Tripartite ATP-independent Periplasmic (TRAP) Transporters Use an Arginine-mediated Selectivity Filter for High Affinity Substrate Binding*

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Background: Haemophilus influenzae requires a substrate-binding protein (SBP)-dependent TRAP transporter to acquire sialic acid.
Results: A conserved arginine residue in the SBP is essential for the high affinity and carboxylate specificity of the TRAP transporter.
Conclusion: The arginine/carboxylate interaction in TRAP SBPs restricts substrate range to carboxylate-containing substrates.
Significance: The study reveals the mechanism by which a key bimolecular interaction underpins bacterial virulence.

Tripartite ATP-independent periplasmic (TRAP) transporters are secondary transporters that have evolved an obligate dependence on a substrate-binding protein (SBP) to confer unidirectional transport. Different members of the DctP family of TRAP SBPs have binding sites that recognize a diverse range of organic acid ligands but appear to only share a common electrostatic interaction between a conserved arginine and a carboxylate group in the ligand. We investigated the significance of this interaction using the sialic acid-specific SBP, SiaP, from the Haemophilus influenzae virulence-related SiaPQM TRAP transporter. Using in vitro, in vivo, and structural methods applied to SiaP, we demonstrate that the coordination of the acidic ligand moiety of sialic acid by the conserved arginine (Arg-147) is essential for the function of the transporter as a high affinity scavenging system. However, at high substrate concentrations, the transporter can function in the absence of Arg-147 suggesting that this bi-molecular interaction is not involved in further stages of the transport cycle. As well as being required for high affinity binding, we also demonstrate that the Arg-147 is a strong selectivity filter for carboxylate-containing substrates in TRAP transporters by engineering the SBP to recognize a non-carboxylate-containing substrate, sialylamide, through water-mediated interactions. Together, these data provide biochemical and structural support that TRAP transporters function predominantly as high affinity transporters for carboxylate-containing substrates.

Secondary transporters use preformed electrochemical gradients to energize the concentrative movement of chemicals across biological membranes and are ubiquitous in living cells. Bacteria use secondary transporters primarily for uptake of nutrients from their environment, and often bacterial genomes can encode over 100 different secondary transporters (1), which will likely transport a very wide variety of substrates ranging from sugars, amino acids, nucleotides, fatty acids, inorganic ions, and organic acids (1, 2). Many bacteria and archaea, especially those that live in the sea and other marine environments (7), and also a range of human pathogens, use a particular family of secondary transporters called tripartite ATP-independent periplasmic (TRAP)6 transporters that also have a substrate-binding protein (SBP) component (3–5). The SBP is either free in the periplasm of Gram-negative bacteria or anchored to the cell membrane in Gram-positive bacteria and is a transporter component traditionally found in prokaryotic ATP-binding cassette uptake systems (6, 7). Through the characterization of the SiaPQM sialic acid-specific TRAP transporter from Haemophilus influenzae, which is essential for host colonization, some of the properties conferred by using the SBP have been elucidated at the biochemical level (8–11). TRAP transporters have three subunits, the SBP and two membrane proteins of unequal size. The M subunit is a 12-transmembrane helix (TMH)-containing protein that is a member of the ion transporter superfamily and likely forms the translocation channel, whereas the Q subunit is a 4-TMH protein of essential but undefined function. In the H. influenzae SiaPQM TRAP trans-

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6The abbreviations used are: TRAP, tripartite ATP-dependent periplasmic; SBP, substrate-binding protein; r.m.s., root mean-squared; Neu5Ac, N-acetylneuraminic acid; TMH, transmembrane helix; IPTG, isopropyl 1-thio-β-D-galactopyranoside; LOS, lipooligosaccharide.
porter, the SBP is SiaP, and the two membrane proteins that constitute the TRAP transporter, SiaM and SiaQ, are naturally fused into a single predicted 17-TMH protein, SiaQM. The SBP initiates the transport process by binding the substrate. By being located on the extra-cytoplasmic side of the membrane, it imposes directionality on a secondary transporter by delivering the ligand to the SiaQM membrane domains for subsequent translocation into the cell (12).

