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## Mechanistic diversity in ATP-binding cassette (ABC) transporters

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### Abstract

ABC transporters catalyze transport reactions ranging from high-affinity uptake of micronutrients into bacteria to the extrusion of cytotoxic compounds from mammalian cells. Crystal structures of ABC domains and full transporters have provided a framework for formulating reaction mechanisms of ATP-driven substrate transport, but recent studies suggest a remarkable mechanistic diversity within the protein family. This review evaluates the differing mechanistic proposals and outlines future directions to explore ABC transporter-catalyzed reactions.

### Introduction

ABC transporters are a family of membrane proteins that mediate diverse, ATP-driven transport processes. They contain a pair of conserved, cytoplasmic domains termed ATP-binding cassettes (ABC) or nucleotide-binding domains (NBD). NBDs hydrolyze ATP and drive conformational changes in the attached transmembrane domains (TMDs), allowing substrates to cross the lipid bilayer of the membrane either into the cytoplasm (import) or out (export). The domain arrangement and conserved sequence motifs are summarized in **box 1**, and representative structures of the different sub-families of ABC transporters are shown in Fig. 1.

ABC transporters are found in all domains of life. In bacteria, ABC importers are involved in the uptake of nutrients and micro-nutrients through medium- and high-affinity pathways<sup>1,2</sup>. These reactions are highly specific and require a periplasmic (or lipid-anchored external) substrate binding protein<sup>3,4</sup>. Bacterial ABC exporters have diverse roles, including the extrusion of (often lipid-linked) building blocks required for cell wall assembly<sup>5-7</sup>. Other ABC exporters are poly-specific and contribute to drug resistance by exporting certain toxic substances<sup>8,9</sup>.

There are 48 distinct human ABC transporters that belong to various sub-families<sup>10</sup>. Most are thought to be exporters, and some display a remarkable poly-specificity for substrates. Some human ABC transporters have functions other than substrate translocation, e.g. in regulating potassium channels (SUR1 and SUR2)<sup>11,12</sup> or acting as an ATP-gated chloride channel (CFTR)<sup>13</sup>. Many human ABC transporters are bio-medically and clinically relevant: For example, multidrug transporters such as ABCB1, ABCG2, or ABCC1 protect various organs from toxic insult, but also contribute to drug resistance of cancer cells<sup>14-17</sup>; In liver, a series of ABC transporters (ABCB11, ABCB4, ABCG5/G8) catalyzes the generation of bile, with dysfunction of either transporter leading to liver disease<sup>18-22</sup>; The transporter of antigenic peptides (TAP, a heterodimer of ABCB2 and ABCB3) has an important role in the adaptive immune response<sup>23,24</sup>. Archaea, yeasts, plants, and human parasites also have ABC transporters that catalyze vital functions (reviewed in<sup>25-27</sup>).

In addition to the fundamental question of how they translocate large substrates across lipid bilayers, a detailed structural and mechanistic understanding of ABC transporters may contribute to drug discovery, for example in the case of multidrug or liver ABC transporters<sup>28-31</sup>.

It has been established that the closed NBD dimer conformation is essential for ATPase activity of full ABC transporters, because physical NBD separation results in the transporter inactivation<sup>32-34</sup>. Given the tight interaction between the NBDs and the coupling helices, it is generally accepted that the conserved power stroke of ABC transporters is the pushing together and pulling apart of the

cytoplasmic ends of the TMDs during the ATP hydrolysis cycle (reviewed in <sup>35-38</sup>). However, recent structural and mechanistic studies challenged the notion that a single, conserved mechanism can explain the reactions even within individual sub-families of ABC transporters. This review evaluates transport mechanisms derived from structures of full transporters as well as from functional, spectroscopic, and biophysical studies.

### **Alternating access in ABC importers**

**Type 1 ABC importers.** Type I ABC importers facilitate medium-affinity uptake of diverse nutrients including ions, sugars, amino acids, short peptides, and oligosaccharides, into bacteria. The *E. coli* maltose transporter has been extensively studied (REFS) and can serve as a mechanistic model of a generic mechanism of type 1 ABC importers (Fig. 2a). The transport cycle can be thought to start in an inward-facing conformation, with substrate-loaded binding protein approaching the external side of the transporter. There is a possibility of futile ATPase activity at this state, resulting in a "basal ATPase rate". However, this activity is slow in type 1 ABC importers and often additionally inhibited. Docking of the substrate binding protein stimulates the basal ATPase rate of the maltose<sup>39</sup> and alginate transporters<sup>40</sup>, but not of the molybdate/tungstate transporters<sup>32,41</sup>. The key conformational transition during the transport cycle (state 1 to 2) is the closing of the NBDs, which pushes the coupling helices towards each other and converts the TMDs to an outward-facing conformation. This generates a tunnel from the attached binding protein to a low-affinity substrate-binding pocket in the translocation pathway, halfway across the membrane. In the maltose transporter, two effects were found to ensure the release of substrate from the binding protein: First, the binding pocket is slightly distorted. Second, periplasmic TMD loops (in particular a "scoop loop") cause a partial steric clash with bound substrate<sup>42</sup>. Although the affinity of maltose to the temporary binding site between the TMDs is only moderate (Kd in the millimolar range), the substrate populates this pocket because the local concentration in the tunnel is at least 2 orders of magnitude above the dissociation constant. In the subsequent, irreversible step (state 2 to 3) ATP is hydrolyzed and inorganic phosphate released. This causes the NBD dimer to open, pulling the coupling helices outwards and triggering the conversion to an inward-facing conformation. The substrate can now dissociate into the cytoplasm. Finally, the exchange of ADP for ATP resets the system (state 3 to 1).

