Structural basis for the glycosyltransferase activity of the Salmonella effector SseK3

Diego Esposito¹, Regina A. Gunster², Luigi Martino¹, Kamel El Omari³, Armin Wagner³, Teresa L. M. Thurston², Katrin Rittinger¹*

¹Molecular Structure of Cell Signalling Laboratory, The Francis Crick Institute, 1 Midland Road, London NW1 1AT, United Kingdom; ²Section of Microbiology, Medical Research Council Centre for Molecular Bacteriology and Infection, Imperial College London, London, United Kingdom; ³Diamond Light Source, Harwell Science and Innovation Campus, Chilton, Didcot OX11 0DE, United Kingdom

Running title: SseK3 is a retaining type-A glycosyltransferase

*To whom correspondence should be addresses: Katrin Rittinger: Molecular Structure of Cell Signalling Laboratory, The Francis Crick Institute, 1 Midland Road, London NW1 1AT, United Kingdom; katrin.rittinger@crick.ac.uk; Tel (44) 020 37962274
Teresa L. M. Thurston: Section of Microbiology, Medical Research Council Centre for Molecular Bacteriology and Infection, Imperial College London, London, United Kingdom; t.thurston@imperial.ac.uk; Tel (44)

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ABSTRACT

The Salmonella secreted effector SseK3 translocates into the host cells, targeting innate immune responses including NF-κB activation. SseK3 is a glycosyltransferase that transfers an N-acetylgalactosamine (GlcNAc) moiety onto the guanidino group of a target arginine, modulating host cell function. However, a lack of structural information has precluded elucidation of the molecular mechanisms in arginine and GlcNAc selection. We report here the crystal structure of SseK3 in its apo form and in complex with hydrolysed UDP-GlcNAc. SseK3 possesses the typical glycosyltransferase type-A (GT-A) family fold and the metal-coordinating DXD motif essential for ligand binding and enzymatic activity. Several conserved residues were essential for arginine-GlcNAcylation and SseK3-mediated inhibition of NF-κB activation. Isothermal titration calorimetry revealed SseK3’s preference for manganese coordination. The pattern of interactions in the substrate-bound SseK3 structure explained the selection of the primary ligand. Structural re-arrangement of the C-terminal residues upon ligand binding was crucial for SseK3’s catalytic activity and NMR analysis indicated that SseK3 has limited UDP-GlcNAc hydrolysis activity. The release of free N-acetyl α-D-glucosamine, and the presence of the same molecule in the SseK3 active site, classified it as a retaining glycosyltransferase. A glutamate residue in the active site suggested a double-inversion mechanism for the arginine N-glycosylation reaction. Homology models of SseK1, SseK2, and the Escherichia coli orthologue NleB1, disclosed differences in the surface electrostatic charge distribution possibly accounting for their diverse activities. This first structure of a retaining GT-A arginine N-glycosyltransferase provides an important step towards a better understanding of this enzyme class and their roles as bacterial effectors.

Salmonella enterica is an intracellular Gram-negative pathogen for which strains of the various serovars cause many diseases in humans and animals worldwide. In immune-competent humans, non-typhoidal Salmonella (NTS) serovars, including Salmonella enterica serovar Typhimurium, typically cause self-limiting gastroenteritis (1). However, in immune-compromised individuals, NTS serovars frequently cause...
an invasive disease that results in significant morbidity (2). Following invasion or phagocytic uptake into the host cell, one of the key virulence determinants is the *Salmonella* pathogenicity island 2 (SPI-2)-encoded type III secretion system (T3SS), which delivers approximately 28 effector proteins into the host cell (3-5). A number of *Salmonella* effectors target and modify host proteins that have a role in mediating host inflammatory responses (3,6). Three highly related *Salmonella* effectors, SseK1, SseK2 and SseK3, orthologues of the enteropathogenic and enterohemorrhagic *Escherichia coli* T3SS effector NleB1 (7), translocate via the T3SS into the host cell (8). Similarly to NleB1 (9,10), SseK1 and SseK3 are N-acetylglucosamine (GlcNAc) transferases that modify the TNFR1-associated death domain protein TRADD and inhibit activation of the pro-inflammatory transcription factor NF-κB, as well as necroptotic host cell death (8). In in vitro experiments, SseK2 modifies the FAS-associated protein with death domain (FADD) (11) but despite detectable translocation into host cells, SseK2 is only able to inhibit NF-κB when highly overexpressed in 293ET cells (8). This suggests that either a loss of catalytic activity or of substrate interaction occurs after delivery of physiologically relevant levels of SseK2. Whether SseK2 has unidentified functions during infection that are mediated by different targets or a different sugar modification remains unknown.