The first structure of a TRAP SBP (5) revealed that the charge of the carboxylate group of the ligand, sialic acid, was neutralized by an arginine residue, Arg-147, in SiaP, which sequence analysis had revealed to be the most highly conserved residue, being present in 98% of TRAP SBPs of the common DctP family (5). On this basis, the Arg/carboxylate interaction was proposed to be a defining feature of the DctP-TRAP SBPs that might have additional functions in the overall transporter mechanism than simply this electrostatic interaction with the ligand (4), such as triggering of domain closure in the protein. Additional structures of DctP-type SBPs from TRAP transporters that recognize (hydroxy-)ectoine, pyroglutamate, and monocarboxylate keto acids as substrates all have a similarly functioning arginine residue (13–17). By comparing the substrate-binding sites in these different SBP structures, it is clear that each binding site is highly adapted to its particular carboxylate-containing substrate, with the notable exception of the conserved Arg/carboxylate interaction that is observed between the equivalent residues in Arg-147 of SiaP and the carboxylate group of the different substrates (4). A remarkable recent study has added another 29 unique DctP-TRAP SBP structures to the Protein Data Bank, each with an equivalent arginine (18). These data reinforce the key role of this residue in defining the biological function and substrate range of TRAP transporters, but the structural and biochemical consequences of its disruption have not been investigated to date.

To investigate how SiaP recognizes its sialic acid substrate, Johnston et al. (8) examined the biological phenotypes of a number of site-directed mutants of SiaP. Residue Arg-147 was mutated to lysine and to alanine. Using a monoclonal antibody that is unable to recognize sialylated lipooligosaccharide (LOS), they demonstrated a complete lack of complementation of a siaP strain of non-typable H. influenzae by genes that contained an Arg-147 mutation (8). In contrast, other mutations of the binding site resulted in complete or partial complementation, suggesting that they were not important for transport and hence the sialylation phenotype. The basis for the lack of LOS sialylation for Arg-147 mutants was not determined but is consistent with Arg-147 playing an important role in transporter function.

In this study we determined the role of the arginine/carboxylate interaction using SiaP. We demonstrate that it is essential for high affinity binding of substrate and transport under physiological conditions of low substrate concentrations but also that its loss can be tolerated at high ligand concentrations. By disrupting this key interaction in a rational way, we show that we can change the substrate specificity of the SBP from binding a carboxylate to binding an amide. Together these data demonstrate a key role for the Arg-147/carboxylate interaction in TRAP transporter function.

**Experimental Procedures**

**Bacterial Strains and Culture Conditions**—For cloning and transformation, Escherichia coli DH5α, BW25113, and MC1061 were grown in LB broth (10 g/liter tryptone, 5 g/liter powdered yeast, 10 g/liter NaCl). Protein production and growth on specific carbon sources used M9 minimal medium salts (19) supplemented with 0.4% d-glucose, Neu5Ac, or other carbon source at the indicated concentrations. Antibiotic selection, where appropriate, used 30 μg/ml chloramphenicol, 100 μg/ml ampicillin, and 50 μg/ml kanamycin.

**Mutagenesis of the siaP Gene**—The wild-type siaP gene (HI0146) was amplified by PCR from genomic DNA of H. influenzae Rd using the primers siaPfor 5′-cgctgctacaaaaagagacatatagatgaaattgacaaac-3′ and siaPhis6rev 5′-ccgctgaagtttatcttgcgttac-3′. The product was digested with Ndel and Xhol and cloned into pBlueScriptII KS, verified by DNA sequencing, and then subcloned using the same sites into pET21b creating pAH16. To mutate the siaP gene in pAH16 directly, we used partially overlapping oligonucleotide primers with a silent Stul restriction site in the reverse primer (20); the primers for R147A were 5′-aaatgtgctgccaaatgcagcaaaac-3′ (forward) and 5′-ggccacacaagtggctgcctcatctc-3′ (reverse); R147K primers were 5′-aaatgatgaaattgacaaac-3′ (forward) and 5′-ggccacacacatctgctcatctc-3′ (reverse); and R147E primers were 5′-aaatgacagaataggctgcctcatctgct-3′ (reverse), creating plasmids pAH35 (R147A), pAH36 (R147E), and pAH37 (R147K). For low copy number expression in *in vivo* growth and uptake experiments, we used pWKS30 (21) as the vector. Either wild-type or mutated *siaP* alleles from pAH16 and pAH35-37 were amplified and digested with Ndel and Xhol before being cloned in a three-way ligation with Xhol-siaQM-BamHI and Ndel-, BamHI-cut pWKS30, pAH15 (siaP-His6), pES15 (R147A), pES16 (R147K), and pES17 (R147E). To mutate the *siaP* gene in pAH16 directly, we used partially overlapping oligonucleotide primers with a silent Ncol restriction site in the reverse primer (19); F170Wfor 5′-atggcaatggttgctgataatctgc-3′ and F170Wrev 5′-tttctgagattcttcctgtaatc-3′ and F170Wrev 5′-tttctgagattcttcctgtaatc-3′, creating pAH45. The same primers were used to introduce this point mutation in the pAH35-37 series creating pAH66, pAH64, and pAH65, respectively. The clones were verified by DNA sequencing.