There are mechanistic variations of the theme in certain type 1 ABC importers. Because of their role in medium-affinity pathways, efficient coupling of ATP hydrolysis to transport is important, and the observed basal ATPase activities are generally low. While some transporters such as OpuA have strict coupling of 2 ATP per transport cycle<sup>43</sup>, additional regulatory strategies have been observed for other transporters in the form of trans-inhibition<sup>32,33,44</sup> or the binding of regulatory proteins<sup>45</sup>.

**Type 2 ABC importers.** These are generally part of high-affinity uptake pathways for metal chelates including heme and other iron-containing complexes, and cobalamin (reviewed in <sup>46,47</sup>). These substrates are not only larger and more hydrophobic than those of type I ABC importers, but available at low concentrations only. The mechanism proposed for the *E. coli* vitamin B12 transporter BtuCD-F can serve as a model for other type 2 ABC importers. Unlike observed in the maltose transporter, there is no measurable affinity (and thus no substrate binding pocket) for B12 in the TMDs of BtuCD-F. The obtained crystal structures revealed three gate regions in the transmembrane BtuC subunits, two on the cytoplasmic side (cyto-gates 1 and 2) and one on the periplasmic side (peri-gate)<sup>48,49</sup>. The key difference to type 1 ABC importers is that in BtuCD-F, ATP binding and hydrolysis does not generate an inward-facing conformation by itself. Rather, the two cytoplasmic gates separate the cytoplasm from the central translocation cavity both in the ATP-bound and nucleotide-free states, as long as transport substrate is absent. Figure 3b shows a schematic of the transport mechanism deduced from structural, biochemical, and spectroscopic data<sup>48-52</sup>. The cycle can be thought to start in

an ATP-bound state (state 1) with a closed NBD dimer conformation, and coupling helices pushed together, which causes cyto-gate 2 to close. Because the peri-gate is open, state 1 is an outward-facing conformation. Futile ATPase activity (arrows to the left) has been observed in all type 2 ABC importers studied to date, although the measured rates are very low and the loss of cytoplasmic ATP apparently manageable for the cell. State 2 is reached upon docking of the substrate binding protein and the release of substrate into a hydrophobic ("teflon") cavity with no measurable affinity. As in type 1 importers, the high-affinity pocket in the substrate binding protein is distorted upon docking, and loops of the TMD cause a steric clash, "scooping" the substrate from its pocket and enclosing it in the translocation pathway between the TMDs. In BtuCD, trapping of substrate in this cavity could only be demonstrated in liposomes. In detergent, the substrate escapes from the cavity, probably due to increased protein dynamics or competition for binding with detergent molecules. The following, irreversible step (state 2 to 3) is the hydrolysis of ATP and the release of phosphate, which requires (at least partial) opening of the NBD dimer, pulling the coupling helices apart and opening cyto-gate 2. Due to the size of the trapped transport substrate, cyto-gate 1 is now unable to close, which probably causes a strained conformation and pressure from the sides onto the substrate. This resembles a peristaltic force and may contribute to substrate dissociation into the cytoplasm. Upon substrate release, BtuCD relaxes into an asymmetric conformation (state 4) that does not feature a central cavity anymore, because both cyto-gate 1 and the peri-gate are now closed<sup>51</sup>. This state is very stable *in vitro* and probably prevents potential ion leakage during the subsequent re-setting of the system. The asymmetric conformation of state 4 appears unique to type 2 importers because the structural elements involved are not present in type 1 importers. Notably, the mechanistic proposal does not assign any role to the BtuCD conformation first observed, an apo-state with outward-facing TMDs<sup>53</sup>. Although such a conformation may transiently occur in the cell (before ADP is replaced by ATP) and contribute to futile ATP hydrolysis, it is not essential to formulate a productive transport cycle. Another important point is that nucleotide-bound BtuCD with a closed NBD dimer could only be stably trapped and crystallized upon introduction of a disulfide cross-link that covalently linked the D-loops of the two NBDs.

### **Distinct mechanistic proposals for B-family ABC exporters**

Despite a number of high-resolution structures reported, no common mechanism has been established for B-family ABC exporters. One problem is that most structures do not reveal bound substrates, a consequence of their hydrophobicity and low binding affinity in detergent solution. Another issue is that when removed from the membrane and in the absence of ATP (or ATP-ADP mixtures reflecting average cytoplasmic concentrations), B-family ABC exporters are prone to adopting inward-facing conformations with occasionally very large separations of the NBDs. When bound to nucleotides, the conformations tend to show a smaller separation of the NBDs as well as occluded or outward-facing conformations. Figure 3 shows a selection of structures of B-family ABC exporters, demonstrating the wide conformational range.

Early mechanistic proposals stated that alternating access in B-family ABC exporters is reached by converting the observed, inward-facing conformations to outward-facing states with bound ATP. However, several crystal structures are inconsistent with this simplified view<sup>54-58</sup>. The relevance of the observed inward-facing conformations has therefore been controversially discussed (see **box 2**), and the discussion is not simply about semantics. The critical issue is whether an observed structure reflects a functionally relevant state. If it is concluded that a conformation is not physiologically relevant (as were the inward-facing conformation of the LLO flippase PglK), there would be no point in attempting to dock a transport substrates into the observed cavities. The two extreme views outlined in **box 2** could be reconciled by viewing B-family ABC exporters as springs, whose architecture works to push their NBDs apart. In cellular membranes, the combined effects of lipids, ATP, and substrate

would allow the transporters to reach outward-facing conformations. Inward-facing conformations might reflect physiological states if they provide substrate binding sites or may occur transiently because the NBD dimer needs to release the ATP hydrolysis products. Inward-facing conformations without NBD contact or substrate binding pockets are likely unphysiological.

