F., Palacín, M., Biber, J., Markovich, D., Murer, H. and Lang, F. (1994) Opposite directed currents induced by the transport of dibasic and neutral amino acids in *Xenopus* oocytes expressing the protein rBAT. *J. Biol. Chem.*, **269**, 25581–25586.
- Calonge, M.J. *et al.* (1994) Cystinuria caused by mutations in rBAT, a gene involved in the transport of cystine. *Nature Genet.*, **6**, 420–425.
- Chairoungdua, A. *et al.* (1999) Identification of an amino acid transporter associated with the cystinuria-related Type II membrane glycoprotein. *J. Biol. Chem.*, **274**, 28845–28848.
- Chillarón, J. *et al.* (1996) Obligatory amino acid exchange via systems b⁰+ like and y⁺L-like. A tertiary active transport mechanism for renal reabsorption of cystine and dibasic amino acids. *J. Biol. Chem.*, **271**, 17761–17770.
- Chillarón, J., Estévez, R., Samarzija, I., Waldegger, S., Testar, X., Lang, F., Zorzano, A., Busch, A. and Palacín, M. (1997) An intracellular trafficking defect in type I cystinuria rBAT mutants M467T and M467K. *J. Biol. Chem.*, **272**, 9543–9549.
- Chillarón, J., Roca, R., Valencia, A., Zorzano, A. and Palacín, M. (2001) Heteromeric amino acid transporters: biochemistry, genetics and physiology. *Am. J. Physiol. Renal. Physiol.*, **281**, F995–F1018.
- Coady, M.J., Jalal, F., Chen, X., Lemay, G., Berteloot, A. and Lapointe, J.Y. (1994) Electrogenic amino acid exchange via the rBAT transporter. *FEBS Lett.*, **356**, 174–178.
- Coady, M.J., Chen, X.Z. and Lapointe, J.Y. (1996) rBAT is an amino acid exchanger with variable stoichiometry. *J. Membr. Biol.*, **149**, 1–8.
- Devés, R. and Boyd, C.A. (2000) Surface antigen CD98(4F2): not a single membrane protein, but a family of proteins with multiple functions. *J. Membr. Biol.*, **173**, 165–177.
- Feliubadaló, L. *et al.* (International Cystinuria Consortium) (1999) Non-Type I cystinuria caused by mutations in *SLC7A9*, encoding a subunit (b⁰+AT) of rBAT. *Nature Genet.*, **23**, 52–57.
- Fernández, E., Carrascal, M., Rousaud, F., Abián, J., Zorzano, A., Palacín, M. and Chillarón, J. (2002) The rBAT-b⁰+AT heterodimer is the main apical reabsorption system for cystine in kidney. *Am. J. Physiol. Renal Physiol.*, **283**, F540–F548.
- Folch, J., Lees, M. and Sloane Stanley, G.H. (1957) A simple method for the isolation and purification of total lipids from animal tissues. *J. Biol. Chem.*, **226**, 497–509.
- Font, M. *et al.* (International Cystinuria Consortium) (2001) Functional analysis of mutations in *SLC7A9* and genotype/phenotype correlation in non-Type I cystinuria. *Hum. Mol. Genet.*, **10**, 305–316.

- Frimpter,G.W., Horwith,M., Furth,E., Fellows,R.E. and Thompson,D.D. (1962) Inulin and endogenous amino acid renal clearances in cystinuria: evidence for tubular secretion. *J. Clin. Invest.*, **41**, 281–288.
- Fukasawa,Y., Segawa,H., Kim,J.Y., Chairoungdua,A., Kim,D.K., Matsuuo,H., Cha,S.H., Endou,H. and Kanai,Y. (2000) Identification and characterization of a Na⁺-independent neutral amino acid transporter that associates with the 4F2 heavy chain and exhibits substrate selectivity for small neutral D- and L-amino acids. *J. Biol. Chem.*, **275**, 9690–9698.
