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The Escherichia coli multidrug transporter MdfA catalyzes both electrogenic and electroneutral transport reactions

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The resistance of cells to many drugs simultaneously (multidrug resistance) often involves the expression of membrane transporters (Mdrs); each recognizes and expels a broad spectrum of chemically unrelated drugs from the cell. The Escherichia coli Mdr transporter MdfA is able to transport differentially charged substrates in exchange for protons. This includes neutral compounds, namely chloramphenicol and thiophenicol, and lipophilic cations such as tetraphenylphosphonium and ethidium. Here we show that the chloramphenicol and thiophenicol transport reactions are electrogenic, whereas the transport of several monovalent cationic substrates is electroneutral. Therefore, unlike with positively charged substrates, the transmembrane electrical potential (negative inside) constitutes a major part of the driving force for the transport of electroneutral substrates by MdfA. These results demonstrate an unprecedented ability of a single secondary transporter to catalyze discrete transport reactions that differ in their electrogenicity and are governed by different components of the proton motive force.

Multidrug resistance (Mdr) transporters that expel drugs from the cytoplasm or cytoplasmic membrane to the external medium constitute one of the major causes of Mdr. Mdr transporters are widely distributed in bacteria, including pathogenic strains (1–4). The prokaryotic Mdr transporters are able to extrude a variety of dissimilar lipophilic compounds, many of which are positively charged under physiological conditions. However, there are some bacterial Mdr proteins that also interact with neutral and zwitterionic drugs, and some transporters export lipophilic anionic drugs (5–8).

We investigated the Mdr phenomenon by using Escherichia coli Mdr transporter MdfA (9) as a model for secondary Mdr transporters. MdfA is a 410-aa-long membrane protein of which close homologues have been identified thus far only in the pathogenic bacteria Salmonella enterica serovar Typhimurium (90% identity) (10) and Yersinia pestis (73% identity) (11). Cells expressing MdfA from a multicopy plasmid exhibit Mdr resulting from active drug extrusion driven by the proton electrochemical gradient. Previous studies have proposed that MdfA is a drug/proton antiporter (12). As predicted from the hydropathy plot of the protein and confirmed by gene-fusion studies, the putative 12 transmembrane regions of MdfA have only one charged amino acid residue, glutamate at position 26, which is embedded in the membrane, in the middle of putative transmembrane segment 1 (13). As demonstrated by mutational analysis, the amino acid residue at this position is important for substrate recognition and binding by MdfA but not for proton translocation (ref. 14 and J.A., O.L., and E.B., unpublished data). As mentioned, MdfA recognizes compounds that are not only structurally unrelated but also dissimilar in charge, and recent work has demonstrated simultaneous binding of such substrates to the transporter (15). This property poses an interesting question regarding the electrogenicity of the substrate/proton exchange with positively charged compounds versus electroneutral ones.

In E. coli, secondary transport is driven mostly by energetically downhill influx of protons. Depending on the proton/substrate stoichiometry of the reaction and the charge on the substrate, exchange of protons with a substrate by antiporters may result in a net movement of electric charges (electrogenic transport) or no net movement of charges (electroneutral transport). Electrogenic transport will be driven by both components (Δψ and ΔpH) of the proton electrochemical gradient (ΔμH+), but when transport is electroneutral, ΔpH should be the primary driving force. Importantly, whereas ΔμH+ remains relatively constant at external pH values ranging from 5.0 to 8.0, the electrical (Δψ) and chemical (ΔpH) components change. Because internal pH is kept constant at 7.6 as external pH is raised, ΔpH (inside alkaline) drops and even becomes inverted (inside acid) above pH 7.6 (16), whereas Δψ (inside negative) increases. Accordingly, ΔpH-driven electroneutral transport processes should be inhibited at alkaline external pH values, whereas electronegenic transport processes should be less affected. Of the secondary transport proteins described to date, they catalyze either electronegenic or electroneutral translocation reactions. In this regard, Mdr transporters represent a unique subset of secondary transporters. Because of their ability to transport differentially charged substrates, they are theoretically capable of catalyzing transport reactions that are electrically distinct. Most probably, a transport process that involves ΔμH+–dependent active extrusion of neutral drugs is electrogenic, but the degree to which Δψ or ΔpH (inside negative and alkaline) contributes to the uptake or extrusion of a given solute may vary (17).