As was observed for ABC importers, substrates and inhibitors can modulate the ATPase activity of B-family ABC exporters. The molecular basis of this phenomenon is unknown, but stimulation or inhibition is often used to identify specific interactions. Two distinct mechanisms of B-family ABC exporters are shown in Figure 4.

**Alternating access.** An essential component of this mechanistic proposal (Fig. 4a) is the binding of transport substrate to an inward-facing pocket of the TMDs (state 1). The relevant binding site may not only be accessible from the cytoplasm, but also from the inner leaflet of the lipid bilayer. Highly hydrophobic substrates (drugs, phospholipids) could thus be directly recruited from within the membrane. The inward-facing conformation is linked to an open NBD conformation. As for ABC importers, futile ATP hydrolysis may occur in the absence of substrate, albeit at a slower rate than when substrate is bound. For poly-specific transporters (including the multidrug transporter ABCB1), it is unclear if there is only one binding pocket<sup>59</sup>, as some findings suggest multiple sites<sup>60,61</sup>. In the case of TAP, it was concluded that the N- and C-termini of all transported peptides are firmly bound, whereas the central aminoacids may only weakly interact with the transporter<sup>62</sup>. Once a substrate is recruited, the transporter probably converts to an occluded conformation (state 1 to 2), with ATP bound between closed NBDs, but without access to the central cavity from either side of the membrane. Such a conformation was observed in the structures of AMPPNP-bound McjD<sup>63</sup> and ATP<sub>γ</sub>S-bound PCAT<sup>57</sup>, albeit without bound substrate. It is unclear if the occluded conformation is a mandatory intermediate. The following step (state 2 to 3) reflects the conversion to the outward-facing conformation, allowing substrate release into the external medium or into the outer leaflet of the lipid bilayer. Given that there is a re-arrangement of the TM helices from states 1 to 3, with different sets of helices surrounding the translocation pathway and forming the two external wings (Fig. 4), there is a change in the cavity surface surrounding the substrate. In the AMPPNP-bound, outward-facing Sav1866 structure, a rather hydrophilic cavity was observed, suggesting that a chemical mismatch to the hydrophobic drugs could be driving substrate release<sup>64</sup>. This notion was in line with observations that the apparent affinity of multidrug ABC transporters for substrates was lower from the external than from the cytoplasmic side<sup>65</sup>. If a substrate is highly hydrophobic (such as in the case of certain tumor drugs or phospholipids), it is likely to re-partition into the membrane and may only remain in the external medium when bound to a carrier protein (e.g. BSA) and swept away in the blood stream, or if stabilized in a mixed micelle such as in bile. In the final step (state 3 to 1), hydrolysis of ATP and the release of inorganic phosphate drives the NBDs apart and, by transmitting this motion to the TMDs via the coupling helices, converts the outward-facing into an inward-facing conformation. This step is irreversible given the large amount of energy gained from the hydrolysis of ATP (approaching 50kJ mol<sup>-1</sup> for one ATP). It is unclear if the hydrolysis of two ATP molecules is simultaneous or consecutive, nor is it certain that both ATP molecules are hydrolyzed during every transport cycle (see below).

**Outward-only mechanism.** This mechanism was proposed for the LLO flippase PglK<sup>66</sup> (Fig. 4b). As its name implies, the outward-only mechanism does not invoke an inward-facing cavity to interact with the substrate. Although two distinct inward-facing conformations were determined for PglK, the observed cavities did not appear to contribute to the interaction with the substrate. The cycle of the outward-only mechanism can be thought to start in an ATP-bound state and an outward-facing TMD conformation (state 1). Whereas the lipidic polyprenyl tail of the substrate probably remains attached to the lipid-facing surface of the transporter, the pyrophosphate-glycan head-group is proposed to be transferred directly into the outward-facing cavity, where strong, electrostatic interactions between

arginines and the pyrophosphate group are formed (state 1 to 2). Once the head-group is bound in the outward-facing cavity, ATP hydrolysis and the release of inorganic phosphate drives the NBDs apart and the external wings of the TMDs together (state 3). The extrusion of the substrate appears essential to avoid a steric clash, similar to the peristaltic component proposed for type 2 ABC importers. The important concept is that upon release of the LLO head-group, the transporter adopts a conformation with insufficient size to fit a substrate. In the final step (state 3 to 1), dissociation of the substrate, re-binding of a new substrate, and exchange of ADP for ATP resets the system. Unlike the alternating-access mechanism, the outward-only mechanism does not rely on a large separation of NBDs, although a certain opening of the sandwich dimer is required for phosphate release and nucleotide exchange. The mechanism appears suitable to rationalize not only the flipping of LLOs, but the export of very large substrates (e.g. proteins or long polysaccharides, see also below), as repeated ratcheting through inward-facing conformations while constantly hydrolyzing ATP molecules appears sterically impossible. It is particularly attractive to consider the outward-only mechanism not only for lipid-linked oligosaccharides, but also for phospholipid and cholesterol export (transported by ABCB4, ABCG5/G8, and potentially ABCG1 and ABCA1).

It should be noted that the alternating access and outward-only mechanisms are not mutually exclusive and may be used by a single transporter for different substrates. For example, one could imagine that a multidrug ABC transporter might employ the alternating-access mechanism for certain substrates while using the outward-only mechanism for others.