**Expression and Purification of SiaP Proteins**—A flask containing 50 ml of M9 minimal medium containing 0.4% d-glucose was inoculated with *E. coli* BL21 (DE3) pLysS pAH16 or other pET21 derivatives and grown at 37 °C overnight with shaking at 180 rpm. This was inoculated into 625 ml of M9 minimal medium containing 0.4% d-glucose to an *A*0.600 0.1, grown at 25 °C to an *A*0.600 0.2–0.3, expression induced by the addition of 1 mM IPTG, and incubated overnight at 25 °C. Induced cells were spun at 4000 × g for 20 min at 4 °C, resuspended in 25 ml of 5 mM EDTA, 50 mM Tris, 0.5 mM succrose, pH 8.0, and incubated with 12 mg of lysozyme (chicken egg white; Sigma) at 30 °C for 2 h to prepare the periplasmic fraction. The sample was then centrifuged at 17,000 × g for 10 min at 4 °C, and the supernatant was dialyzed against 20 mM Tris/HCl, 300 mM NaCl, pH 7.5 (TBS). This was clarified by centrifugation at
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17,000 $\times$ g for 10 min at 4 °C, and imidazole was added to 12 mM. A 1-ml HisTrap HP column (GE Healthcare) was washed with TBS, 12 mM imidazole using a P-1 peristaltic pump (Amersham Biosciences). The dialyzed and clarified periplasmic fraction was loaded onto the column at a 2–5 ml min$^{-1}$, washed with 20 column volumes of TBS, 20 mM imidazole, and the protein eluted with TBS 400 mM imidazole. Protein-containing fractions were visualized using SDS-PAGE, pooled, and concentrated using Vivaspin 2 ultrafiltration spin columns with a 5-kDa molecular mass cutoff (Sartorius).

**Protein Fluorescence Spectroscopy**—Protein fluorescence experiments were performed in a 3-ml quartz cuvette (Starna) using a FluoroMax2 (Instruments SA, Inc.) with an LTD6 water bath (Grant) controlled with the supplied software, DataMax-Std version 2.20. SiaP contains no tryptophans, and so the protein was excited for tyrosine fluorescence. 0.05 $\mu$M protein in 50 mM Tris/HCl, pH 8.0, was excited at 281 nm with slit widths of 5–10 nm to give a signal intensity of 2–3 $\times$ 10$^5$ units. Ligand was added at concentrations and times to produce spectra and time course titrations. For titrations, the cumulative fluorescence change was plotted using SigmaPlot (version 10.0), and the $K_D$ value was determined using a fit to a simple hyperbolic curve. For the F170W/R147K mutant titrations, protein was using 10-fold control was run in parallel to remove dilution effects from the calculation of $K_D$ values.

**Circular Dichroism**—CD spectra were determined using a J-810 spectropolarimeter (Jasco) controlled by the supplied software Spectra Manager version 1.53.00 and maintained at constant temperature by the Peltier unit PFD-425S. The spectrum of 0.1 mg/ml protein in 10 mM potassium phosphate buffer, pH 8.0, was determined in a 1-mm pathlength quartz cuvette (Starna) between 240 and 180 nm at 100 nm/min with a 1-nm pitch.

**Isothermal Titration Calorimetry**—Both the protein and ligand preparations were degassed at 2 °C below the experimental temperature immediately prior to analysis using a VP-ITC microcalorimeter (MicroCal), controlled by VPViewer2000 version 1.4.24 (MicroCal LLC). The titration pattern was an initial injection of 3 $\mu$l of the ligand, followed by 6-$\mu$l injections of 14 s and separated by 180 s. The titrations were then analyzed using Origin 7SR2 version 7.0383(B383) (OriginLab Corp.) by fitting to one set of binding sites.

**Whole Cell [$^{14}$C]Neu5Ac Uptake Assay**—These were performed in a similar manner to Severi et al. (22). Briefly, overnight cultures in minimal medium with the carbon source of interest were diluted to an $A_{650}$ of 0.1 in the same medium and grown to an $A_{650}$ of 0.5. These cells were harvested, washed four times, and resuspended in M9 salts to an $A_{650}$ of 3.0. For Neu5Ac uptake assays, cells were diluted 10-fold in M9 at 37 °C and allowed to acclimatize for 2 min with stirring, before initiating the assays by adding varying amounts of [$^{14}$C]Neu5Ac (Sigma) appropriately diluted with unlabeled Neu5Ac. The uptake assay and total protein quantification were then performed as described in Severi et al. (11), except that 200 $\mu$l of cell suspensions were immobilized instead of 400 $\mu$l. $V_{\text{max}}$ and $K_m$ values were calculated by fitting to a hyperbolic Michaelis-Menten equation using SigmaPlot.