Here we have examined the intriguing possibility that different MdfA-mediated transport reactions exist, each accommodating a distinct group of substrates. The results demonstrate that the MdfA-catalyzed transport of neutral substrates is indeed electrogenic, whereas transport of cationic substrates is electroneutral. Thus, in addition to their promiscuity regarding substrate recognition, Mdr transporters are also versatile regarding the component of ΔμH+ used for transport.

Materials and Methods

Materials. E. coli lipids were from Avanti. [3H]Tetraphenylphosphonium (TPP+) [32 Ci/mmol (1 Ci = 37 GBq)] was purchased from Amersham Pharmacia, and [3H]chloramphenicol (20 Ci/mmol) was purchased from NEN. Nigericin, valinomycin, carbonyl cyanide m-chlorophenyl hydradzone (CCCP), ATP, ethidium bromide (EtBr), chloramphenicol, kanamycin, Hoechst 33342, DNase, phosphatidyl choline, and ampicillin all were purchased from Sigma. Benzoalkonium chloride was obtained from Calbiochem, and thiophenicol was obtained from

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Abbreviations: Mdr, multidrug resistance; CCCP, carbonyl cyanide m-chlorophenyl hydradzone; EtBr, ethidium bromide; TPP+, tetraphenylphosphonium.

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ICN. Oxonol V and 9-amino-6-chloro-2-methoxyacridine were purchased from Molecular Probes. N-dodecyl-maltoside was purchased from Anacrace (Maumee, OH). His Bind resin was obtained from Novagen and used for purification of MdfA-6His. Cellulose acetate and nitrocellulose filters (0.2 μm) were from Schleicher & Schüll. Polycarbonate filters (0.4 μm) were from Osmonics (Minnetonka, MN). All other materials were reagent-grade and obtained from commercial sources.

**Drug-Resistance Assays.** E. coli UTmdfA::kan (14) was transformed with the indicated plasmid (pT7-5 or pT7-5 mdfA-6His), and after 1 h of recovery at 37°C the transformants were diluted as indicated and plated over LB agar plates. The plates were supplemented with 75 mM bis-Tris-propane at the desired pH and ampicillin (100 μg/ml) kanamycin (30 μg/ml), and the indicated concentrations of the test compounds (chloramphenicol, thiamphenicol, benzalkonium, or EtBr). The ability of the transformants to form single colonies was recorded after 36 h at 37°C.

**Transport Assays in Whole Cells.** EtBr-efflux assays (6) and chloramphenicol or TPP⁺ uptake assays (15) were conducted as described with the following modifications: For EtBr efflux, E. coli UTL2mfdA::kan cells harboring pT7-5 or pT7-5 mdfA-6His (15) were grown to an OD₆₀₀ of 0.8 units in LB broth and kept on ice. Two-milliliter aliquots (OD₆₀₀ of 0.3 units) were pelleted and resuspended in 2 ml of 50 mM potassium phosphate at the desired pH and supplemented with 100 μM CCCP and 5 μM EtBr. Cells were incubated for 5° at 35°C, pelleted, and resuspended in the same buffer but without CCCP. Transport was initiated by the addition of 0.2% glucose. For chloramphenicol uptake, E. coli UTLmdfA::kan cells harboring pT7-5 or pT7-5 mdfA-6His were grown to an OD₆₀₀ of 0.5 units in LB broth, washed once in 50 mM potassium phosphate at the desired pH, resuspended in the same buffer, at an OD₂₈₀ of 30 units, and aliquoted (50 μl). After 15 min recovery at 35°C in the presence of 0.2% glucose, transport was initiated by the addition of an equal volume (50 μl) of the same buffer containing [³²H]-chloramphenicol. [³²H]TPP⁺ uptake was conducted essentially the same as with chloramphenicol. UTL2mfdA::kan cells were resuspended to an OD₂₈₀ of 13.5 units. In experiments with the ionophores valinomycin (5 μM) or nigericin (2 μM), the outer-membrane permeability mutant was made further permeable to the ionophores by a 15-min incubation at 35°C followed by centrifugation (100,000 × g) and resuspended in the same buffer at a protein concentration of 8 mg/ml. Proteoliposomes were aliquoted, frozen in liquid nitrogen, and stored at −80°C for up to 6 months.

