

## STRUCTURAL BIOLOGY

# Structural snapshots of human PepT1 and PepT2 reveal mechanistic insights into substrate and drug transport across epithelial membranes

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The uptake of peptides in mammals plays a crucial role in nutrition and inflammatory diseases. This process is mediated by promiscuous transporters of the solute carrier family 15, which form part of the major facilitator superfamily. Besides the uptake of short peptides, peptide transporter 1 (PepT1) is a highly abundant drug transporter in the intestine and represents a major route for oral drug delivery. PepT2 also allows renal drug reabsorption from ultrafiltration and brain-to-blood efflux of neurotoxic compounds. Here, we present cryogenic electron microscopy (cryo-EM) structures of human PepT1 and PepT2 captured in four different states throughout the transport cycle. The structures reveal the architecture of human peptide transporters and provide mechanistic insights into substrate recognition and conformational transitions during transport. This may support future drug design efforts to increase the bioavailability of different drugs in the human body.

## INTRODUCTION

The plasma membrane forms a natural barrier for amino acids, short peptides, and other hydrophilic or charged nutrients. To preserve the distinct intracellular milieu, a large number of membrane transporters for these molecules have emerged during evolution to maintain the nutrient homeostasis of cells. For efficient uptake of individual amino acids and small peptides, specific amino acid transporters together with the promiscuous peptide transporter 1 (PepT1) are expressed in the mucosa of the small intestine (1). PepT1 belongs to the solute carrier family 15 (SLC15), also known as the proton-coupled oligopeptide transporter (POT) family, which consists of four members in eukaryotes: PepT1 (SLC15A1), PepT2 (SLC15A2), PhT1 (SLC15A4), and PhT2 (SLC15A3). PepT1 and PepT2 are best characterized and mediate the uptake, distribution, and resorption of di- and tripeptides in the body. These transporters are highly promiscuous and accept almost any di- and tripeptide, independent of their side-chain composition, but with substantial differences in affinity (2–6). PepT1 is the predominant paralog in the apical membrane of the intestinal epithelial cells, while PepT2 has a broad expression pattern and is mainly found not only in the kidney but also in various other tissues including the brain, neurons, lung, and choroid plexus (7–10).

PepT1 and PepT2 are secondary active transporters, which are energized by the inward-directed electrochemical proton gradient. This provides a driving force for transport and accumulation of nutrients above extracellular concentrations (11, 12). Besides natural di- and tripeptides, PepT1 and PepT2 recognize and transport chemically diverse drug molecules such as  $\beta$ -lactam antibiotics,

angiotensin-converting enzyme inhibitors, and antiviral drugs, thus affecting their availability, clearance, and distribution in the body (13–19). PepT1 accounts for ~50% of all known clinically relevant drug transporters in the small intestine and represents one of the main routes for oral drug absorption (20). PepT2 reduces the clearance of exogenous molecules via renal tubular reabsorption (21) and enables drug efflux from the cerebrospinal fluid to the choroid plexus, thus influencing drug disposition, dynamics, and toxicity in the brain (22–24). In human disease, colonic expression of PepT1 leads to bacterial di- and tripeptide uptake in epithelial cells, causing downstream chronic inflammation and is associated with numerous gastrointestinal tract disorders including inflammatory bowel disease (IBD) and colonic cancer (25, 26). PepT1-mediated uptake of tripeptides has been shown to reduce nuclear factor  $\kappa$ B (NF- $\kappa$ B) and mitogen-activated protein (MAP) kinase inflammatory signaling pathways and proinflammatory cytokine secretion and reduced the incidence of colitis in mice, raising the use of anti-inflammatory oligopeptides as attractive therapeutic strategy against IBD (27–30).

On a structural level, human PepT1 (*HsPepT1*) is 708 and human PepT2 (*HsPepT2*) is 729 amino acids long. Both polypeptides consist of a core transporter unit of predicted 12 transmembrane helices (TMs) of the major facilitator superfamily (MFS) fold and an extracellular immunoglobulin-like domain (ECD) placed between TM9 and TM10. *HsPepT1* and *HsPepT2* share overall high sequence similarity (>70%), which is even higher for the transporter core units (>85% sequence similarity, ~65% sequence identity), and a highly conserved substrate binding site (fig. S1). Currently, it is postulated that substrate transport occurs via the “rocker-switch” alternating access mechanism, which involves conformational transitions between at least three different states: (i) outward open, (ii) occluded, and (iii) inward open (31).

Structures of different bacterial POT homologs have been determined over the past 10 years in apo, peptide-bound, and drug-bound states, exclusively representing inward-facing open or inward-facing partially occluded structures (32–45). This strongly limits our molecular understanding of the conformational transitions, required to complete an entire transport cycle. In particular, it is not clear whether the available inward-facing structures of the binding site

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are representative of the outward-facing state, where substrate recognition occurs.

Here, we present multiple cryogenic electron microscopy (cryo-EM) structures of *HsPepT1* and *HsPepT2* representing different stages of the transport cycle of the SLC15 family. *HsPepT1* was captured in the outward-facing conformation, both in apo and substrate-bound states, and in a substrate-bound outward-facing occluded structure. *HsPepT2*, on the other hand, has been trapped in a substrate-bound inward-facing partially occluded state. This work reveals the architecture of human POTs, which differ from the bacterial homologs, and elucidates substrate coordination in the centrally located binding cavity. Because of the availability of different conformational states of these highly similar transporters, we obtained molecular insights in conformational changes occurring during the entire transport cycle required for substrate recognition and transport. Our work will form the basis for future drug design and modification approaches using peptide transporters as shuttle systems.

## RESULTS

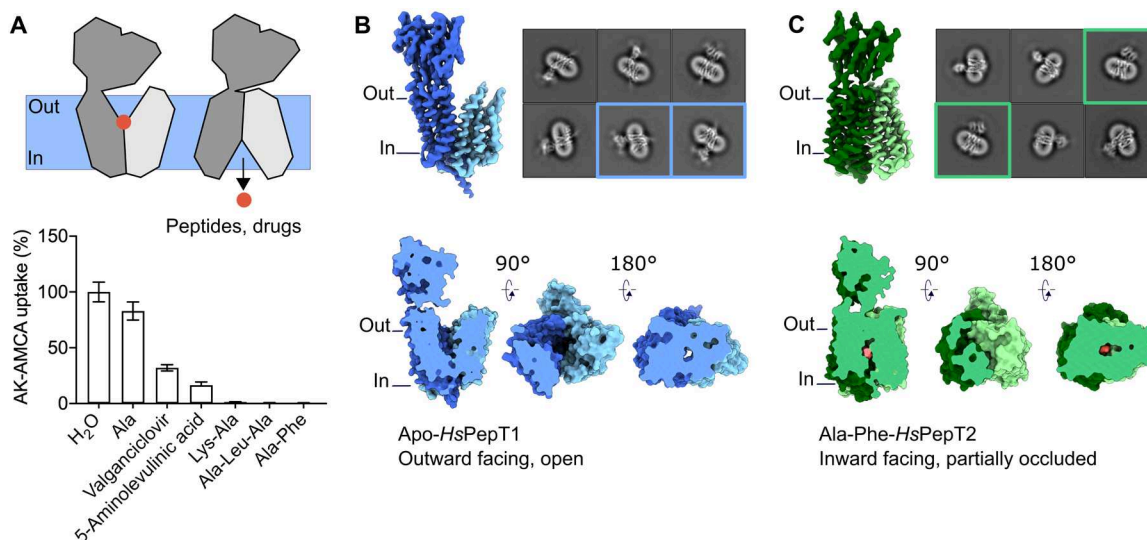
### Expression, purification, and structure determination of *HsPepT1* and *HsPepT2*