Unlike mammalian O-linked GlcNAcylation, where GlcNAc is attached to the oxygen of the hydroxyl group of serine and threonine residues of numerous cytosolic and nuclear proteins, including those that mediate signalling (12), GlcNAcylation mediated by NleB from pathogenic *E. coli* or SseK bacterial effectors results in addition of GlcNAc to arginine residues with a N-glycosidic linkage (9,10). This distinct modification can be recognized in mammalian cells using an antibody that does not detect O-linked GlcNAcylation (13). Both, mammalian GlcNAcylation and bacterial-mediated arginine-GlcNAcylation, involves the transfer of sugar from an activated uridine-diphosphate carbohydrate donor substrate, UDP-GlcNAc. NleB and SseK proteins have a conserved DXD motif, typical of GT-A family of glycosyltransferases, where the aspartic side chains are required for the coordination of a metal divalent cation necessary for their enzymatic activity (14).

Whilst, to date, there are a number of structures describing O-glycosyltransferases, and the O-GlcNAcylation catalytic event, the only known structure of a bacterial arginine-specific glycosyltransferase is that of the B pattern-type inverting glycosyltransferase EarP, which modifies translation elongation factor P (EF-P) by arginine rhamnosylation (15,16). Another study reported glucosylation of arginine by the sweet corn protein amylogenin, which mediates self-glucosylation in vitro (17), but there have been no follow up reports.

Sequence analysis suggests a structure for the catalytic core of NleB1 protein similar to that of *Photorhabdus asymbiotica* tyrosine-glycosyltransferase protein toxin (18) and *Clostridium difficile* toxin A and toxin B (19,20), which glycosylate host Rho GTPases involved in the regulation of the host cytoskeleton, and the *Legionella pneumophila* Lgt1 that targets a serine residue of eEF1A, blocking protein biosynthesis (21). As well as the conserved DXD motif in SseK effectors, additional amino acids are conserved, including a tyrosine and glutamic acid residue, equivalent to Y219 and E253 in NleB1, both required for NleB1-mediated arginine-GlcNAcylation of the death-domain containing protein, FADD (22). Arginine-GlcNAcylation of target proteins is not observed in un-infected host cells and is irreversible by host enzymes (23) and therefore represents a potent bacterial-mediated virulence mechanism. However, in the absence of structural information, the molecular details underlying arginine and UDP-GlcNAc selection are unknown.

This study presents the crystal structures of the N-glycosyltransferase SseK3 in its free form and bound to UDP, GlcNAc and manganese. SseK3 adopts the classical GT-A glycosyltransferases family fold and is able to hydrolyse UDP-GlcNAc in the absence of a protein substrate. Mutational
analysis of amino acids predicted to be important for substrate binding identified several conserved residues in SseK3 that are essential for arginine-GlcNAcylatation of TRADD and SseK3-mediated inhibition of TNFα-induced NF-κB activation, directly correlating enzymatic activity to virulence function. Structural analysis reveals the presence of an active site glutamate residue, which is conserved only in NleB and SseK proteins but not in other structurally related glycosyltransferases, and is essential for enzymatic activity. The close proximity of this residue to the reactive anomic carbon supports a double inversion catalytic mechanism where an intermediate enzyme-sugar is stabilised for the nucleophilic attack of the acceptor arginine side chain of the host substrates. This is the first structure of a retaining arginine glycosyltransferase bound to a hydrolysed form of its donor substrate representing an important step towards the understanding of this class of effector proteins.