**Preparation of inverted Membrane Vesicles.** E. coli UTL2mfdA::kan cells overexpressing MdfA were grown and treated as described (15). Inverted membrane vesicles were prepared from these cells as described (18). Cells were washed once in 50 mM potassium phosphate, pH 7.5/5 mM MgSO₄. Cells (0.2 g/ml) in 50 mM potassium phosphate, pH 7.5/5 mM MgSO₄/30 μg/ml DNase/0.5 mM Pefablock/10 mM 2-mercaptoethanol were disrupted by French Press at 8,000 psi (1 psi = 6.89 kPa) (three passages). Vesicles in 50 mM potassium phosphate, pH 7.5/10% glycerol were frozen in liquid nitrogen and kept at −80°C.

**Hoechst 33342 Transport in Inverted Membrane Vesicles.** Transport of Hoechst 33342 was conducted as described (19) with the following modifications: 4 μl of vesicles (30 mg of protein per ml) were diluted into 2 ml of 50 mM Hepes, pH 7.0/50 mM potassium gluconate/10 mM magnesium sulfate/0.2 μM Hoechst 33342. Transport was initiated by the addition of 0.2 mM ATP. Nigericin and valinomycin (2 μM each) were added as indicated.

**Membrane Potential Measurements in Inverted Membrane Vesicles.** Forty microliters of vesicles (30 mg of protein per ml) were diluted into 2 ml of 50 mM Hepes, pH 7.0/50 mM potassium gluconate/10 mM magnesium sulfate/0.5 μM oxonol V. A transmembrane potential was generated by the addition of 0.4 mM ATP, CCCP (10 μM) was used to completely abolish the membrane potential. The fluorescence was measured in a Perkin–Elmer SL50 fluorimeter with excitation and emission wavelengths of 599 and 634 nm, respectively, and a slit width of 10 nm.

**Purification and Reconstitution of MdfA.** MdfA was purified to near homogeneity as described (15). For reconstitution, a combination of methods was used (20–22). Seventy-five milligrams of 3:1 E. coli lipids (acetone/ether washed) and egg-yolk phosphatidylcholine were dissolved in 2 ml of 20 mM sodium phosphate, pH 7.0/100 mM sodium gluconate/2 mM magnesium sulfate/0.3% glycerol/0.03% N-dodecyl-maltoside. Dithiothreitol (1 mM) was added to the argon-flushed lipids, which were then subjected to two rounds of freezing (in liquid nitrogen) and thawing (undisturbed at room temperature). Next, preformed liposomes were prepared by extrusion (10 times) through a double layer of 0.4-μm polycarbonate filters, and purified MdfA (0.5 ml of 2 mg/ml protein) was added in 20 mM Tris-HCl, pH 7.9/0.5 M NaCl/250 mM imidazole/0.1% N-dodecyl-maltoside/1% glycerol/0.2 mg/ml lipids. As a control, 0.5 ml of the same buffer but without protein was added to identically treated lipids. The mixture (2.5 ml) was agitated gently at 4°C for 30 min and then transferred (at room temperature) into vigorously stirred 210 ml of argon-flushed 20 mM sodium phosphate, pH 7.0/100 mM Na-glucanote/2 mM magnesium sulfate. After 20 min of stirring at room temperature, proteoliposomes were collected by 1 h of centrifugation at 185,000 × g, and resuspended in the same buffer at a protein concentration of 0.8 mg/ml. Proteoliposomes were aliquoted, frozen in liquid nitrogen, and stored at −80°C for up to 6 months.