*HsPepT1* and *HsPepT2* were expressed in human embryonic kidney (HEK) 293 cells. To monitor transport, we made use of the fluorescently labeled dipeptide ( $\beta$ -Ala-Lys-AMCA) (46) and confirmed specific uptake of this compound after *HsPepT2* overexpression (Fig. 1A and fig. S2). Different di- and tripeptides efficiently compete for the same binding site resulting in reduced uptake of the fluorescent reporter. The same expression system has been used for *HsPepT1* transport assays in the past (47). Initially, we reconstituted *HsPepT2* in saposin-lipid nanoparticles to better mimic a lipid bilayer (48, 49). This approach yielded a homogeneous sample, but after grid preparation and imaging, it became clear that the particles adopted a preferred orientation on the cryo-EM grid (mainly top views) resulting in a poorly interpretable volume (fig. S3, C to F).

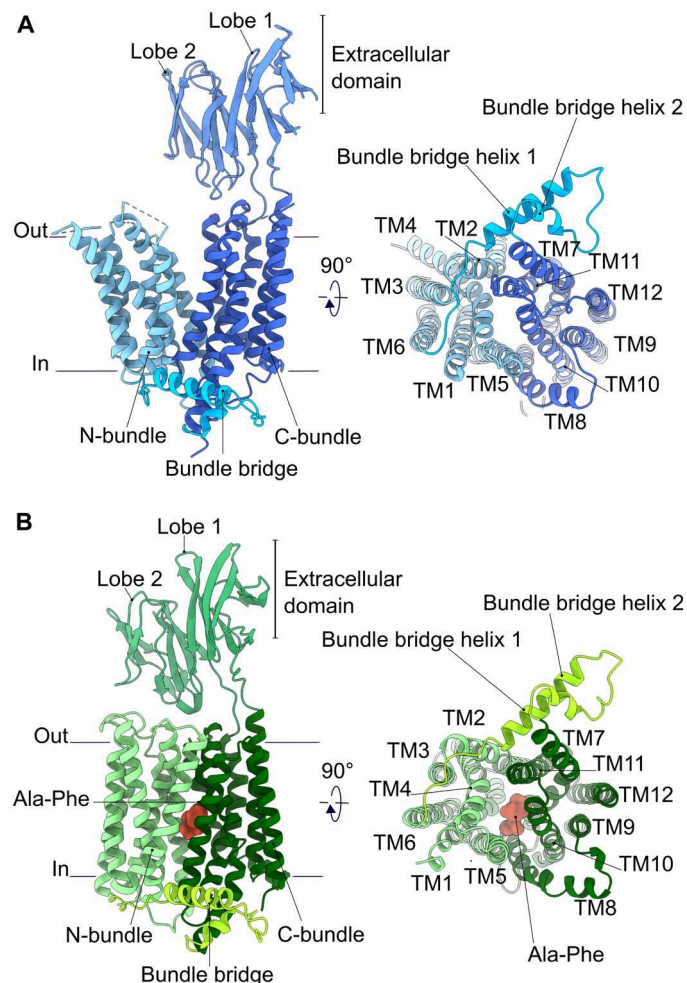
Therefore, we imaged monomeric full-length *HsPepT1* and *HsPepT2* extracted in detergent (Fig. 1 and fig. S3). In a first step, the structure of *HsPepT1* was investigated in its apo form, while *HsPepT2* was preincubated with one of its natural substrates. Distinct conformational states were already noticeable from two-dimensional (2D) class averages between the two paralogs, with *HsPepT1* representing an outward-facing conformation and *HsPepT2* representing an inward-facing state. We obtained 3D reconstructions for apo *HsPepT1* at a nominal resolution of 3.9 Å (fig. S4) and for *HsPepT2* bound to the dipeptide Ala-Phe at a nominal resolution of 3.8 Å with an estimated local resolution of up to 3.2 Å within the transporter unit (fig. S5 and table S1).

### Architecture of human peptide transporters

The 3D reconstructions allowed de novo modeling of most of the MFS transporter units, which consist of 12 TMs and a long linker connecting both helical bundles (figs. S6 and S7). Concerning the *HsPepT1* structure, 11 residues at the N terminus, four loops connecting TM1-TM2 (residue 49), TM3-TM4 (residues 109 to 116), TM5-TM6 (residues 189 to 194), and TM11-TM12 (residues 638 to 641), and the last 25 residues at the C terminus could not be modeled because of poor density likely caused by intrinsic dynamics and partial disorder. The *HsPepT2* model is lacking the first 40 residues at the N terminus and the last 32 C-terminal residues, which are predicted to be disordered. The transmembrane domains of *HsPepT1* and *HsPepT2* adopt the canonical MFS fold formed by 12 TMs organized in two six-helix bundles with both N and C termini facing the cytoplasm (Fig. 2). The bundles are connected via a long cytoplasmic linker, which encompasses two  $\alpha$  helices that interact with each other. This linker, which we have termed the “bundle bridge,” is likely common for all SLC15 transporters (figs. S1 and S8). The bundle bridge is of amphipathic nature and is associated with the inner leaflet of the plasma membrane (fig. S8). Its role is currently unclear, but it might take part in lipid sensing or in providing a platform for other interacting proteins.



**Fig. 1. Cryo-EM structures of apo *HsPepT1* and *HsPepT2* bound to Ala-Phe.** (A) Whole-cell transport competition assays of the  $\beta$ -Ala-Lys peptide coupled to the fluorescent AMCA moiety (AK-AMCA) in *HsPepT2*-transfected HEK293 cells showing reduced AK-AMCA uptake in the presence of 5 mM of the competing substrate. (B and C) Three-dimensional reconstructions of (B) *HsPepT1* and (C) *HsPepT2* with corresponding 2D class averages and surface representation highlighting the (B) outward-facing open and (C) inward-facing partially occluded conformations.



**Fig. 2. Overall architecture of human POTs.** (A) Apo-*HsPepT1* and (B) substrate-bound *HsPepT2* models shown as cartoon representation. The different architectural elements are labeled. Loops that could not be modeled because of poor density are shown as dashed lines.