- Geering,K. (2001) The functional role of beta subunits in oligomeric P-type ATPases. *J. Bioenerg. Biomembr.*, **33**, 425–438.
- Kagawa,Y. and Racker,E. (1971) Partial resolution of the enzymes catalyzing oxidative phosphorylation. *J. Biol. Chem.*, **246**, 5477–5487.
- Kanai,Y., Segawa,H., Miyamoto,K., Uchino,H., Takeda,E. and Endou,H. (1998) Expression cloning and characterization of a transporter for large neutral amino acids activated by the heavy chain of 4F2 antigen (CD98). *J. Biol. Chem.*, **273**, 23629–23632.
- Kanai,Y. *et al.* (2000) Transport properties of a system y⁺L neutral and basic amino acid transporter. Insights into the mechanisms of substrate recognition. *J. Biol. Chem.*, **275**, 20787–20793.
- Kirk,P., Wilson,M.C., Heddle,C., Brown,M.H., Barclay,A.N. and Halestrap,A.P. (2000) CD147 is tightly associated with lactate transporters MCT1 and MCT4 and facilitates their cell surface expression. *EMBO J.*, **19**, 3896–3904.
- Malathi,P., Preiser,H., Fairclough,P., Malett,P. and Crane,R.K. (1979) A rapid method for the isolation of kidney brush border membranes. *Biochim. Biophys. Acta*, **554**, 259–263.
- Mastroberardino,L., Spindler,B., Pfeiffer,R., Skelly,P.J., Loffing,J., Shoemaker,C.B. and Verrey,F. (1998) Amino-acid transport by heterodimers of 4F2hc/CD98 and members of a permease family. *Nature*, **395**, 288–291.
- Meier,C., Ristic,Z., Klausner,S. and Verrey,F. (2002) Activation of system L heterodimeric amino acid exchangers by intracellular substrates. *EMBO J.*, **21**, 580–589.
- Mizoguchi,K. *et al.* (2001) Human cystinuria-related transporter: Localization and functional characterization. *Kidney Int.*, **59**, 1821–1833.
- Mykkanen,J. *et al.* (2000) Functional analysis of novel mutations of y⁺LAT-1 amino acid transporter gene causing lysinuric protein intolerance (LPI). *Hum. Mol. Genet.*, **9**, 431–438.
- Nakauchi,J. *et al.* (2000) Cloning and characterization of a human brain Na⁺-independent transporter for small neutral amino acids that transport D-serine with high affinity. *Neurosci. Lett.*, **287**, 231–235.
- Palacín,M., Borsani,G. and Sebastio,G. (2001a) The molecular bases of cystinuria and lysinuric protein intolerance. *Curr. Opin. Genet. Dev.*, **11**, 328–335.
- Palacín,M., Goodyer,P., Nunes,V. and Gasparini,P. (2001b) Cystinuria. In Scriver,C.R., Beaudet,A.L., Sly,S.W. and Valle,D. (eds), *Metabolic and Molecular Bases of Inherited Diseases*. 8th edn. McGraw-Hill, New York, pp. 4909–4932.
- Pfeiffer,R., Rossier,G., Spindler,B., Meier,C., Kuhn,L. and Verrey,F. (1999a) Amino acid transport of y⁽⁺⁾L-type by heterodimers of 4F2hc/CD98 and members of the glycoprotein-associated amino acid transporter family. *EMBO J.*, **18**, 49–57.
- Pfeiffer,R., Loffing,J., Rossier,G., Bauch,C., Meier,C., Eggermann,T., Loffing-Cueni,D., Kuhn,L.C. and Verrey,F. (1999b) Luminal heterodimeric amino acid transporter defective in cystinuria. *Mol. Biol. Cell*, **10**, 4135–4147.