**SiaQM-Proteoliposome [$^{14}$C]Neu5Ac Uptake Assay**—In vitro [$^{14}$C]Neu5Ac uptake by reconstituted SiaPQM was measured using the method of Mulligan et al. (12). N-terminally tagged SiaQM was purified from the E. coli MC1061 pBADQNM membrane fraction using nickel-nitrilotriacetic acid resin (Qiagen) and reconstituted into proteoliposomes with E. coli lipids by rapid dialysis. Proteoliposomes containing 200 $\mu$g of SiaQM were resuspended in Inside buffer (100 mM potassium acetate, 20 mM potassium phosphate, 2 mM MgSO$_4$, pH 7.0) and extruded 11 times through a 400-nm polycarbonate filter (Avestin Inc.). The extruded proteoliposomes were collected by ultracentrifugation and resuspended in 50 $\mu$l of Inside buffer. For each uptake assay, 5 $\mu$m of the binding protein of interest and 5 $\mu$m [$^{14}$C]Neu5Ac were added to 300 $\mu$l of the reaction (Outside) buffer (100 mM sodium acetate, 2 mM MgSO$_4$, 20 mM sodium PIPES, pH 7.0, 1 $\mu$m valinomycin) and incubated at the reaction temperature of 30 °C for 1 min. 6 $\mu$l of 1.15 $\mu$m SiaQM-proteoliposomes were added to start the reaction, and 50-$\mu$l samples were taken as indicated. Each of these samples was mixed with 50 $\mu$l of reaction buffer containing 1 mM unlabeled sialic acid for 10 s, added to a 0.22-$\mu$m nitrocellulose filter (Milipore), and washed with 2 ml of 50 mM potassium phosphate buffer, pH 7.0. The radioactivity associated with the filters was determined using liquid scintillation counting.

**Bacterial Growth Experiments**—E. coli BW25113 ΔnanT containing pAH15 or pES15-17 was inoculated to an $A_{650}$ 0.01 in 700 $\mu$l M9 minimal medium with Neu5Ac and 1 mM IPTG in a 24-well glass-bottom plate closed by an air-permeable sterile lid. Growth at 35 °C with shaking at 250 rpm was monitored using a prototype incubated plate shaker (EnzyScreen). Shaking was halted every 30 min for about 1 min so that a flat-bed scanner could capture an image of the base of the plate. The increasing whiteness of the growing cultures was converted to a G value by the associated software (ImageAnalysisGIU version 1.0.0.0), which was correlated to an $A_{650}$ value ($A_{650}^G$) using a stand curve. The density of all growth experiments was within the linear range of this.

For growth at low (1 mM) concentrations of Neu5Ac, we used a Tecan Infinite M200 Pro microplate reader. Bacteria were inoculated from a single colony into 4 ml of M9 glucose (0.4% w/v) medium and grown overnight. For strains harboring plasmids, ampicillin (100 $\mu$g/ml) and IPTG (1 mM) were added. The $A_{650}$ of each culture was measured and adjusted to 1 by addition of 1× M9 salts. 150 $\mu$l of M9 minimal medium with the appropriate concentration of Neu5Ac was aliquoted into the 96-well plate (Corning Costar 3595), and 1.5 $\mu$l of bacterial cell suspension was added and mixed by pipetting. For strains harboring plasmids, ampicillin (100 $\mu$g/ml) and IPTG (1 mM) were added. The outer wells of the 96-well plate were filled with 150 $\mu$l of sterile distilled H$_2$O to prevent evaporation from the internal wells. The plate reader was set to monitor $A_{650}$ every 30 min for 48 h and to maintain a constant temperature of 37 °C. Data were exported to Magellan data analysis software for analysis. An average of two media-only control wells was taken from every 30-min time point. The data were normalized by subtracting this value from the $A_{650}$ reading for all cultures at