The mechanistic proposals shown in Fig. 4 are generic and there are multiple variations and complications. Arguably the most important is the fact that some B-family ABC exporters contain only one consensus ATP binding site, while the other site is formed by degenerate motifs that reduce or abolish ATP binding or hydrolysis at that site. Transporters with published structures include Tm287/8, TmrAB, and TAP<sup>55,56,67,68</sup>. Of those, Tm287/8 has been resolved at sufficiently high resolution to allow side chain placement, and although a coupling of the ATPase sites via the D-loops was proposed, a definitive role of the asymmetry for the transport mechanism was not established<sup>55,56</sup>. EPR studies comparing a homodimeric ABC exporter (MsbA, containing two functional ATPase sites) with a heterodimeric one (BmrAB, containing one functional, one degenerate site) concluded that the presence of a degenerate ATPase site may prevent a wide opening of the NBDs in BmrAB, possibly indicating a fundamental difference between symmetrical vs. asymmetrical transporters<sup>69</sup>. It should be noted that ATPase asymmetry may not be restricted to heterodimeric ABC transporters. Even ABC transporters with two functional, consensus ATPase sites may reach an asymmetric state<sup>70-72</sup> or even hydrolyze only one ATP molecule per transport cycle. The asymmetry of ATPase sites extends beyond the B-family of ABC transporters: The cystic fibrosis transmembrane conductance regulator (ABCC7) also features a degenerate site in addition to the consensus site and it is currently assumed that binding of ATP to the degenerate site of CFTR is essential for chloride conductance, even though nucleotide is only hydrolyzed at the consensus site<sup>73</sup>. Another detail omitted in Fig. 4 is the role of the D-loops: A replacement of the conserved D-loop aspartates was found to convert TAP from an active transporter into a passive facilitator, suggesting a role in the coupling of ATP hydrolysis to vectorial transport<sup>74</sup>. The structural basis of this observation remains to be established, and it is currently unknown if other ABC transporters have similar phenotypes when their D-loops are mutated.

### **Mechanisms: Does one size fit all?**

A number of attempts have been made to formulate generally applicable mechanisms for ABC transporters. These include, among others, the "switch model", the "constant contact model", or the "reciprocating twin-channel model"<sup>38,75</sup>. However, given the structural and biochemical differences discussed above, mechanistic similarities in the entire superfamily appear restricted to the events at

the NBDs. This is in line with the observation that not all ABC transporters have the same phylogenetic origin<sup>76,77</sup>. Rather, distinct TMD folds appear to have evolved separately. Because the different TMDs all feature coupling helices, they can interact with NBDs to harness the conformational power provided by ATP hydrolysis and facilitate import or export in a given ABC transporter. The TMDs contain distinct binding sites adapted to recognize substrates of highly diverse mass and chemical nature and feature gates at different locations in the TMDs. In conclusion: No one size fits all. Rather than attempting to formulate unified transport mechanisms, I believe the goal must be to define the specifics of every ABC system under study. This requires a combination of structural, biochemical, and biophysical (including single molecule spectroscopic<sup>78</sup>) studies, and ultimately computational analysis, which, despite progress with select systems<sup>79-82</sup>, has thus far struggled with the fact that ABC transporters are very slow enzymes, with turnover rates approaching seconds in some cases.

### **ABC transporter research: Golden days ahead**

Significant progress in understanding reaction mechanisms has been made since the first structures of ABC transporters were determined. However, many details of how these transporters work remain to be discovered, and this search may yield surprises and unexpected findings even for B-family exporters. For example, new mechanistic concepts may have to be established for rationalizing phospholipid flipping by ABCB4. In addition, there are structurally uncharacterized ABC transporters that will likely reveal new TMD folds. These include the human G-family ABC transporters, but also bacterial ABC exporters of capsular polysaccharides<sup>83</sup>.

Some ABC transporters are part of larger cellular assemblies. For example, TAP is part of the endoplasmic peptide loading complex, where protein-protein interactions control and influence the transport activity. SUR1 and SUR2 (ABCC8 and ABCC9) form complexes with inward-rectifying potassium channels<sup>12</sup>, generating assemblies approaching a megadalton in mass and featuring almost a dozen ATP-binding sites for fine regulation of channel activity. In bacteria, there are ABC transporters involved in the secretion of toxic hemolysins<sup>84</sup>, which is achieved by assembly of an ABC transporter (HlyB) with periplasmic and outer membrane components (HlyD and TolC). To gain insight into these larger assemblies, single particle cryo-electron microscopy will likely be indispensable. However, given the recent resolution revolution in single particle cryo-EM<sup>85</sup>, even single ABC transporters may soon be studied at high resolution without the need to grow three-dimensional crystals. At present, the resolution of the cryo-EM structures of two ABC transporters is in the range of 6-8Å (TmrAB, Tap)<sup>67,68</sup>, but further developments in technology and sample preparation may soon allow resolutions sufficient for side chain placement to be obtained. This will have a large impact on the field, as transport intermediates and samples that have not yielded to crystallization efforts may be visualized, allowing biochemists and structural biologists to probe further into the intricacies of ABC transporter reactions. It's an exciting time indeed for ABC transporter research.

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### Box 1: Architectures and conserved motifs of ABC transporters

ABC transporters contain four core domains, two cytoplasmic NBDs and two TMDs. The NBDs point their conserved motifs towards each other, which generates two ATP-binding sites between the P-loops of one NBD and the signature motif of the other. In certain heterodimeric transporters (and in some full transporters), there are degenerate ATPase sites, where key sequence motifs carry mutations. Although nucleotide binding to these degenerate motifs may be preserved, hydrolysis at the modified ATPase sites is severely inhibited or abolished (reviewed in <sup>86</sup>).