Results

A divalent cation is necessary for binding of the ligand to SseK3

Mass spectrometry analysis of bacterially expressed full length SseK3 revealed limited proteolysis, with the protein losing its first 13 and last 2 amino acids (data not shown). For the majority of T3SS effector proteins, the N-terminal 15-25 residues, often predicted to be unstructured, are required for translocation and remain uncleaved following delivery into the host cell cytosol (24,25). This region is frequently removed prior to crystallisation, as in the case of the E. coli effector protease NleC (26), or disordered, as in the case of E. coli virulence factor NleE (27). We therefore, prepared two forms of SseK3, 14-333 and 14-335, for subsequent structural and functional characterization. Glycosyltransferases require the presence of a divalent cation for activity. To test the selectivity of SseK3 for divalent cations and UDP-glucose derivatives we compared the ability of SseK3 to interact with Mg\(^{2+}\), Mn\(^{2+}\) and 3 UDP-carbohydrate derivatives: UDP-GlcNAc, UDP-Glc (glucose) and UDP-Gal (galactose) by isothermal titration calorimetry (ITC) (Figure 1 and S1, Table 1). No binding was observed for UDP-GlcNAc to SseK3\textsubscript{14-335} in the absence of a coordinating metal cation (Figure S1). When the titration experiments were performed in the presence of MgCl\(_2\) or MnCl\(_2\), the resulting binding affinity for UDP-GlcNAc in the presence of Mn\(^{2+}\) (K\(_d\) = 1.9 \(\mu\)M) showed a 5-fold increase compared to that for Mg\(^{2+}\) (K\(_d\) = 10 \(\mu\)M) (Figure 1 and S1). This binding affinity is similar to that observed for C. difficile toxin A for UDP-Glc and manganese (K\(_d\) = 11.4 \(\mu\)M) (28). The binding studies showed that ligand interaction is influenced by the presence of the two C-terminal residues in the longer SseK3 construct (W334 and R335), but both SseK3\textsubscript{14-333} and SseK3\textsubscript{14-335} bind with a lower affinity to UDP-Gal, compared to UDP-Glc and UDP-GlcNAc. UDP-GlcNAc has the highest affinity for SseK3\textsubscript{14-335}, although only binds marginally stronger than UDP-Glc, and manifests the largest increase in affinity due to the presence of the two C-terminal residues W334 and R335. Interestingly, the change in affinity appears to be associated with an increase in binding enthalpy and a loss of entropy suggesting the occurrence of a structural reorganization upon ligand binding to SseK3 (Table 1).

Structure of SseK3

Crystals of SseK3\textsubscript{14-333} enriched with seleno-methionine diffracted to good resolution but most of them exhibited severe anisotropy, were non-isomorphous and showed translational non-crystallographic symmetry. Attempts to solve the structure by molecular replacement using available glycosyltransferase X-ray structures or selenium SAD phasing failed. Instead, we recorded data at the I23 long-wavelength beamline at Diamond Light Source to reveal the positions of the selenium and sulphur atoms within SseK3 based on the small anomalous signals present at long wavelengths to obtain an initial model (29). This model was used as template in a molecular replacement search with a complete 2.2\AA resolution dataset that did not exhibit severe diffraction anisotropy. SseK3\textsubscript{14-333} crystal belongs to space group \(P2_12_12_1\) with 2 molecules in the asymmetric unit (a.u.) (Table 2) that overlap with a root
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mean square deviation (RMSD) (Cα of residues 27-328) of 0.91 Å. The structure of apo SseK3 is reported in Figure 2A. Residues 14-25 are not visible in the electron density for either chain, nor are amino acids 329-333 of chain B and the last serine residue (S333) of chain A.

The structure of SseK3 at 2.20 Å in complex with UDP, GlcNAc and Mn²⁺ (Figure 2B) was solved by molecular replacement using the apo structure. SseK3 crystals belong to the same space group as the shorter unbound construct, with 2 molecules in the asymmetric unit (Table 2). The two chains in the a.u. overlap with a RMSD in Ca positions of 0.88 Å for residues 27-335. No electronic density is visible for amino acids 14-26 for either chain in the a.u. whilst all the C-terminal residues are observable. Residues 27-321 of SseK3 in the apo and ligand-bound forms overlap with a RMSD of 0.84 Å while residues 322-332 of SseK3 are unstructured with an average B factor of 45.5 Å², compared to the average macromolecule B factor of 33.3 Å².