Arg-147 Is Essential for High Affinity Ligand Binding to SiaP in Vitro—A reanalysis of the Pfam database family for DctP-TRAP SBPs (SBP-bac-7) reveals that 5943 of 6142 full-length sequences (96.8%) have an arginine residue in an equivalent position to Arg-147 in SiaP. To investigate the involvement of SiaP residue Arg-147 in transporter function, we mutated it to either alanine (R147A) or lysine (R147K). Binding of N-acetylneuraminic acid (Neu5Ac) to C-terminally hexahistidine-tagged wild-type SiaP-His6 yielded a $K_D$ of 0.14 $\pm$ 0.04 $\mu$M, in close agreement with published data for native or N-terminally His-tagged versions (Fig. 1A) (8, 11). In contrast to this high nanomolar affinity, the R147A and R147K mutants analyzed in version 0.99 and CorelDRAW X3, respectively. Procheck, and the ADIT Protein Data Bank deposition server. The data were checked to fulfill the following criteria: the "Density Fit Graph" and "Check Waters" validation tools in Coot, added automatically using Coot. Finally, water molecules were added in the final iterations of model building and refinement, and Babinet scaling was applied. Structures were refined anisotropically with the exception of the R147K structure (2wx1), which was refined isotropically at a resolution of 2.2 Å. Composite omit maps were calculated with Comit (23) for the placement of low occupancy water molecules that were added automatically using Coot. Finally, water molecules were checked to fulfill the following criteria: the "Density Fit Graph" and "Check Waters" validation tools in Coot, B-factor variance, and H-bond distances. The identity of the compounds within the WT and R147E structure was checked using unrestrained refinement on the basis of their atomic factors and bond lengths. Finally, the data were checked using Coot, Sfcheck, Procheck, and the ADIT Protein Data Bank deposition server. The superpositions and figures were generated with PyMOL version 0.99 and CorelDRAW X3, respectively.

Results

Arg-147 Is Essential for High Affinity Ligand Binding to SiaP

According to Mulligan et al. (12), we could readily measure uptake with the wild-type protein, we were unable to measure uptake with either the R147A or R147K mutants (Fig. 2B). Taken together, these data suggest that physiological high affinity transport by SiaPQM is abolished by the alteration of the Arg-147 residue of SiaP.

Arg-147/Carboxylate Interaction Is Not Essential for SBP Closure—The high level of conservation of the Arg-147 residue, or its equivalent in other DctP-TRAP SBPs, is unusual for a binding protein family, where sequence diversity is normally much greater. Given the almost total loss of detectable function for SiaPQM containing this single point mutant within an otherwise unaltered binding site, we wondered whether the arginine/carboxylate interaction might have additional fundamental roles in the SBP function such as domain closure or productive interactions with the SiaQM proteins. To understand the structural consequences of removing the Arg-147 from SiaP, we crystallized the R147A and R147K mutants in the presence of 10–15 mM Neu5Ac (Table 1). Surprisingly, given the very weak binding observed for these proteins in vitro, both structures contain Neu5Ac bound in the substrate binding pocket, and the proteins are clearly in the closed ligand-bound conformation as opposed to the expected open ligand-free conformation that has been observed for SiaP previously (Fig. 3A) (5, 8). Overall, both mutant structures are structurally indistinguishable from the wild-type structure in complex with Neu5Ac (3B50) (8) with a root mean square deviation (r.m.s.d.) of 0.30 and 0.17 Å for the R147K and R147A mutants, respectively. The ability of the mutant proteins to bind and close around the Neu5Ac in these conditions suggests that the interaction of Arg-147 with the carboxylate is neither essential for protein closure nor Neu5Ac accommodation. Neu5Ac is rec-
Ordered Water Molecules Mediate Ligand Contacts to the SiaP Arg-147 Mutants—We examined the structure in more detail to assess how the loss of the Arg-147 residue is still compatible with binding of Neu5Ac. With the relatively conservative substitution of arginine to an also charged lysine (Fig. 3B), the key guanidinium group of Arg-147 is partially substituted by the NH atom of Lys-147, which occupies an equivalent position (0.3 Å shift) to one of the NH2 groups of Arg-147 and provides one of the interactions with the ligand (Fig. 3B). The other contact is now mediated by a water molecule that provides the missing H-bond to the carboxylate of Neu5Ac (Fig. 3B). The R147A mutant protein also crystallized in a closed Neu5Ac-bound form, despite effectively reducing the highly conserved charged side chain to a methyl group (Fig. 3B). In contrast to the R147K mutant, the replacement of Arg-147 by alanine results in removal of all direct contacts between residue 147 and the Neu5Ac (Fig. 3C). Rather, two water molecules dissipate the charge of the Neu5Ac carboxylate, mediating contacts between ligand and protein. One of these water molecules (water 1) is present in both the R147K and R147A structures, although the other water (water 3) is unique to this structure.