Within the MFS, mammalian PepT1 and PepT2 are the only known transporters with an additional ECD placed between TM9 and TM10. As the extracellular domain of *HsPepT2* was poorly resolved, the predicted structure from AlphaFold (50) was used for template-based model refinement in this region. The higher local resolution in *HsPepT1*-ECD allowed us to identify six N-linked glycans (fig. S9), five of which were experimentally confirmed to be present on murine PepT1 and are likely involved in protein folding, membrane targeting, and protection from proteolytic degradation (51). In the context of the full-length transporter, the arrangement of the two immunoglobulin lobes in *HsPepT1* differs strongly from the previously crystallized isolated murine soluble PepT1-ECD (52) but is very similar to *HsPepT2*-ECD. Although *HsPepT1* and *HsPepT2* display different conformational states, both extracellular domains are positioned similarly with respect to the C-terminal bundle, with small hinge movements in relation to the linker region, and no interaction is observed between the ECDs and the N-bundle. This observation is in agreement with previous work, highlighting that the ECD is not essential for substrate transport but potentially forms an interaction platform for proteases such as trypsin via a conserved

acidic motif (52). Nevertheless, it cannot be excluded that the presence of a lipid environment might have an influence on the position of the ECD with respect to the transporter unit. The overall structures of *HsPepT1* and *HsPepT2* reveal notable architectural differences between human and bacterial homologs, as illustrated in fig. S10. While *HsPepT1* and *HsPepT2* contain 12 TMs, an additional ECD, and the bundle bridge connecting the N- and C-terminal bundles, bacterial POT structures display two additional TMs (HA-HB) of currently unknown function and lack the soluble domain (32).

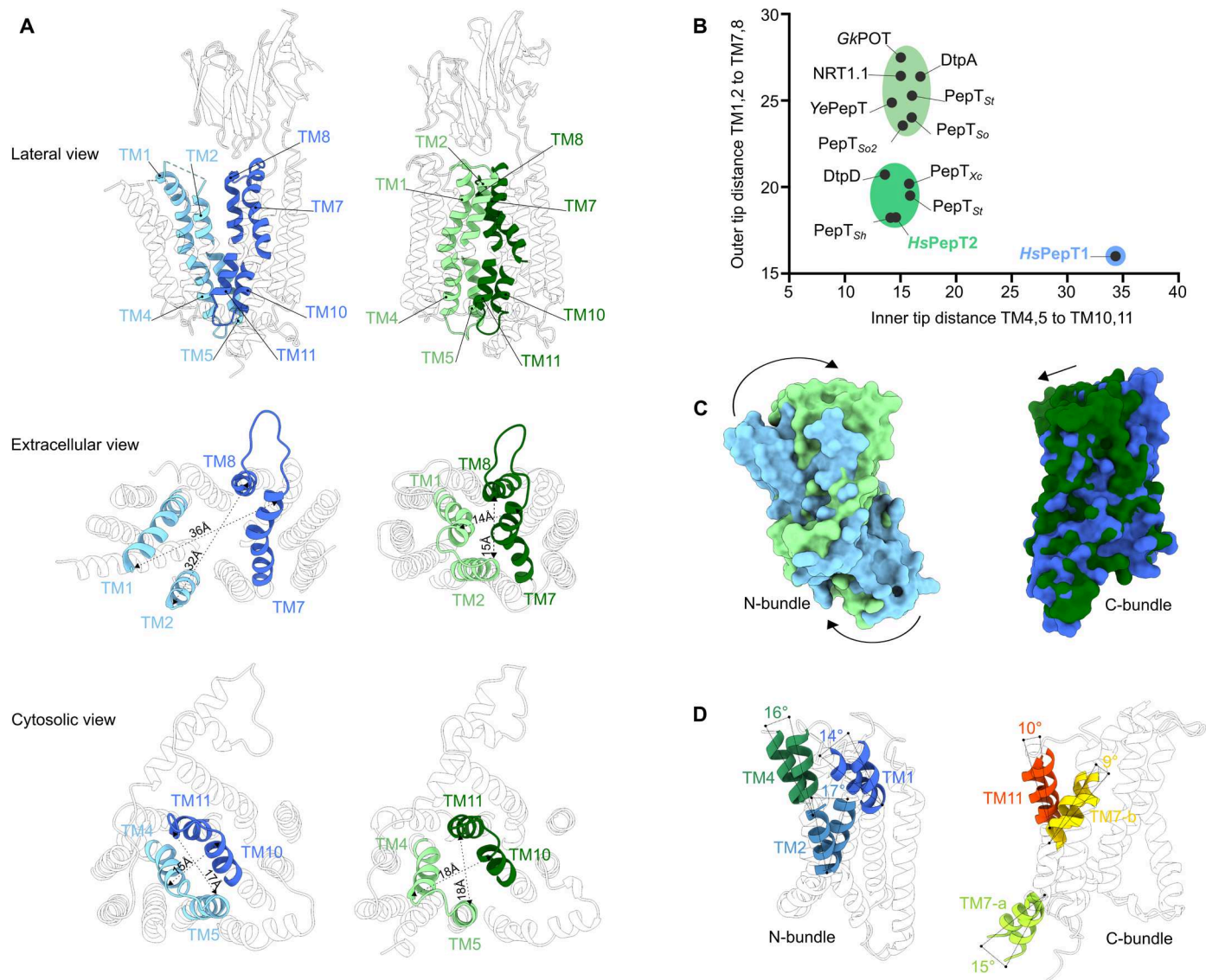
### Major conformational changes between inward- and outward-facing states

In the outward open state of apo *HsPepT1*, the central substrate binding cavity of the transporter unit is widely exposed to solute molecules from the extracellular space with an opening of approximately 30 Å (Fig. 3A). So far, among the available POT structures, no outward-facing conformation has been observed (Fig. 3B). This previously not captured state allowed us to observe the major structural changes occurring between outward- and inward-facing conformations. As the sequence conservation between *HsPepT1* and *HsPepT2* is high (>70% similarity), we superimposed their transmembrane domains to highlight the differences between these two states. The transition from *HsPepT1* outward-facing conformation to the inward-facing state observed in *HsPepT2* can be deconvoluted in two main processes: (i) a large rocking motion of the N-bundle coupled with a smaller movement of the C-bundle, carrying along the extracellular domain (Fig. 3C), and (ii) an additional bending of TM1, TM2, TM4, TM7, and TM11, allowing for a further opening and closing of the cytoplasmic and extracellular side, respectively (Fig. 3D).