The TMDs form a single, central pathway for their cognate substrate(s). Whereas NBDs are structurally similar, the folds of TMDs are diverse (reviewed in <sup>1,87,88</sup>). In bacterial ABC importers, TMDs are expressed as separate protein subunits that either belong to type I or type II based on whether the fold is similar to that first observed in *A. fulgidus* ModBC<sup>41</sup> (type 1) or *E. coli* BtuCD<sup>53</sup> (type 2). Recently, ECF-type transporters have revealed another fold that can be counted to the ABC importer family. ECF-type transporters facilitate high-affinity uptake of certain vitamins into bacteria. Although shown in Fig. 1 for completeness, they will not be discussed here, and the reader is referred to recent reviews<sup>89,90</sup>. There is currently only one structurally defined fold of ABC exporters, first revealed by the bacterial Sav1866 protein<sup>54</sup> and subsequently found in all reported crystal structures of ABC exporters<sup>34,55,57,58,63,66,91-96</sup>. It will be referred to as the B-family ABC exporter fold. In this fold, the NBDs are fused to the C-termini of TMDs, which generates half-transporters that assemble to full transporters by forming homodimers or heterodimers. In some B-family ABC exporters, all four domains are fused to a single polypeptide (e.g. in ABCB1 or ABCB4).

An important observation was that although the three TMD folds discussed here show no structural similarity, they all feature a short, cytoplasmic helix roughly parallel to the membrane plane and facilitating critical contacts with the NBDs (Fig. 1b). Due to its key role transmitting ATP-driven conformational changes from the NBDs to the TMDs, it has been termed "coupling helix"<sup>54,97</sup>. Coupling helices allow distinct and evolutionarily unrelated TMD folds to harness the conformational power delivered by NBDs.

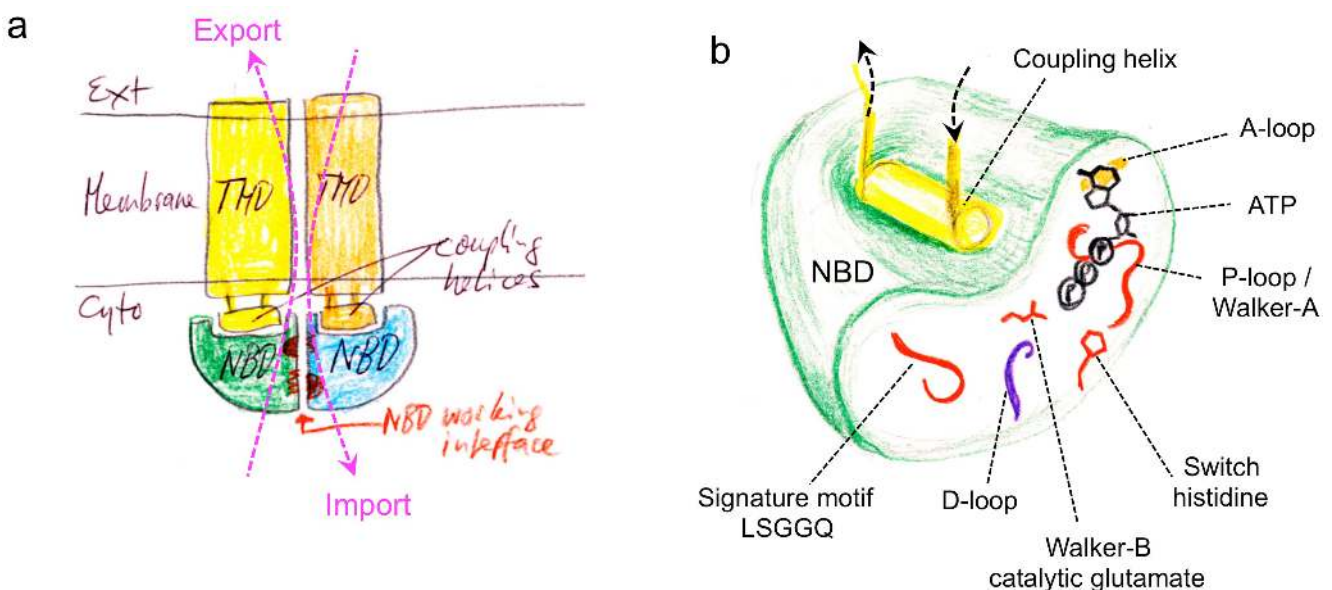


Figure legend box 1: **a**, Domain arrangement of ABC transporters: At the NBD interface, red half-circles and lines indicate the P-loops and ABC signature motifs, respectively. Any given ABC transporter is either an importer or an exporter. Coupling helices transmit conformational changes from the NBDs to the TMDs and vice versa. **b**, Schematic of a single NBD. The location of the conserved and functionally critical motifs are indicated and labeled. They include the P-loop (or Walker-A motif) that binds the  $\alpha$ - and  $\beta$ -phosphates of ATP, the A-loop that provides an aromatic side chain that packs against the purine ring of adenine, a Walker-B motif providing the catalytic glutamate, a signature LSGGQ motif that pins and orients ATP during hydrolysis, a "switch histidine" that stabilizes the transition state geometry, a Q-loop providing contacts to the TMD (not shown), and a dimerization or D-loop that has a role in coupling hydrolysis to transport (reviewed in <sup>87</sup>). A groove in the NBD surface forms the contact interface with the coupling helix of the TMD. While the coupling helix is not the only contact between TMDs and NBDs, it is the only architecturally conserved one among distinct TMD folds and provides the majority of contacts between domains.