Despite this significant change in the binding pocket, the structure of SiaP around the missing arginine is only very mildly perturbed. In both structures (2xwi and 2xwk) slightly elevated B-factors for the carboxylate group of Neu5Ac compared with the rest of the ligand indicate that this part of the Neu5Ac is less constrained in the absence of the strong interaction with the Arg-147 residue, compared with the wild-type protein. Water 3 also has a higher B-factor than water 1, which manifests as an anisotropic motion perpendicular to the carboxylate. This mobility translates to the neighboring water molecule (water 4) that moves by ~1 Å with respect to the wild type, concomitant with a 0.5-Å translation of the β-strand that bears residue 147. There are no additional waters present in the void left by the short side chain of Ala due to the aliphatic patch behind the arginine that drives it into the binding site (4). Hence, although Arg-147 provides important contacts that are essential for high affinity binding, the required coordination of the Neu5Ac carboxylate in the binding pocket can be fulfilled by water molecules allowing these mutants to both bind and close around the Neu5Ac under these conditions.

High Ligand Concentrations Restore in Vivo Transporter Function for SiaP Arg-147 Mutants—Given that the structural data suggested that the Arg-147 mutant can close and the struc-
ture of the ligand-bound forms for the wild-type and Arg-147 mutants are indistinguishable apart from the missing Arg/carboxylate interaction, we reasoned that transporter function might be restored under similar substrate concentrations to those used for crystallography. To test this hypothesis, we used a genetic system that requires the presence of a functional sialic acid transporter to allow growth of *E. coli* in liquid media with sialic acid as the sole carbon source. In an M9 minimal medium, *E. coli* will grow on 3.2 mM (1 mg/ml) Neu5Ac as the sole carbon source, which is dependent on the NanT sialic acid transporter, a classical secondary transporter of the major facilitator superfamily (22, 24). A ΔnanT strain cannot grow on Neu5Ac unless a functional sialic acid transporter is provided in trans, and siaPQM can function in this role (22). Using the ΔnanT strain complemented with either the wild-type siaPQM or equivalent genes containing the specific Arg-147 mutants, we were indeed able to detect growth for all of the strains (Fig. 2). Interestingly, we observed differential growth phenotypes for the Arg-147 mutants, with the more conservative changes in the R147K mutant exhibiting growth more similar to the strain expressing the wild-type SiaP (doubling times of 120 min), than the R147A mutation, which resulted in an ∼50% decrease in the growth rate, giving a doubling time of about 180 min.

We repeated these experiments using a microplate reader with lower (1 mM) Neu5Ac concentrations and saw a similar pattern of results, although the R147A mutant growth is severely attenuated (Fig. 4). These data suggest that in the presence of relatively high external concentrations of sialic acid and in the context of transporter function in whole cells, the Arg-147 residue is not essential and in fact links the differential phenotypes for substitutions of the Arg-147 to the nature of the side-chain modifications introduced and observed in the crystal structures.

**Weakening Protein and Ligand Interactions Simultaneously Abolish Transporter Function**—*E. coli* can grow on the related nonulosonic acid KDN using siaPQM expressed in trans (25), but this compound binds to SiaP with over 300-fold lower affinity than Neu5Ac (KD of 42 μM) (5). To test whether wild-type and mutant versions of SiaPQM could function with this weaker binding substrate, we repeated the above experiments but with KDN replacing Neu5Ac as the sole carbon source. Although we could detect growth for the wild-type on KDN, there was no detectable growth of the Arg-147 mutants (Fig. 2). Together these data suggest that for the physiological substrate of SiaPQM, the loss of the function of the Arg-147 can be tolerated in *vivo* at high substrate concentrations. However, for a ligand that already binds relatively weakly to the wild-type protein, the resulting drop in ligand binding affinity in the Arg-147 mutants is too great to support any physiological function-
ing of the transporter in the low micromolar concentrations of free Neu5Ac found in the body.

**Engineered Arg-147 Mutant Can Accommodate Non-carboxylate-containing Ligands**—The coordination of Neu5Ac in the R147A mutant suggests that waters can essentially replace the guanidinium group of the arginine and retain the ability of the protein to coordinate the carboxylate of the ligand. To investigate whether this water network adds plasticity, i.e. ligand promiscuity, we engineered an R147E mutant to maintain the hydrophobic part of the arginine side chain and yet have the capacity to coordinate water 1 and water 3 observed in the R147A structure. In addition, by using the charge-swap from Arg to Glu, we can also assess the ability of the water molecules to dissipate the apparent charge clash. In vivo data confirm the in silico predictions as growth of bacteria containing SiaP with the R147E mutation is more impeded than the R147K mutation but less than the R147A mutation (Fig. 2). These data suggest that the two waters are able to accommodate two carboxylate groups pointing directly toward each other, while retaining sufficient function to enable bacterial growth.