The SseK3 catalytic core domain (ca. 250 residues) shows the classic features of the large glycosyltransferase type-A family of enzymes (GT-A) consisting of a single module composed of a central parallel β-sheet core flanked by a number of α-helices (14). The structure contains an additional small domain, spanning residues 134-171, containing two extra α-helices that are here named α-helical insertion (Figure 2). This protruding region, especially the position of the residues in the loop 148-154, is differently tilted in the two SseK3 copies present in the crystallographic asymmetric unit, most likely due to non-identical crystal contacts.

Using the structure of SseK3 as template, a DALI (30) search for homologous structures yielded high scores of 17.4, 16.9, 16.7 and 16.2 for the GT-A family members lethal toxin (LT) from C. sordellii in complex with UDP-Glc (PDB: 2VKD) (31), P. asymbiotica ToxG bound to UDP-GlcNAc (PDB: 4MIX) (18), C. difficile toxin A bound to UDP-Glc (PDB: 3SRZ) (20) and C. difficile toxin B bound to UDP and Glc (PDB: 2BVL) (19), respectively. According to the Carbohydrate-Active Enzymes (CAZY) database (32), all these enzymes are classified as sugar stereochemistry retaining glycosyltransferases. Due to the length of the crystallization process, and to the hydrolysis capabilities of SseK3, in the structure crystallized in the presence of UDP-GlcNAc and Mn²⁺, the ligand is hydrolysed with the UDP and N-acetyl α-D-glucosamine still present in the active site (Figure 2B and S2). A similar observation was made for C. difficile toxB co-crystallized in the presence of UDP-Glc (19). A comparison with the apo structure, reveals that the C-terminal amino acids in the ligand-bound SseK3 fold back onto the ligand to produce a short segment of α-helix, with the last two residues W334 and R335 directly interacting with the UDP and GlcNAc (Figure 3B and 3C).

To investigate the conformation of this C-terminal region in the apo state we recorded a 1D proton NMR spectrum of apo SseK3 (Figure S2C), likely corresponding to the side chain Heps of the C-terminal residue W334. The sharp signals in the spectrum of the free protein are broadened when UDP-GlcNAc and MgCl₂ are added to the solution suggesting a conformational rearrangement and loss of flexibility. Reorganization of the -terminus in the SseK3 structure upon ligand binding correlates both with the NMR downfield resonances collapse in the 1D proton spectrum and with the loss of entropy upon binding detected by the ITC experiments.

The active site

SseK3 contains the classic GT-A DXD motif (D²AD²⁸) that coordinates the divalent manganese cation in an octahedral geometry (Figures 3A and 3B). Mn²⁺ interacts with the carboxylate oxygens of
D228 and D325, the UDP diphosphate and the hydroxyl group of S327, as well as an ordered water molecule, which bridges, via hydrogen bonds, the metal atom and the carboxyl group of D226 of the DXD motif (Figure 3B). All the metal coordinating residues are conserved across this effector family apart from SseK3 D325 that is an asparagine residue in the other family members (Figure 3A). Most C-terminal residues participate in interactions that lock the phosphate groups in the correct orientation. S333, W334 and R335 all interact via hydrogen bonds with the oxygen atoms of the diphosphate. The last residue visible in the structure of SseK314,333, serine 332, translates 17 Å from its position in the apo structure to interact directly with a β-oxygen of the UDP (Figure 3B).