In the *HsPepT2* structure, the cytoplasmic side ends of TM4 and TM5 in the N-bundle are in close proximity to TM10 and TM11 in the C-bundle, narrowing the exit route from the peptide binding site to the cytosol, although without fully closing it as instead was observed in outward-facing *HsPepT1* (Figs. 1C and 3A). Such a conformation is referred to as “partially occluded.” As previously shown in bacterial homologs studies, the transition from fully occluded to fully inward open conformation occurs via bending of TM10 and TM11 toward TM4 and TM5 (fig. S11) (34, 39, 41, 42). Upon switching to the outward-facing conformation, the modest rocking movement of the C-bundle translates the TM10-TM11 hairpin in closer proximity to TM4-TM5, resulting in tight sealing of the substrate binding pocket from the cytosol (Fig. 3A).

While a network of glutamine residues in TM7, TM9, and TM10 restricts conformational changes within the C-bundle in proximity to the extracellular domain in both *HsPepT1* and *HsPepT2*, other interactions between polar residues conserved among mammalian PepT1 and PepT2 are formed and broken between outward- and inward-facing conformations (Fig. 4 and fig. S1). In *HsPepT1*, the large bending of TM2 is stabilized by a hydrogen bond between Y64 (TM2) and N630 (TM11) in the extracellular region of the transporter unit. The sealing of the cytosolic side is stabilized by two interbundle salt bridges between R159 (TM5)–E604 (TM10) and R161 (TM5)–D341 (TM8) (Fig. 4A).

In the partially occluded inward-facing state of *HsPepT2*, these interbundle salt bridges on the cytosolic side are disrupted (Fig. 4B), allowing the large rocking movement of the N-bundle (Fig. 3C). In the upper part of the transporter unit, the interaction between Y94 in TM2 (Y64 in *HsPepT1*) and N657 (N630 in *HsPepT1*) is disrupted



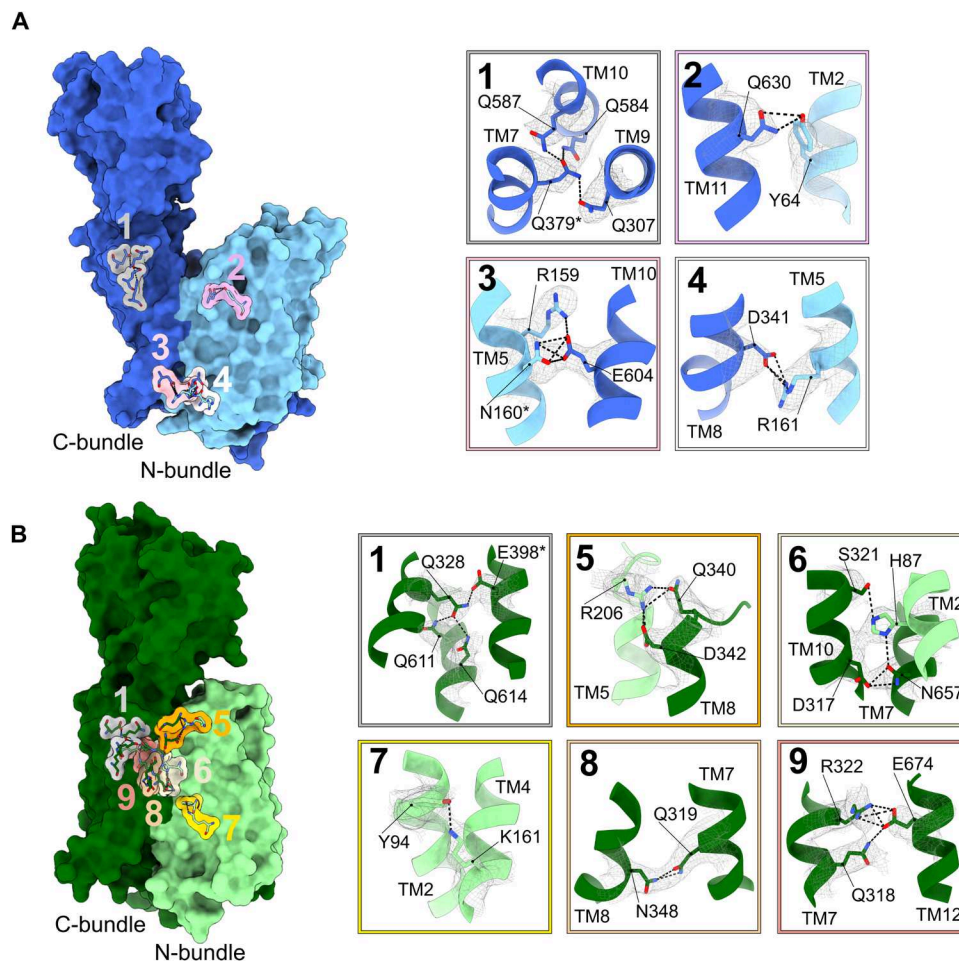
**Fig. 3. Structural comparison between the outward- and inward-facing states observed in apo *HsPepT1* and substrate-bound *HsPepT2*.** (A) Opening and closing of the substrate binding site to the extracellular and intracellular milieu observed in *HsPepT1* (blue) and *HsPepT2* (green). (B) The distances between C $\alpha$  atoms of the relevant pairs of helix tips from all bacterial POTs determined by x-ray crystallography were measured and compared to the human transporters. (C) Rocking motions of the N-bundle (*HsPepT1*: light blue and *HsPepT2*: light green) and C-bundle (*HsPepT1*: dark blue and *HsPepT2*: dark green) after structural alignment of both transporter units. (D) Bending of TMs with measured tilt angles observed in the N-bundle (left) and C-bundle (right) between *HsPepT1* and *HsPepT2*.

and replaced by an interaction between Y94 and K161 (TM4), restricting bending of TM2. Additional intrabundle contacts including Q319 (TM7)–N348 (TM8) and Q316–R322 (TM7)–E674 (TM12) further stabilize the partially occluded inward-facing state. Last, the interbundle salt bridge between R206 (TM5) and D342 (TM8), together with the polar interaction network around H87 involving TM2, TM7, and TM8, tightly seals the central cavity from the extracellular space. These interactions need to be disrupted so that the transporter can cycle back to the outward-facing state observed in *HsPepT1*. The interaction networks are illustrated in Fig. 4 (A and B). Mutational studies on *HsPepT1* identified H57 (H87 *HsPepT2*) as a critical residue for substrate translocation in both transporters (53), in agreement with our data presented here. This residue, as well as four of the five interbundle interaction networks mentioned above,

is conserved in mammalian POTs, but not in bacterial homologs (fig. S1). This might indicate an evolutionary divergence in the mechanism of conformational changes throughout the transport cycle of the POT family.