To investigate further the ability of the waters to relax the selectivity of SiaP for its ligands, the R147E mutant was crystallized in the presence of the non-cognate ligand sialylamide, which differs from Neu5Ac by a change of the carboxylate to an amide (Fig. 5A). This neutral molecule was previously reported to bind wild-type SiaP with around a thousand-fold lower affinity than Neu5Ac ($K_D$ of 240 μM (5)). The R147E-sialylamide co-crystal structure (Fig. 5A) again reveals a closed conformation with an r.m.s.d. of 0.2 Å from the wild-type SiaP in complex with Neu5Ac. The amide and the opposing carboxy-amine (Fig. 5 which differs from Neu5Ac by a change of the carboxylate to an amide (Fig. 5A) as the NH$_2$ group donates rather than accepts hydrogen bonds.

We attempted to crystallize the wild-type protein with this non-cognate sialylamide ligand and were able to collect data to atomic resolution of 1.05 Å where ab initio phasing was applied to remove any model bias. Strikingly, the ligand appeared to be coordinated exactly as Neu5Ac, including the positioning of all waters in the binding pocket, and careful refinement of the structure revealed that the bound species was in fact Neu5Ac, based on analysis of H-bonding patterns (interaction with waters), B-factors of the carboxylate, and also considering expected bond lengths for unrestrained refinement of oxygen and nitrogen atoms (26). This final structure (2xwv) is indistinguishable (r.m.s.d. 0.1 Å) from the reference SiaP-Neu5Ac complex of Johnston et al. (8) and has the highest resolution for a TRAP SBP to date (Fig. 5). Mass spectrometry data confirmed the presence of trace amounts of Neu5Ac in the sialylamide sample despite purification after synthesis using silica and ion-exchange chromatography and checked by one-dimensional $^1$H NMR. The finding that the wild-type protein was able to selectively bind this minor contaminating component, while R147E protein exposed to the same stock crystallized with the bound sialyl-
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![A](image1)

**FIGURE 3. Ligand binding induces closure of the mutants with waters substituting missing direct interactions.** A, structural superposition of mutants R147K (magenta) and R147A (green) onto WT (gray, shown with side chains) highlights that all structures exhibit a closed conformation in the presence of Neu5Ac (blue) and are structurally indistinguishable. B, close-up of the binding pocket of the R147K mutant solved to 2.2 Å resolution, with electron density (2mFo−DFo, omit maps rendered at 1σ) shown, reveals that two water molecules 1 and 2 (red spheres labeled with blue font) take up interactions with opposite ends of the ligand enabled by the alternative conformation of Asn-10 and the Lys mutation. C, close-up of the binding pocket of R147A (2mFo−DFo, omit maps rendered at 1.5σ). The Ala mutation opens space for another water (3) in an equivalent position to the missing guanidinium group, but without altering the surrounding backbone in comparison with the wild type (shown in gray lines).

amid, confirms the important role of the Arg-147 residue in the selectivity of SiaP for its cognate ligand.

**Discussion**

As we learn about the diversity of bacterial physiology through genome sequencing, it is clear that bacterial TRAP transporters form significant components of the encoded repertoire of solute transporter in the genomes of bacteria that live in nutrient-poor environments (7). The best examples of this are in marine environments that contain a wide range of dissolved organic compounds at low concentrations, requiring organisms that live in these environments to be able to scavenge effectively for many diverse nutrients (27). TRAP transporters appear to be enriched in these environments as they have high affinity, use Na⁺ as a coupling ion for transport, and have specificity for diverse organic acids that appear to be major carbon sources in these niches (28, 29). When TRAP transporters were first discovered, they were as part of C₁-C₆-dicarboxylate transporters, but subsequent work has identified a range of other ligands, united by containing a negatively charged group, mostly carboxylic acid, or in the case of ectoine a sulfonate group (3, 7, 13, 14). More recently, around 30 novel TRAP SBPs of the DctP family had their structures solved with co-purified ligands, revealing a whole plethora of new organic acid ligands, but also examples of zwitterionic amino acids and a glycerol 3-phosphate molecule bound. In all of these structures, the carboxylate (or rarely a sulfonate or phosphate) forms an electrostatic interaction with the conserved arginine in these proteins, equivalent to Arg-147 in SiaP (18). We have now investigated using SiaP, the structural and biochemical role of this residue in TRAP transporter function.