## Box 2: Inward-facing conformations in B-family ABC exporter structures

The physiological relevance of inward-open conformations with wide separation of the NBDs is controversially discussed. Those arguing in favor would point out that extreme conformational flexibility has been observed in certain transporters by DEER-EPR<sup>69,98</sup>. Furthermore, the exchange of ADP for ATP following a hydrolysis cycle requires an apo-state to be populated, even if only briefly. In addition, inward-facing conformations (and to some degree NBD separation) are required if alternating access is postulated, because an inward-facing cavity would form the substrate binding site. The only structures of B-family ABC exporters with bound transport substrates are those of ScAtm1 and the bacterial homologue NaAtm1<sup>93,95</sup>. The two structures are very similar and revealed nucleotide-free, inward-facing conformations with substrates (GSH and GSSG) bound to an inward-facing cavity at the level of the inner membrane leaflet. In both structures, however, there is physical contact between the NBDs, afforded through long helices at the C-terminal ends of the NBDs.

Those arguing against the functional relevance argue that the absence of nucleotides or the presence of ADP alone used in structural and spectroscopic studies do not reflect physiological conditions, because the cytoplasm of viable cells contains multi-millimolar concentrations of ATP, as well as ADP at about a tenth of the ATP concentration. Apo-states or ADP-states are therefore likely short-lived and transient *in vivo*. Furthermore, many B-family exporters require a lipid bilayer for substrate-stimulated ATP hydrolysis, whereas only low (and often non-stimulated) ATPase rates are observed in detergent. Moreover, P-glycoprotein has been found to be functional even when its two halves were cross-linked and a wide, inward-facing conformation as observed in crystal structures was prevented<sup>99</sup>. Also, separating the NBDs has been demonstrated to inhibit the activity of B-family ABC exporters: ABCB1 is inhibited by a nanobody between the NBDs<sup>34</sup> while TAP is inhibited by a viral peptide wedged between the TMDs<sup>68</sup>, also preventing NBD dimer closure. The LLO flippase PgIK is currently the only B-family exporter crystallized in two distinct apo-states, revealing inward-facing conformations and NBD separations that differ in distance and relative orientation<sup>66</sup>. This argues against a functional relevance of the, inward-facing conformations. Rather, lattice contacts and detergent effects can dictate the degree of NBD separation.

Figure 1

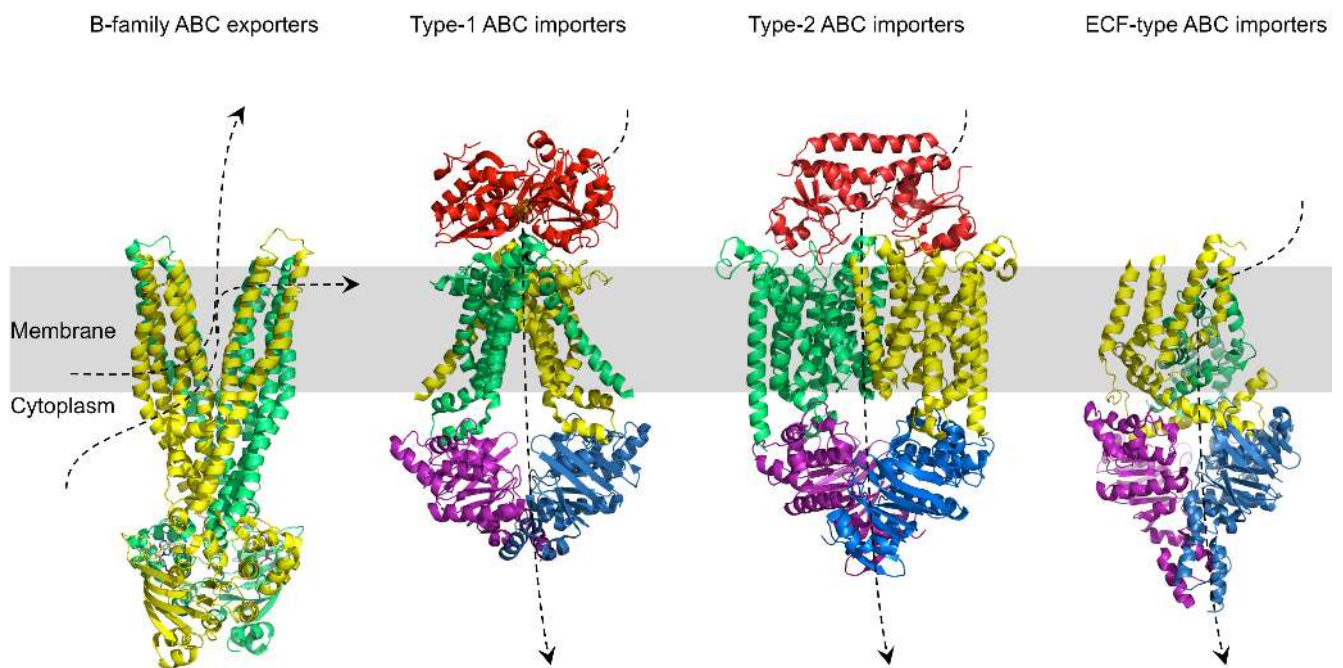


Figure 1: ABC transporter structures. The molecules shown in the figure defined the TMD folds of the indicated sub-families: *Staphylococcus aureus* multidrug transporter Sav1866<sup>54</sup> (B-family ABC exporters); *Archaeoglobus fulgidus* molybdate / tungstate transporter ModBC-A<sup>41</sup> (type 1 ABC importer); *Escherichia coli* vitamin B12 transporter BtuCD-F<sup>53</sup> (type 2 ABC importer); *Lactobacillus brevis* folate importer EcfAST<sup>100</sup> (ECF-type ABC importer). TMDs are colored yellow and green, periplasmic substrate binding proteins are in red, NBDs in ABC importers are purple and blue. Note that the TMDs and NBDs are fused in B-family exporters. Whereas bacterial B-family ABC exporters are homodimeric or heterodimeric, many eukaryotic homologs contain all four domains on a single polypeptide chain. Dashed arrows indicate the known or suspected substrate paths during translocation. One of the TMDs in ECF-transporters is variable, and there are no coupling helices similar to those of canonical ABC transporters.