### Dynamics in the transport cycle induced upon substrate binding

Initially, we determined the structures of *HsPepT1* in the absence of a substrate and of *HsPepT2* in the presence of the dipeptide Ala-Phe. We used the fluorescently labeled dipeptide  $\beta$ -Ala-Lys-AMCA as a transport reporter in cell-based uptake assays to confirm binding of the dipeptide Ala-Phe, other peptides, and known drugs to *HsPepT2*. Concentration-dependent competition experiments yielded median inhibitory concentration (IC<sub>50</sub>) values in the micromolar range,

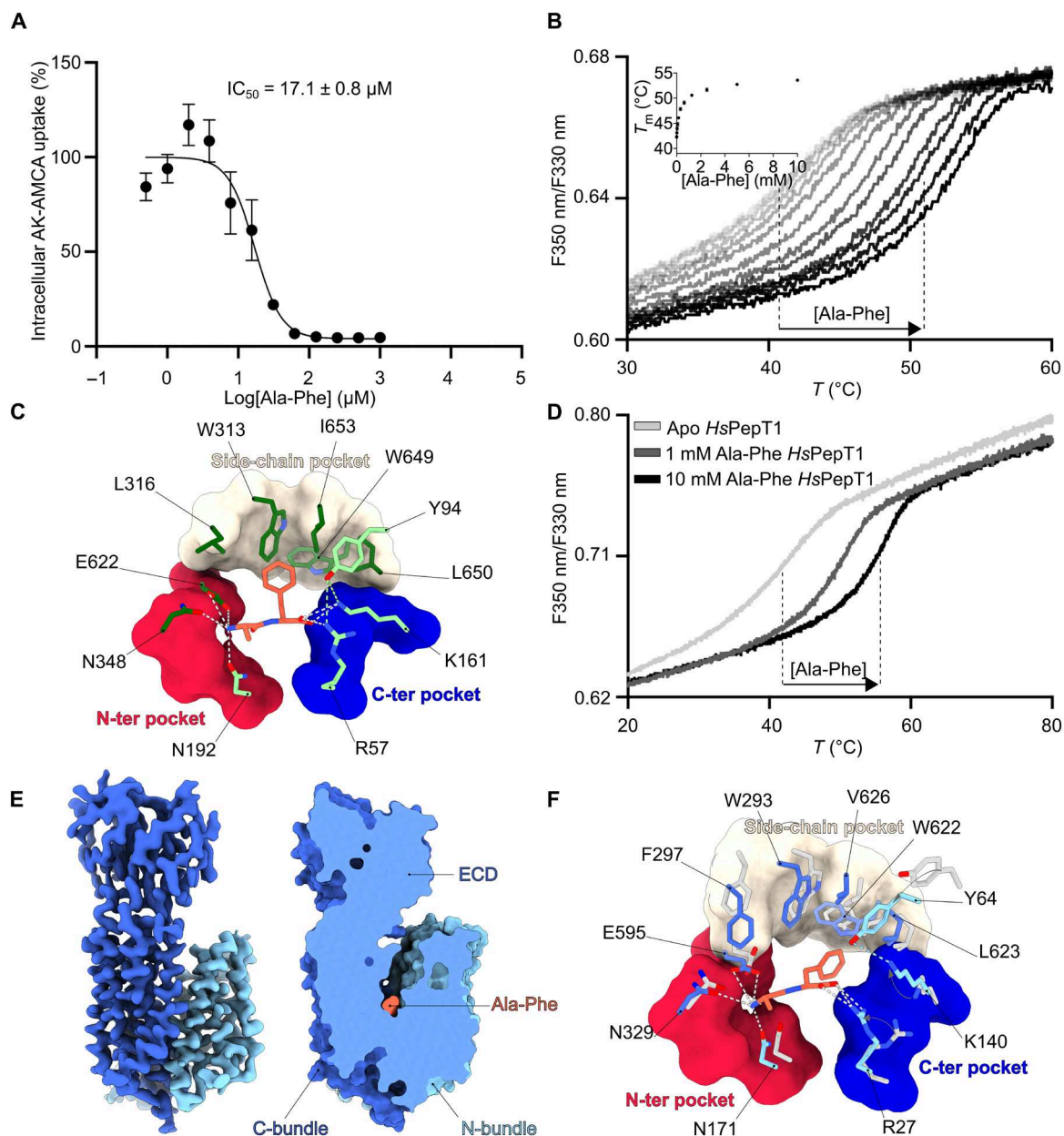


**Fig. 4. Interactions stabilizing the outward-facing open state of *HsPepT1* and the inward-facing partially occluded state of *HsPepT2*.** The locations of key interactions are shown and labeled on (A) *HsPepT1* and (B) *HsPepT2*. Corresponding close-up views show the cryo-EM densities of the side chains forming the interactions as indicated by dashed lines.

with naturally occurring substrates displaying higher affinities ( $IC_{50} = 17.1 \pm 0.8 \mu\text{M}$  to  $45.5 \pm 0.1 \mu\text{M}$ ) compared to the tested known transported drugs, such as valganciclovir or 5-aminolevulinic acid ( $IC_{50} = 368.9 \pm 1.5 \mu\text{M}$  and  $373.5 \pm 1.6 \mu\text{M}$ ) (Fig. 5A and fig. S2). Recombinantly expressed and purified *HsPepT2* used for structure determination was stabilized against heat unfolding in the presence of the dipeptide Ala-Phe in a concentration-dependent manner, confirming substrate binding of detergent extracted *HsPepT2* (Fig. 5B). The higher local resolution in the substrate binding site of *HsPepT2* enabled us to unambiguously assign the extra density to the dipeptide Ala-Phe and model all coordinating *HsPepT2* side chains (Fig. 5C and fig. S12A). The N and C termini of the dipeptide are clamped by electrostatic interactions with N192, N348, E622, R57, and K161, respectively. The phenyl group side chain is accommodated in a hydrophobic pocket formed by Y94, W313, L316, W649, L650, and I653, referred to as the side-chain pocket. The interactions between the peptide backbone and *HsPepT2* are in agreement with previous biochemical and mutational studies in mammalian POTs (13, 53, 54) and structural studies on inward-facing open or partially occluded states of bacterial homologs (33, 34).

Subsequently, to assess whether the switch from the outward-facing open state (as observed in the apo *HsPepT1* structure) to the

inward-facing partially occluded state (observed in the Ala-Phe-bound *HsPepT2* structure) had been triggered by substrate binding, we also imaged and determined the structure of *HsPepT1* in the presence of the same dipeptide, which also stabilized the transporter against heat unfolding in a concentration-dependent manner (Fig. 5D). However, even in the presence of the Ala-Phe dipeptide, the conformation of *HsPepT1* remained outward facing. Nevertheless, this time, we could differentiate between two distinct conformations in the same dataset, both different from the apo outward-facing open state (figs. S13 and S14). The data resulted in a lower resolution reconstruction for the substrate-bound outward-facing occluded state (nominal resolution of 4.1 Å) and a substrate-bound outward-facing open state at a nominal resolution of 3.5 Å, with an estimated local resolution of up to 3.0 Å within the transporter unit (Fig. 5E and figs. S13 and S15). This allowed us to compare the binding sites of outward-facing open *HsPepT1* in its apo and substrate-bound forms (Fig. 5F). The structural overlay reveals that the negatively charged N-terminal pocket of the binding site is already positioned to coordinate the C terminus of the substrate in the central cavity. On the other hand, the positively charged C-terminal pocket undergoes larger rearrangements upon substrate binding, resulting in tightening of the central cavity around the substrate (fig. S16).