First, the binding affinity of SiaP is significantly reduced in the absence of Arg-147. All three Arg-147 mutants have major defects in ligand binding in vitro and in vivo using physiological (micromolar) concentrations of substrate. The Arg-147 mutations do not alter the overall structure of SiaP significantly (r.m.s.d. of 0.12, 0.14, and 0.23 Å between wild-type and R147A, R147E, and R147K, respectively), and the impact of these structural changes is seen through the loss of the ionic salt bridge between the protein and the ligand. Andrews et al. (30) reported the average intrinsic binding energy of a carboxylate group to be 8.2 kcal/mol (in the range 7.3–10.3 kcal/mol at 298 K), and hence an important component of driving high affinity in a ligand-binding site. Given that a change in the equilibrium constant for binding (K_d) by 1 order of magnitude corresponds to a change in Gibbs free energy (ΔG) of ~1.36 kcal/mol, the loss of the salt bridge could lead to a reduction in K_d of over 5 orders of magnitude. This corresponds to a decrease in binding affinity from 120 μM to ~10 mM, which is very similar to our experimental observations. It supports the hypothesis that Arg-147 is essential for physiological function of TRAP transporters that bind carboxylate-containing ligands. Further support for the “affinity” hypothesis comes from experiments with KDN, a 300-fold weaker binder than Neu5Ac. KDN binding is undetectable when other parts of the interaction with the binding site are changed in addition to the broken Arg/carboxylate interaction. Hence, our data support a conserved role for this arginine in conferring high affinity to TRAP transporters mediated by an electrostatic binding interaction in the SBP subunit alone.

Our data support the essential function of Arg-147 in SiaP for the effective colonization of the host by the H. influenzae (9, 31). The bacterium obligately depends on scavenging sialic acid from the host via SiaPQM to incorporate it into LOS and confer serum resistance (9, 31). In this pathogen, the transporter is co-transcribed with a gene encoding a periplasmic sialic acid mutarotase (NamM, H10148), which helps the bacterium scavenge the limited amount of α-Neu5Ac present in the environment to the β-form that is transported by SiaPQM (32). Our
structural data agree with microbiological work demonstrating that LOS sialylation is lost in a bacterium containing an R147A SiaP mutation in the presence of 100 μM Neu5Ac (8).

The second significant finding from this study is that water molecules can partially replace the function of the missing guanidinium group of Arg-147. Nonetheless, faster growth with the R147K mutant over both other mutants (Fig. 2) suggests an affinity benefit from even one direct charge-mediated interaction with the ligand over losing both direct contacts compared with using two water-mediated interaction (Fig. 3, B and C). Yet this “plasticity” in the binding site can recover function at high non-physiological (millimolar) substrate concentrations. It also allowed us to engineer an R147E mutant that positions two water molecules in the space previously occupied by the guanidinium group of Arg-147. Our crystal structure confirmed that the mutant protein is able to use water-mediated contacts to bind sialylamide, an atypical ligand for SiaP. Crystallization of the wild-type SiaP with the same preparation of sialylamide revealed that the arginine is able to “fish out” even the minimal amounts of Neu5Ac present in this sialylamide preparation, analogous to conditions of limited environmental nutrients. These data also suggest that sialylamide probably is not a true ligand for wild-type SiaP, and binding detected in Muller et al. (5) was likely due to this low level of contamination with Neu5Ac.
High Affinity Substrate Recognition in TRAP Transporters

In addition to the two geometrically engineered water molecules in the R147E/sialylamide structure, there is further evidence of using water to neutralize ligand charge in other SBP subtypes; the bridging of negatively charged aspartate ligands by two water molecules to the protein backbone was suggested to be a general feature of 59 out of 78 Bug proteins (33), which can be components of bacterial tripartite tricarboxylic acid transporters (34).

Although there is now strong support for the arginine being central to TRAP transporter ligand binding, there are small fractions of DctP-TRAP SBPs that do not have an equivalent arginine. In our Pfam analysis of the SBP-bac-7 family of 6142 sequences, 3% (199) lack the Arg-147 equivalent. The Arg is most commonly substituted for a Phe, a hydrophobic aromatic amino acid. Interestingly, the structure of the Tp0957 TRAP SBP from Treponema pallidium revealed a unique hydrophobic binding pocket where the Arg-147 equivalent is an alanine, which may define a small group of TRAP transporters likely binding hydrophobic substrates (35, 36), although the exact nature of these ligands is currently unknown. Within the study by Vetting et al. (18), the structure of Chromohalobacter salexigenus DSM 3043 CsaL_0678 protein, which lacks the Arg-147 equivalent, binds an atypical TRAP ligand, ethanolamine, reinforcing the strong link between the Arg and selectivity of carboxylate-containing ligands as typical TRAP substrates.

In conclusion, our data provide the first structural, biochemical, and mechanistic insight into the importance of Arg-147 in TRAP transporter biology. Its function lies in conferring high ligand specificity and affinity to the TRAP transporters and enables function in environments with limited amounts of carboxylate-containing ligands.

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