- Pineda,M., Fernández,E., Torrents,D., Estévez,R., Lopez,C., Camps,M., Lloberas,J., Zorzano,A. and Palacín,M. (1999) Identification of a membrane protein, LAT-2, that co-expresses with 4F2 heavy chain, an L-type amino acid transport activity with broad specificity for small and large zwitterionic amino acids. *J. Biol. Chem.*, **274**, 19738–19744.
- Pines,G., Danbolt,N.C., Bjoras,M., Zhang,Y., Bendahan,A., Eide,L., Koepsell,H., Storm-Mathisen,J., Seeberg,E. and Kanner,B.I. (1992) Cloning and expression of a rat brain L-glutamate transporter. *Nature*, **360**, 464–467.
- Radian,R. and Kanner,B.I. (1985) Reconstitution and purification of the sodium- and chloride-coupled γ -aminobutyric acid transporter from rat brain. *J. Biol. Chem.*, **260**, 11859–11865.
- Rossier,G., Meier,C., Bauch,C., Summa,V., Sordat,B., Verrey,F. and Kuhn,L.C. (1999) LAT2, a new basolateral 4F2hc/CD98-associated amino acid transporter of kidney and intestine. *J. Biol. Chem.*, **274**, 34948–34954.
- Sato,H., Tamba,M., Ishii,T. and Bannai,S. (1999) Cloning and expression of a plasma membrane cystine/glutamate exchange transporter composed of two distinct proteins. *J. Biol. Chem.*, **274**, 11455–11448.
- Sato,H., Tamba,M., Kuriyama-Matsumura,K., Okuno,S. and Bannai,S. (2000) Molecular cloning and expression of human xCT, the light chain of amino acid transport system x_c⁻. *Antioxid. Redox Signal.*, **2**, 665–671.
- Segawa,H., Fukasawa,Y., Miyamoto,K., Takeda,E., Endou,H. and Kanai,Y. (1999) Identification and functional characterization of a Na⁺-independent neutral amino acid transporter with broad substrate selectivity. *J. Biol. Chem.*, **274**, 19745–19751.
- Silbernagl,S. (1988) The renal handling of amino acids and oligopeptides. *Physiol. Rev.*, **68**, 911–1007.
- Torras-Llort,M., Torrents,D., Soriano-García,J.F., Gelpi,J.L., Estévez,R., Ferrer,R., Palacín,M. and Moreto,M. (2001) Sequential amino acid exchange across b^{0,+}-like system in chicken brush border jejenum. *J. Membr. Biol.*, **180**, 213–220.
- Torrents,D., Estévez,R., Pineda,M., Fernández,E., Lloberas,J., Shi,Y.B., Zorzano,A. and Palacín,M. (1998) Identification and characterization of a membrane protein (y⁺L amino acid transporter-1) that associates with 4F2hc to encode the amino acid transport activity y⁺L. A candidate gene for lysinuric protein intolerance. *J. Biol. Chem.*, **273**, 32437–32445.
- Verrey,F., Jack,D.L., Paulsen,I.T., Saier,M.H., Jr and Pfeiffer,R. (1999) New glycoprotein-associated amino acid transporters. *J. Membr. Biol.*, **172**, 181–192.
- Verrey,F., Meier,C., Rossier,G. and Kuhn,L.C. (2000) Glycoprotein-associated amino acid exchangers: broadening the range of transport specificity. *Pflügers Arch.*, **440**, 503–512.
- Wagner,C.A., Lang,F. and Bröer,S. (2001) Function and structure of heterodimeric amino acid transporters. *Am. J. Physiol. Cell Physiol.*, **281**, C1077–C1093.
- Wang,Y. and Tate,S.S. (1995) Oligomeric structure of a renal cystine transporter: implications in cystinuria. *FEBS Lett.*, **368**, 389–392.
- Wilson,M.C., Meredith,D. and Halestrap,A.P. (2002) Fluorescence resonance energy transfer studies on the interaction between the lactate transporter MCT1 and CD147 provide information on the topology and stoichiometry of the complex *in situ*. *J. Biol. Chem.*, **277**, 3666–3672.

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