Figure 2

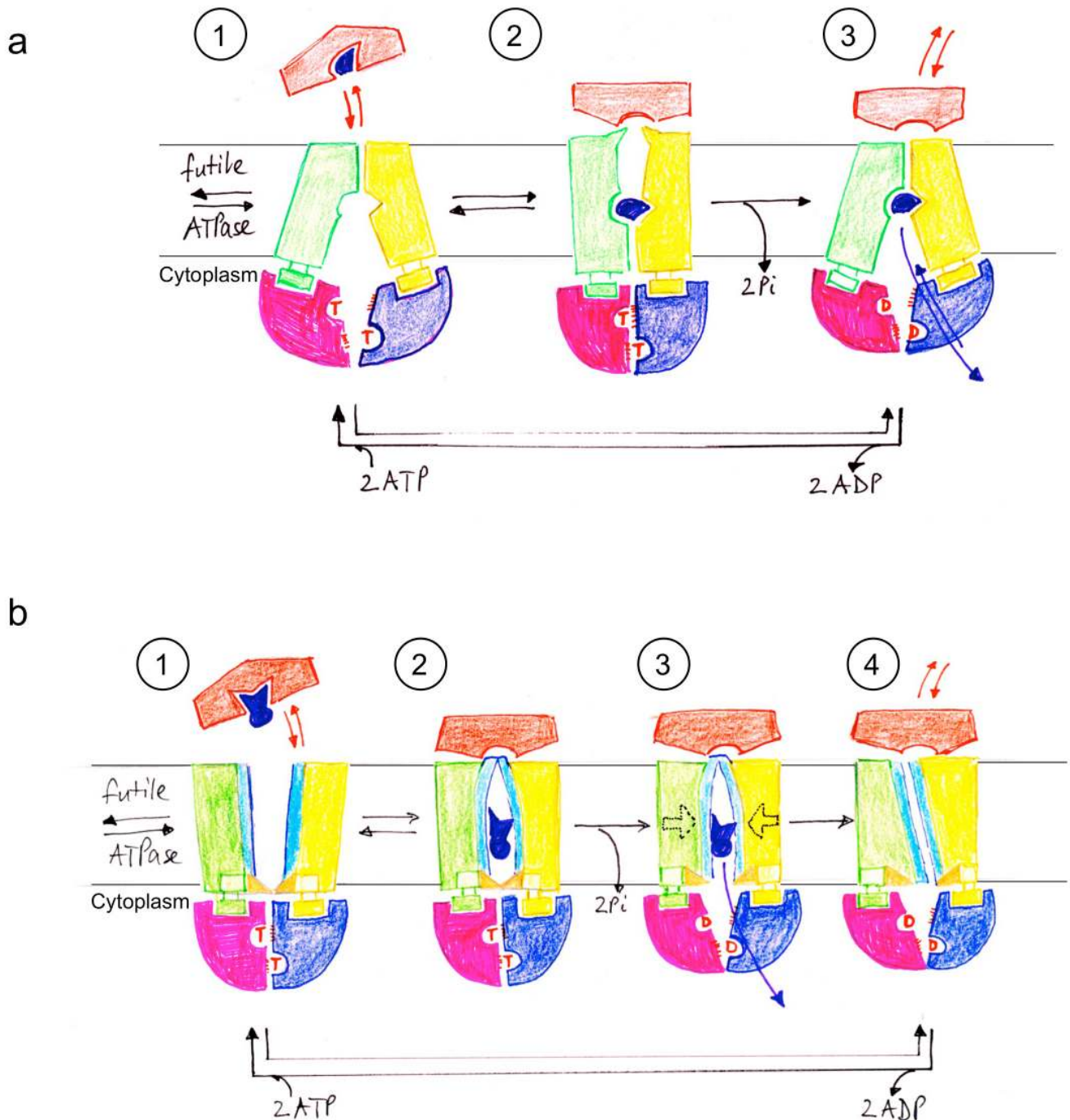


Figure 2. Distinct mechanisms of type 1 and type 2 ABC importers. TMDs are colored green and yellow, NBDs are colored pink and purple, binding proteins are colored orange, with substrates in dark blue. "T" and "D" denote ATP and ADP, respectively. Dotted red lines in the NBDs depict the ABC signature motifs. Numbers above schematics indicate states. See text for detailed explanations. **a**, Schematic of a type 1 importer importer mechanism. **b**, Schematic of a type 2 importer containing three distinct gates. The blue lines in the TMDs represent TM5 forming cyto-gate 1 and the peri-gate, whereas the orange triangles represent cyto-gate 2, formed by the loop between TM2 and TM3. Small arrows in TMDs of state 3 indicate forces acting on the translocation pathway and entrapped substrate.