The aromatic portion of the ligand is held in place by a series of interactions that include hydrogen bonds of the backbone amide and carbonyl of F53 with the N3 and O2 of the ring and the π-stacking of the uracil by W52 and F190 (Figure 3C). The ribose C2 hydroxyl group interacts with both the backbone carbonyl of Q51 and the hydroxyl group of Y224, whilst the C3 –OH group forms a hydrogen bond with the backbone amide of A227. The N-acetyl α-D-glucosamine in the active site is stabilized by a series of interactions that involve the anomeric C1 hydroxyl group, at a distance of 3.6 Å from the phosphorous atom to which it was bound, and the guanidine group of R335, with hydrophobic interactions also established between the sugar and the aromatic ring of W334 (Figure 3C). The C2 acetyl group oxygen, absent in UDP-Glc, forms a hydrogen bond (3.13 Å) with the backbone amide of G260. The same interaction can be seen between G405 and the sugar acetyl group in the structure of the pneumococcal transferase GtFA in complex with UDP and GlcNAc (PDB 4PQG) (33) and might contribute to the binding preference of SseK3 for UDP-GlcNAc compared to UDP-Glc. The structural position of G260 is substituted by a bulky glutamine side chain in clostridial toxin A and toxin B (Figure 3C) suggesting an explanation for preference of these toxins for UDP-Glc (19,20).

The GlcNAc C4 and C6 hydroxyl groups in the structure of ligand bound SseK3 are locked in the active site by the side chain carboxylate of D191, which may explain why UDP-Gal, which has a C4 inverted stereochemistry compared to glucose, shows the lowest affinity for SseK3.

The UDP and GlcNAc present in the SseK3 catalytic pocket are in an arrangement similar to UDP-Glc bound to toxA and the hydrolysed UDP and Glc in toxB (Figure S2) (19,20). Differently from the clostridial toxins, the active site in SseK3 contains a glutamic acid residue, E258, 3.2 Å away from the anomeric carbon. This residue is conserved in all SseK effectors and NleB, but is replaced by an isoleucine in toxA, toxB and LT (19,20,31). Previous data suggested an important role of this residue for NleB1 function (22).

**Analysis of UDP-GlcNAc hydrolysis**

The presence of hydrolysed UDP-GlcNAc in the active site shows that, in the absence of a protein substrate, SseK3 can function as a hydrolase. To further explore the ability of SseK3 to hydrolyse UDP-GlcNAc we used 31P and 1H-NMR spectroscopy. At time 0 (Figure 4), the 1D 31P-spectrum of UDP-GlcNAc shows two doublets signals for the Pα, at -13.06ppm, and Pβ, at -11.41ppm, for the diphosphate group of the UDP. A well-resolved quadruplet signal system in the 1D proton spectrum of the intact UDP-GlcNAc at 5.43ppm, can also be observed for the anomeric H1 proton of the glucose moiety, resulting from the H1-H2 (3.56 Hz) and H1-Pβ (7.02 Hz) coupling (34,35). Addition of SseK314-335 results in a progressive reduction of both NMR signals with time. The 31P spectrum (Figure 4A) shows the emergence of the Pα and Pβ doublet signals from UDP in a free form, whilst the quadruplet signal of the sugar C1 proton (Figure 4B) gradually disappears, substituted by a rising doublet at 5.12ppm corresponding to the same proton no longer attached to the β-phosphorus of the UDP. The doublet has a \( J_{H1,H2} = 3.58 \) Hz, a value typical for an α-anomeric equatorial-axial stereochemistry for the sugar C1 (the β-form has a value
around 7-8 Hz) (36,37). The chemical shift and proton-proton coupling constant of the glucosamine derived from the hydrolysis of the UDP-GlcNAc are identical to those in the spectrum of free N-acetyl α-D-glucosamine. Moreover, mass spectrometry analysis of the reaction mixture does not show SseK3 modification (data not shown) excluding the possibility of intramolecular N-GlcNAcylation.

The UDP-GlcNAc hydrolysis proceeds with a retention of the C1 chirality suggesting that SseK3 is a retaining glycosyltransferase enzyme. Although able to hydrolyse the ligand in the absence of its target substrate, SseK3 does it slowly. There is a reduction of 36 ± 2% of the signal intensities in the 31P-spectrum for the α- and β-phosphorus nuclei of the diphosphate group of the UDP-GlcNAc after 90 minutes at 30 °C. The peaks are still visible in the spectrum, albeit very close to the noise level, after the reaction is allowed to proceed for 12 hours. UDP-GlcNAc hydrolysis is not observed with SseK314,333 (Figure S3A) – or in the absence of enzyme (Figure S3B) – indicating that together, the C-terminal residues W334 and R335 are necessary for SseK3 functional activity.