**Fig. 5. Structural basis for substrate recognition in human POTs.** (A) Concentration-dependent competition assay of the β-Ala-Lys peptide coupled to the fluorescent AMCA moiety (AK-AMCA) in *HsPepT2* with the dipeptide Ala-Phe. The average uptake value for each condition was calculated from three independent measurements. The error bars correspond to the SD from these independent measurements. (B) Thermal stabilization of detergent-solubilized *HsPepT2* upon substrate binding measured by nano-differential scanning fluorimetry (DSF) at increasing concentrations of Ala-Phe (inset shows the increase in melting temperature with increasing peptide concentration). (C) Close-up view of the *HsPepT2* peptide binding site. Electrostatic interactions between the peptide (shown in orange) and the transporter (shown in green) are displayed as gray dashes. The interaction between Y94 and the C-ter pocket is shown as green dashes. The different pockets are indicated. (D) Thermal stabilization of detergent solubilized *HsPepT1* upon substrate binding measured by nano-DSF at increasing concentrations of Ala-Phe. (E) Three-dimensional reconstruction of outward-facing open *HsPepT1* bound to Ala-Phe. (F) Overlay of the binding sites of outward-facing open apo (shown in gray) and Ala-Phe-bound *HsPepT1* (shown in blue colors). Electrostatic interactions between the peptide (shown in orange) and the transporter are displayed as gray dashes. The interaction between Y64 and the C-ter pocket is shown as blue dashes.

Notably, the N-terminal pocket consists of residues located in the C-bundle of the transporter, while the more flexible C-terminal pocket comprises residues from the N-bundle, in agreement with the dynamic observations made between outward- and inward-facing *HsPepT1* and *HsPepT2*.

These additional structures, which represent intermediate states between the outward- and inward-facing states described above, allow us to finally get a more detailed, step-by-step understanding of the molecular events required for transport and mediated by rearrangements of the N-bundle (Fig. 6 and movie S1). Upon peptide



agreement with the proposed “clamp and switch model” for MFS transporters (31, 57). A structural overlay of the N- and C-bundles of *HsPepT1* and *HsPepT2* reveals a stronger deviation for the N-bundle, indicating increased flexibility and dynamics in this part of the molecule, while the C-bundle remains rather rigid. This is in contrast to bacterial POTs (39, 41, 42) and sugar transporters (31, 57), where the N-terminal bundle has been identified as the most stable and rigid part of the molecule. This might be caused by the required stabilizing function of the C-bundle for the additional extracellular domain located between TM9 and TM10 in *HsPepT1* and *HsPepT2*. This is supported by the tight interaction network within the C-terminal bundle formed by the glutamine girdle (Fig. 4).

Our work created the framework for understanding the molecular transport mechanism of human POTs, which, in turn, will accelerate structure-based drug design approaches aiming to increase the bioavailability of different compounds in the human body via these transport systems. At the same time, despite available eukaryotic POT structures, the prediction of drug coordination remains challenging and additional transporter structures bound to a diverse set of drug or inhibitor molecules are required to obtain a more detailed understanding of drug recognition of this promiscuous transporter family.

## MATERIALS AND METHODS

### Expression and purification of *HsPepT2*

The N528Q-N587Q *HsPepT2* gene was cloned into a pXLG vector containing an expression cassette composed of an N-terminal Twin-Streptavidin tag followed by the human rhinovirus 3C (HRV-3C) protease recognition sequence. The double mutation in *HsPepT2* was introduced to decrease sample heterogeneity caused by glycosylation and to increase expression levels. HEK293F cells were collected 48 hours after transient transfection as previously described (60) and stored at  $-80^{\circ}\text{C}$  until further use. Frozen cell pellets were resuspended in 300 mM NaCl, 20 mM NaPi (pH 7.5), 0.5 mM tris(2-carboxyethyl)phosphine (TCEP), and 5% glycerol, supplemented with cOmplete EDTA-free protease inhibitors, and were disrupted using an Avestin Emulsiflex homogenizer. The lysate was centrifuged for 10 min at 10,000g, and the supernatant was centrifuged for 90 min at 95,000g (Optima XE-90, Beckman Coulter). The pellet containing the membrane fraction was solubilized in 1% *N*-dodecyl- $\beta$ -D-maltopyranoside (DDM; Anatrace) and 0.1% cholesteryl hemisuccinate (CHS; Tris Salt, Anatrace) for 1 hour at  $4^{\circ}\text{C}$ . The sample was centrifuged for 50 min at 70,000g, and the supernatant was applied to Strep-TactinXT beads (IBA). After 20 min of incubation on a rotating wheel, the suspension was transferred to a gravity column. Following two wash steps with 300 mM NaCl, 20 mM Hepes (pH 7.5), 0.03% DDM, and 0.003% CHS, *HsPepT2* was eluted with 0.03% DDM, 0.003% CHS, 150 mM NaCl, 20 mM Hepes (pH 7.5), and 10 mM desthiobiotin (Sigma-Aldrich). 3C cleavage was performed in 30 min, and the protease was separated from *HsPepT2* by gel filtration using Superose 6 Increase 10/300 (Sigma-Aldrich). The top fraction was concentrated to 10 mg/ml using a 100-kDa cutoff concentrator (Corning Spin-X UF concentrators) and stored at  $-80^{\circ}\text{C}$  until further use.

### Expression and purification of *HsPepT1*

The wild-type *HsPepT1* gene was cloned into a pXLG vector containing an expression cassette composed of an N-terminal

Twin-Streptavidin tag followed by the HRV-3C protease recognition sequence. HEK293F cells were collected 48 hours after transient transfection and resuspended in 300 mM NaCl, 20 mM NaPi (pH 7.5), 0.5 mM TCEP, and 5% glycerol, supplemented with cOmplete EDTA-free protease inhibitors. Whole cells were solubilized overnight in 1% lauryl maltose neopentyl glycol (LMNG; Anatrace) and 0.2% CHS (Tris Salt Anatrace). The sample was then centrifuged for 60 min at 70,000g, and the supernatant was applied to Strep-TactinXT beads (IBA). After 30-min incubation on a rotating wheel, the suspension was transferred to a gravity column. Following two wash steps with 300 mM NaCl and 20 mM Hepes (pH 7.5) supplied with 0.03% DDM, 0.003% LMNG, and 0.006% CHS, *HsPepT1* was eluted with 0.03% DDM, 0.003% LMNG, 0.006% CHS, 150 mM NaCl, 20 mM Hepes (pH 7.5), and 10 mM desthiobiotin (Sigma-Aldrich).