**Transport Assays in Proteoliposomes.** For all experiments, proteoliposomes were thawed undisturbed at room temperature and diluted at least 10-fold with the desired loading buffer. After two cycles of freezing and thawing, the proteoliposomes were extruded (7–10 times until clear) through a double layer of 0.4-μm polycarbonate filters. After centrifugation (7 min at 280,000 × g), the proteoliposomes were resuspended in the same buffer at a protein concentration of 8 mg/ml (for TPP⁺ uptake) or 2 mg/ml (for chloramphenicol uptake). For evaluation of diffusion potential formation, proteoliposomes or liposomes were routinely assayed by fluorescence spectrometry (Perkin–Elmer SL50 fluorimeter) by using oxonol V (0.5 μM; excitation, 599 nm; emission, 634 nm; slit width, 10 nm) for monitoring Δψ and 9-amino-6-chloro-2-methoxyacridine (1 μM; excitation, 409 nm; emission, 474 nm; slit width, 3 nm) for ΔpH. For generation of ΔpH (inside acidic), proteoliposomes were loaded with 150 mM NH₄Cl/50 mM potassium gluconate/5 mM Tris-HCl, pH 6.5 [and diluted 100-fold into the same buffer (control, no energy gradient)], or 150 mM choline chloride/50 mM potassium gluconate/5 mM Tris-HCl, pH 7.5. For generation of Δψ (inside positive), proteoliposomes were loaded with 120 mM sodium gluconate/2 mM magnesium sulfate/5 mM sodium phosphate, pH 7.0 [and diluted 100-fold into the same buffer (control, no energy gradient)], or 120 mM potassium gluconate/2 mM MgSO₄/5 mM potassium phosphate, pH 7.0/5 μM valinomycin. For the generation of ΔμH⁺ (inside positive and acidic), proteoliposomes were loaded with 120 mM sodium gluconate/2 mM magnesium sulfate/150 mM NH₄Cl/5 mM Tris-HCl, pH 6.5 [and diluted 100-fold into the same buffer (control, no energy gradient)], or 120 mM potassium gluconate/2 mM MgSO₄/150 mM choline chloride/5 mM Tris-HCl, pH 7.5/5 μM valinomycin.
Transport was conducted in 50-μl (chloramphenicol) samples and terminated by the addition of 1.5 mL of ice-cold LiCl (0.1 M) and filtration through 0.2-μm filters (nitrocellulose for chloramphenicol or cellulose acetate for TPP⁺) followed by an additional 1.5-mL wash. Experiments were repeated three times, each in triplicate.

Results

The MdfA-Mediated Resistance Toward Cationic and Neutral Drugs Is Affected Differently by Changes in the External pH. Initial evidence suggesting that the transport reactions of differentially charged substrates may differ originated from the results of drug-resistance assays on solid medium. Drug resistance to positively charged and uncharged compounds was tested under various pH conditions and over a broad range of drug concentrations. The results of a representative experiment are shown in Fig. 1. In the absence of drugs, cells harboring either plasmid, pT7-5 or pT7-5 (mdfA), grew equally well in the pH range of 6.5–8.0. However, when the positively charged substrates EtBr or benzalkonium were added to the medium, only cells expressing MdfA were able to grow, but the growth was attenuated at basic pH. In contrast, the MdfA-mediated resistance to the neutral drugs chloramphenicol and thiamphenicol was maintained also at pH 8.0. Because no growth of control cells (cells that do not express MdfA) was observed at the drug concentrations tested (Fig. 1A), the effect of pH on their intrinsic drug resistance was tested at extremely low drug concentrations (2–4% of the drug concentration used in Fig. 1A). As shown (Fig. 1B), Mdr of control cells was also affected by external pH changes. However, unlike MdfA-expressing cells, control cells are generally less resistant at basic pH toward both cationic and neutral drugs.

The Transport of Positively Charged Substrates Is Inhibited at a Basic pH, Whereas the Transport of Chloramphenicol Remains Unaffected. The dissimilar pH-dependent resistance phenotypes described above might reflect a pH-dependent transport phenomenon. The ability of control and MdfA-expressing E. coli cells to extrude drugs against a concentration gradient was tested at a pH range of 6.5–8.5. Transport of each of the positively charged substrates EtBr and TPP⁺ by MdfA-expressing cells was inhibited gradually by increasing the external pH, with a complete arrest of transport at pH 8.5 (Fig. 2A and B Left). In contrast, transport of the uncharged chloramphenicol was affected to a much lesser degree by the external pH, and substantial transport of chloramphenicol was still apparent at pH 8.5 (Fig. 2C Left). Importantly, the uptake of all the test drugs by control cells (that do not express MdfA) is only slightly affected by identical pH changes (Fig. 2B Right and C Right). Therefore, it seems likely that the pH dependence of MdfA-mediated resistance to cationic drugs (Fig. 1) is due to the inhibitory effect of a basic pH on their transport.