Figure 3

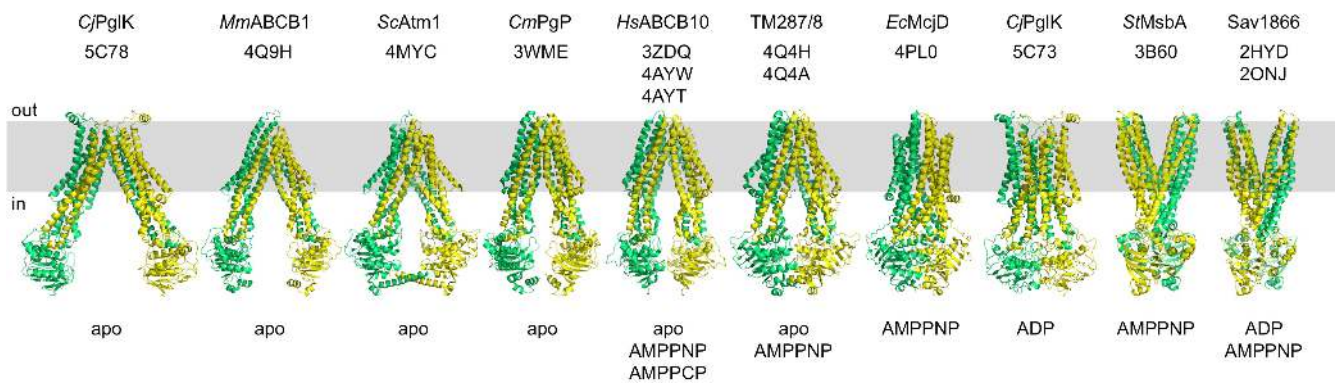


Figure 3: Conformations in selected B-family ABC exporter structures. Species abbreviations and protein names are shown in the top row, relevant pdb codes above the ribbon diagrams, bound nucleotides below. "In" refers to the cytoplasmic side of the membrane. If multiple nucleotide states are indicated (e.g. apo and AMPPNP), the relevant structures did not differ significantly in terms of NBD separation.

Figure 4

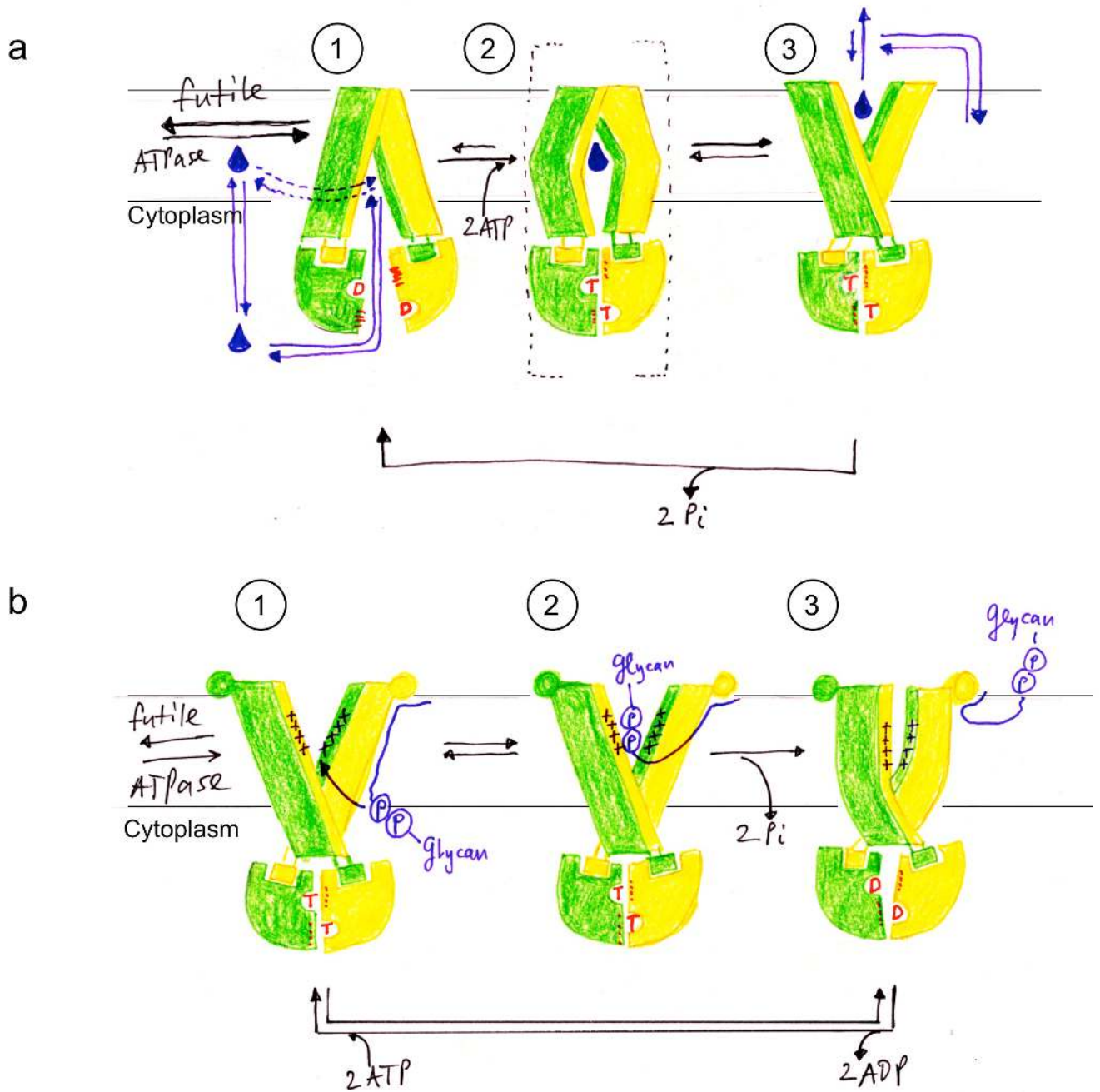


Figure 4: Distinct mechanisms of B-family ABC exporters. Half-transporters are colored green and yellow. "T" and "D" denote ATP and ADP, respectively. Dotted red lines in the NBDs depict the ABC signature motifs. Circled numbers denote states. In outward-open conformations, the TMDs of the transporters form two wings towards the external side of the membrane. Each wing contains two TM helices from one TMD and four TM helices from the other. See text for details. **a**, Alternating access proposed for multidrug or peptide transporters. **b**, Outward-only mechanism with substrate loading directly into outward-facing cavity as proposed for LLO flippase PglK.