The sample was concentrated to 100  $\mu\text{l}$  using a 100-kDa cutoff concentrator (Corning Spin-X UF concentrators) and run directly on a Superdex Increase 200 5/150 gel filtration column for vitrification in 0.015% DDM, 0.0015% LMNG, 0.003% CHS, 150 mM NaCl, 50 mM Hepes (pH 7.5), and 0.5 mM TCEP. The top fraction reached a concentration of 2 mg/ml and was stored at  $-80^{\circ}\text{C}$  until further use.

### Whole-cell uptake assays

The wild-type *HsPepT2* gene was cloned into a pXLG vector (61) containing an expression cassette composed of an N-terminal hexa-histidine tag followed by enhanced green fluorescent protein and a tobacco etch virus protease cleavage site. HEK293F cells grown in suspension in FreeStyle medium were transfected with wild-type *HsPepT2* using a mass ratio of 2:1 polyethyleneimine:DNA. *HsPepT2* was expressed for 48 hours at  $37^{\circ}\text{C}$ , 8%  $\text{CO}_2$  at 220 rpm. For competition assays,  $4 \times 10^6$  cells/ml resuspended in phosphate-buffered saline buffer at pH 6.0 supplemented with 5 mM glucose were incubated in 96-well plates, with 50  $\mu\text{M}$   $\beta$ -Ala-Lys-AMCA in the absence or presence of dipeptides, tripeptides, or drugs for 10 min at  $37^{\circ}\text{C}$ . The reaction was stopped by adding 200  $\mu\text{l}$  of ice-cold buffer, and the cells were then washed three times with the same buffer. Last, the cells were resuspended in 200  $\mu\text{l}$  of buffer, and the fluorescence was measured in an M1000 microplate reader (TECAN) with excitation at 350 nm and emission at 450 nm. All experiments were performed in triplicates. The results were normalized by the fluorescence value of the control (cells overexpressing *HsPepT2* incubated with AK-AMCA in the absence of inhibitor) and plotted as AK-AMCA uptake rate percentage. For concentration-dependent uptake experiments,  $\text{IC}_{50}$  values were processed in GraphPad Prism 9.0 (GraphPad Software) using sigmoidal four-parameter curve fitting.

### Thermal stability measurements

The differential scanning fluorimetry method was used to follow the thermal unfolding event of *HsPepT2* and *HsPepT1* with a Prometheus NT.48 device (NanoTemper Technologies, Munich, Germany). Purified *HsPepT2* was diluted to 0.3 mg/ml and supplemented with decreasing amounts of Ala-Phe dipeptide in a dilution series of 13 points, starting at 10 mM down to 2.4  $\mu\text{M}$ . Purified *HsPepT1* was diluted to 0.3 mg/ml and supplemented with 0, 1, or 10 mM Ala-Phe. The fluorescence at 330 and 350 nm was recorded over a temperature gradient scan from  $15^{\circ}$  to  $95^{\circ}\text{C}$  and processed in GraphPad Prism 9.0 (GraphPad Software).

### Cryo-EM sample preparation and data collection on apo HsPepT1 in the outward-facing open state

Four microliters of purified HsPepT1 at 2 mg/ml was applied to a glow-discharged gold holey carbon 2/1 300-mesh grid (Quantifoil). The grid was blotted for 4 s at 0 force before being plunge-vitrified in liquid propane using Mark IV Vitrobot (Thermo Fisher Scientific). The blotting chamber was maintained at 4°C and 100% humidity during freezing. Movies were collected using a Titan Krios (Thermo Fisher Scientific) equipped with a K3 camera and BioQuantum energy filter (Gatan) set to 20 eV. A total of 22,537 movies were collected at a nominal magnification of  $\times 105,000$  and a physical pixel size of 0.85 Å, with a 70- $\mu\text{m}$  C2 aperture and 100- $\mu\text{m}$  objective aperture at a dose rate of 16  $e^-/\text{pixel}$  per second. A total dose of 66  $e^-/\text{Å}^2$  was collected with 3-s exposure as movies of 50 frames. Data were collected using EPU (Thermo Fisher Scientific).

### Cryo-EM sample preparation and data collection on HsPepT2 bound to Ala-Phe in the inward-facing partially occluded state

One hour before vitrification, purified N528Q-N587Q HsPepT2 was thawed on ice and run on a Superdex Increase 200 5/150 column in 0.015% DDM, 0.0015% CHS, 100 mM NaCl, 10 mM Hepes (pH 7.5), and 0.5 mM TCEP. The top fraction reached a concentration of 1 mg/ml, and 3.6  $\mu\text{l}$  supplemented with 5 mM of the dipeptide alanine-phenylalanine (Bachem) was applied to glow-discharged gold holey carbon 2/1 300-mesh grids (Quantifoil). Grids were blotted for 4 s at 0 force and 2-s wait time before being plunge-vitrified in liquid propane using Mark IV Vitrobot (Thermo Fisher Scientific). The blotting chamber was maintained at 4°C and 100% humidity during freezing. Movies were collected using Titan Krios (Thermo Fisher Scientific) outfitted with a K3 camera and BioQuantum energy filter (Gatan) set to 10 eV. A total of 34,712 movies were collected at a nominal magnification of  $\times 105,000$  and a physical pixel size of 0.85 Å, with a 70- $\mu\text{m}$  C2 aperture and 100- $\mu\text{m}$  objective aperture at a dose rate of 19.5  $e^-/\text{pixel}$  per second. A total dose of 81  $e^-/\text{Å}^2$  was collected with 3-s exposure as movies of 45 frames. Data were collected using EPU (Thermo Fisher Scientific).

### Cryo-EM sample preparation and data collection on HsPepT1 bound to Ala-Phe in the outward-facing open state and in the outward-facing occluded state

Four microliters of purified HsPepT1 at 2 mg/ml supplemented with 20 mM Ala-Phe was applied to a glow-discharged gold holey carbon 2/1 300-mesh grid (Quantifoil). The grid was blotted for 4.0 s at 0 force before being plunge-vitrified in liquid propane using Mark IV Vitrobot (Thermo Fisher Scientific). The blotting chamber was maintained at 4°C and 100% humidity during freezing. Movies were collected using Titan Krios (Thermo Fisher Scientific) equipped with a K3 camera and BioQuantum energy filter (Gatan) set to 15 eV. A total of 37,822 movies were collected in two separate sessions (16,522 in the first session and 21,300 in the second session) using EPU (Thermo Fisher Scientific) at a nominal magnification of  $\times 130,000$  and a physical pixel size of 0.67 Å, with a 70- $\mu\text{m}$  C2 aperture and 100- $\mu\text{m}$  objective aperture at a dose rate of 15  $e^-/\text{pixel}$  per second. A total dose of 55  $e^-/\text{Å}^2$  was collected with 1.7-s exposure as movies of 40 frames.