The Dissipation of ΔΨ or ΔpH Has Distinct Effects on the Transport of Differentially Charged Substrates. By gradually raising the extracellular pH, the ΔpH (inside alkaline) across the cytoplasmic...
The membrane potential is dissipated only by the transport of electroneutral drugs. Thus far, the results have suggested that the MdfA-mediated transport of cationic substrates differs from that of uncharged substrates with regard to the driving force. To explore the electrogenicity of these presumably distinct transport reactions, we measured the effect of the transport of various drugs on a preestablished Δψ. Inverted membrane vesicles were energized by the addition of ATP, and the formation of a Δψ (inside positive) was followed by fluorescence spectroscopy using the Δψ-sensitive dye oxonol V. The Δψ generated by vesicles prepared from MdfA-expressing cells or from control cells was indistinguishable (Fig. 4). The Δψ attained a steady-state level, various substrates were added in concentrations that had no effect on the oxonol V fluorescence or on the Δψ generated by control vesicles (Fig. 4A, data not shown). The addition of benzalkonium to MdfA-containing vesicles did not dissipate the Δψ but rather caused a small but reproducible increase in its magnitude (Fig. 4D). A similar effect on Δψ was observed by the addition of nigericin (Fig. 4E). In both cases, dissipation of the Δψ resulted in an increase of Δψ. However, in contrast to the effects of benzalkonium or nigericin, adding chloramphenicol or thiamicphenicol to MdfA-containing vesicles resulted in a marked dissipation of Δψ (Fig. 4B and C). Taken together, these results suggest that the transport of cationic substrates is electroneutral, whereas that of neutral substrates is electrogenic.

Reconstituted MdfA Catalyzes both Electrogenic and Electroneutral Transport. To determine whether the MdfA protein suffices to catalyze both electrogenic and electroneutral transport, we purified and reconstituted the transporter in proteoliposomes. The reconstituted MdfA is functional, as demonstrated by transport assays with chloramphenicol and TPP⁺ (Fig. 5). Transport assays were driven by artificially imposed diffusion potentials (see Materials and Methods). The established potentials were evaluated routinely by using oxonol V and 9-amino-6-chloro-2-methoxyacridine fluorescence for Δψ and ΔpH, respectively. The uptake of chloramphenicol into proteoliposomes was driven not only by an artificially imposed Δψ but also by an imposed ΔpH (inside positive), suggesting that, by definition,
transport is electrogenic. As expected, ΔpH alone (inside acid) also drove chloramphenicol uptake by MdfA-containing liposomes (Fig. 5A). With the positively charged substrate TPP⁺, the ΔµH⁺-driven transport was similar to that observed when only a ΔpH (inside acid) was imposed. However, generating only a Δψ (inside positive) failed to drive TPP⁺ transport into the proteoliposomes (Fig. 5B). These results indicate that unlike with the neutral substrate chloramphenicol, the Δψ is not sufficient for the active transport of TPP⁺, whereas ΔpH suffices for the uptake. Therefore, MdfA catalyzes both the electrogenic and electroneutral transport reactions without the need for accessory components.

Discussion

Most secondary transporters are specific regarding their substrate recognition properties and transport driving forces. In contrast, Mdr transporters are remarkably promiscuous in various basic aspects of active transport. Unlike the former transporters that specialize in the transport of a well-defined set of molecules, Mdr transporters are able to extrude a large number of chemically dissimilar compounds. In addition to being dissimilar in their chemical structure, the substrates of many Mdr transporters also differ in their charge. Moreover, it is demonstrated here with MdfA, the remarkable versatility of these transporters is exemplified further by their ability to catalyze dissimilar transport reactions.

Theoretically, in extruding a positively charged compound across the cell membrane, the transporter must move the charges against the transmembrane electrical potential, Δψ (inside negative), unlike with uncharged compounds. Accordingly, the transport of uncharged compounds seems more energetically favorable than the transport of charged substrates. Nevertheless, lipophilic cations clearly make up the majority among the substrates of MdfA. From a bioenergetic standpoint, a transporter that has to deal with differentially charged substrates has at least three theoretical modes of action: (i) the transporter may use different drug/proton stoichiometries with neutral versus positively charged, thus balancing the charge difference between the substrates with the appropriate number of counter ions; (ii) the transporter may cotransport an additional charged molecule with some of its substrates to reach the same net charge movement in each of the transport reactions; and (iii) the transporter may use an equal drug/proton stoichiometry with all the substrates, and because the charge of the substrates differs, this must result in different electrogenicities of the different transport reactions. Here we investigated this issue by using MdfA as a model Mdr transporter. The results clearly demonstrate the existence of two types of transport reactions that differ in their electrogenicity. With the monovalent cationic substrates, because their transport is electroneutral, we suggest a proton/substrate stoichiometry of 1 for these substrates. Also for the electroneutral substrates, because their transport is electrogenic, we suggest a proton/substrate stoichiometry of 1, although higher values cannot be ruled out at this stage.

Initial indications of dissimilar transport reactions could already be deduced from experiments examining the in vivo...
function of MdfA as an Mdr protein. By increasing the external pH, the resistance of cells expressing MdfA to cationic drugs was hampered, yet resistance to uncharged drugs remained unaffected (Fig. 1). The differential effect of pH was shown to be specific for MdfA-expressing cells. Direct evidence for a strong correlation between the export of positively charged drugs and the ΔpH (inside alkaline) came from transport assays with whole cells. In contrast, the transport of an uncharged drug proved to depend less on the ΔpH (Fig. 2), probably because of the MdfA ability to use also Δψ as a driving force for transport of uncharged drugs. Similar conclusions were derived from transport experiments of differentially charged substrates in the presence of valinomycin or nigericin (Fig. 3), which strongly indicated that the transport of positively charged substrates depends solely on a ΔpH and completely independent of the Δψ. In contrast, the transport of uncharged substrates depends on both components of the ΔμH+. The experiments described in Fig. 3C clearly demonstrate the versatile ability of MdfA in using either component of ΔμH+ for transport of an uncharged substrate. At pH 7.0, Δψ constitutes ≈70% of the proton motive force, whereas ΔpH constitutes ≈30% (23). Accordingly, the addition of valinomycin abolishes a major part of the MdfA-mediated chloramphenicol transport, whereas addition of nigericin exerts a minor effect (Fig. 3C Left). At pH 6.5, the contribution of Δψ and ΔpH to the overall driving force is approximately equal (23), as also reflected by the inhibitory effect of the ionophores on chloramphenicol transport (Fig. 3C Right). Transport experiments with reconstituted MdfA (Fig. 5) suggested that its ability to perform dissimilar transport reactions is independent of accessory components and is an inherent property of the transporter. This phenomenon was established further in inverted membrane vesicles and preformed Δψ (inside positive). The addition of electroneutral drugs consumed the Δψ, whereas the transport of positively charged drugs led to an increase in the magnitude of the Δψ (Fig. 4). Taking into account that ΔμH+ is a relatively constant value, the consumption of ΔpH through the electroneutral exchange of a positively charged drug and a proton will bring about a concomitant increase in Δψ. A similar effect on Δψ was exerted by nigericin, which operates as an electroneutral H+/K+ exchanger, hence converting ΔpH to Δψ (Fig. 4).

Because for secondary transport systems described to date transport has been characterized as being either electronegative or electroneutral, MdfA provides an example of a secondary transporter able to catalyze both types of transport. These results do not only offer an explanation for a basic phenomenon, how Mdr transporters deal with substrates differing in their charge, but also raise clinically oriented considerations in the treatment of multidrug-resistant bacterial strains; e.g., the use of certain positively charged drugs may be advantageous in a basic environment.

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