### Cryo-EM image processing of apo HsPepT1 in the outward-facing open state

Movies were motion-corrected using Relion-3.1 (62) own implementation of MotionCor2 (63). Contrast transfer function parameters

were calculated using CTFFIND4 (64). A total of 2,091,726 coordinates were extracted from 22,537 micrographs using CrYOLO (65), with a 200-pixel box and binning to 50 pixels, and were subjected to multiple rounds of 2D classification in Relion-3.1. A total of 1,459,348 particles were selected and reextracted with a 200-pixel box size without binning. A 3D ab initio reconstruction was generated in CryoSPARCv2 with all particles and low pass-filtered at 30 Å for 3D classification in Relion-3.1. After multiple rounds of 3D classification using  $T = 10$  and  $K = 4$ , the best classes were selected for additional 3D classification without image alignment in Relion-3.1, focusing on the protein and excluding the micelle. The selection of 593,757 particles from 3D classes with strong signal inside the micelle and in the extracellular domain led to a reconstruction of 4.6 Å in Relion-3.1. CTF refine (per particle defocus and beam tilt) and Bayesian polishing (using optimized trained parameters on a subset of 20,000 particles) were performed in Relion-3.1. The shiny particles were then imported in CryoSPARCv3 for nonuniform refinement (66) that led to a 3.9-Å reconstruction estimated in cryoSPARCv3 using the Fourier shell correlation (FSC) = 0.143 cutoff. Local resolution estimations were calculated in CryoSPARCv3 using 0.5 FSC cutoff. Postprocessing in DeepEMhancer (67) using the two half maps as input and the default tightTarget model resulted in a more interpretable map, which was used only for illustration purposes in Fig. 1B, while the model was build and refined in a map postprocessed using default parameters in the Phenix Autosharpen utility (68).

### Cryo-EM image processing of HsPepT2 bound to Ala-Phe in the inward-facing partially occluded state

Movies were motion-corrected using Relion-3.1 (62) own implementation of MotionCor2 (63). Contrast transfer function parameters were calculated using CTFFIND4 (64). A total of 4,388,314 coordinates were extracted from 34,712 micrographs using CrYOLO (65), with a 200-pixel box and binning to 50 pixels, and were subjected to multiple rounds of 2D classification in Relion-3.1. A total of 2,944,737 particles were selected and reextracted with a 200-pixel box size without binning. A 3D ab initio reconstruction was generated in CryoSPARCv2 with a subset of particles and low pass-filtered at 30 Å for 3D refinement in Relion-3.1 on the 2,944,737 particles, yielding a 7.5-Å reconstruction. After multiple rounds of 3D classification without image alignment in Relion-3.1, using  $T = 4, 8, 10, 20, 30,$  and 40 focusing on the protein and excluding the micelle, the selection of 454,149 particles from 3D classes with strong signal inside the micelle and in the extracellular domain led to a reconstruction of 4.3 Å in Relion-3.1 using SIDESPLITTER (69) and a soft mask covering the micelle and the extracellular domain. Bayesian polishing was performed in Relion-3.1 using optimized trained parameters on a subset of 20,000 particles. The shiny particles were then imported in CryoSPARCv3 for CTF Refinement per particle (defocus and beamtilt). Nonuniform refinement (66) led to a 3.8-Å reconstruction estimated in cryoSPARCv3 using the FSC = 0.143 cutoff. Local resolution estimations were calculated in CryoSPARCv3 using 0.5 FSC cutoff. Postprocessing in DeepEMhancer (67) using the two half maps as input and the default tightTarget model resulted in a more interpretable map, which was used only for illustration purposes in Fig. 1C, while the model was build and refined in a map postprocessed using default parameters in the Phenix Autosharpen utility (68).

## Cryo-EM image processing of HsPepT1 bound to Ala-Phe in the outward-facing open and occluded states

Movies were motion-corrected using Relion-3.1 (62) own implementation of MotionCor2 (63). Contrast transfer function parameters were calculated using CTFFIND4 (64). A total of 6,046,602 coordinates were extracted from 37,822 micrographs using CrYOLO (65), with a 200-pixel box and binning to 50 pixels, and were subjected to multiple rounds of 2D classification in Relion-3.1. A total of 4,247,238 particles were selected and reextracted with a 200-pixel box size without binning. The 3D volume of apo HsPepT1 in its outward-facing open conformation was low pass-filtered at 40 Å for 3D classification in Relion-3.1, resulting in a reconstruction at 4.1 Å with 486,562 particles for the first dataset, and 4.5 Å with 599,754 particles for the second, after performing Bayesian polishing using individually optimized trained parameters on a subset of 30,000 particles for each dataset. In both datasets, further 3D classifications allowed to separate two distinct conformational states, including the substrate-bound outward-facing open state and the substrate-bound outward-facing occluded state differing by the opening or closure of TM2. Focused 3D classification without alignment on the transmembrane domain in Relion-3.1 allowed to improve the resolution of the substrate-bound outward-facing open state from 3.7 to 3.5 Å after nonuniform refinement (66) in cryoSPARCv3. The substrate-bound outward-facing occluded state reached a modest resolution of 4.1 Å after nonuniform refinement in cryoSPARCv3 but led to a clear resolution of all TMs, including the closed TM2. For the 3.5-Å resolution substrate-bound outward-facing open reconstruction, postprocessing was done in Phenix using default parameters in the Autosharpen utility (68). This map was subsequently used for model building and refinement. For illustration purposes in Fig. 5E only, the half maps were also subjected to postprocessing in DeepEMhancer (67) using the two half maps as input and the default tightTarget model. For the 4.1-Å resolution substrate-bound outward-facing occluded reconstruction, postprocessing was done in cryoSPARCv3.

## Model building and refinement

The transmembrane domain of HsPepT2 was built manually in Coot (70), guided by secondary structure predictions from PSIPRED (71). The resolution in the extracellular domain of HsPepT2 did not allow de novo model building. Instead, the structure of HsPepT2-ECD predicted by AlphaFold (50) was docked in the cryo-EM density and linked to the transmembrane domain. The model was iteratively adjusted and refined using Isolde (72) and Phenix real-space refine (73). An inward open partially occluded model of HsPepT1 was generated in SWISSMODEL (74), using HsPepT2 as a reference template. The model was then subjected to multiple rounds of refinement in NAMDINATOR (75), Coot, and Phenix real-space refine (73) to fit in the apo outward-facing open reconstruction. Once the model fitted the density, the extracellular domain was replaced by the AlphaFold HsPepT1-ECD prediction, which improved the quality of the model significantly. This model was then subjected to adjustments in Isolde and a final refinement using Phenix real-space refine. The substrate-bound outward-facing open and occluded models were generated using the final apo outward-facing open model, and manual adjustments in Coot and Isolde before real-space refinement in Phenix. Validation of the models was performed using MolProbity in Phenix.

## SUPPLEMENTARY MATERIALS

Supplementary material for this article is available at <https://science.org/doi/10.1126/sciadv.abk3259>

[View/request a protocol for this paper from Bio-protocol.](#)

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