**Structure based analysis of SseK3 function**

To test the role of active site residues in the enzymatic activity of SseK3 in a functional framework, multiple amino acid substitutions were tested for their ability to modify the SseK3 substrate TRADD, as well as prevent NF-κB activation. We focused on amino acids from our structural analysis of SseK3 (Figure 3) that are required for coordination of UDP and GlcNAc and hence enzymatic activity. In addition, we were particularly interested to test whether E258 was critical for enzymatic activity as this residue is only conserved in NleB and SseK glycosyltransferases and has been shown to be required for the function of NleB1 (22).

Individual SseK3 mutants were co-transfected into 293ET cells together with FLAG-TRADD, and following anti-FLAG immuno-precipitation the GlcNAcylation of TRADD was analysed by immuno-blotting. As a control, transfected SseK3 K251A remained active like wild-type SseK3. In contrast, and as expected, mutation of the manganese coordinating DXD (D226A/D228A) motif ablated SseK3-induced arginine-GlcNAcylation in cell lysates and the modification of immuno-precipitated TRADD (Figure 5A) (8). The conserved glutamate (E258, predicted to coordinate GlcNAc), which is present in SseK and NleB proteins but not other structurally related glycosyltransferases, was also essential for arginine-GlcNAcylation of TRADD.9 In addition, mutations predicted to be required for coordination of uracil (W52A and Y224A) and GlcNAc (D191A, N259A and W334A) also eliminated SseK3-induced protein arginine-GlcNAcylation and more specifically, no modification of TRADD was detected (Figure 5A). Transfected SseK3 R194A (predicted to disrupt GlcNAc binding) appeared inactive but this mutant was not stably expressed and should be approached with caution (Figure 5A). Surprisingly, SseK3 with a mutation in the C-terminal residue R335 (to alanine) retained the ability to modify numerous proteins, including TRADD. However, the pattern of GlcNAcylation was different, with significantly fewer proteins modified when compared to wild-type SseK3 (Figure 5A).

Next, to test if enzymatic activity correlated with the ability of SseK3 to inhibit TNF-α-induced NF-κB activation, variants of SseK3 were transfected into 293ET cells together with an NF-κB-dependent luciferase reporter plasmid. We predicted that Q51A would still be catalytically active as it is the backbone carbonyl group and not the side chain that is involved in interactions with the ligand (Figure 2C). Interestingly, whilst we were not able to detect arginine-GlcNAcylation of TRADD, SseK3 Q51A was still a potent inhibitor NF-κB activation (Figure 5), suggesting that catalytic activity is retained but greatly reduced to below the level of detection. The side chain Nε of residue H247 is 4.4 Å away from the oxygen of the GlcNAc acetyl group, yet a similar finding was observed for SseK3 H247A, with inhibition of NF-κB activation occurring without detectable modification of TRADD (Figure 5). This suggests that very low and non-detectable levels of arginine-GlcNAcylation
Discussion
Here we report the structure of the Salmonella arginine N-glycosyltransferase, SseK3, in its apo and ligand bound forms. The protein belongs to the GT-A family of glycosyltransferase enzymes and binds the ligand in a metal ion dependent manner via a DXD motif. The dynamic features of the last 15 residues of SseK3 are important as they experience a large conformational change upon ligand and metal binding. As suggested by solution NMR spectroscopy, the C-terminal residues are disordered in the unbound SseK3 structure, with structural reorganization following primary substrate binding, with an open-close catalytic cycle that, as in the case of a number of other glycosyltransferases, is likely to support SseK3 enzymatic activity (38,39). Residue W334 in the active site pocket occupies the same structural position as W520 in the catalytic fragment of the clostridial lethal toxin in complex with UDP-Glc and manganese (PDB 2VKD) (31) and it is conserved in toxin A (W519) (20) and toxin B (W520) (19) of the same organism (Figure 6 and S2). A tryptophan residue (W520) is also present in the active site of the L. pneumophila Lgt1 in complex with UDP-Glc and magnesium (PDB 3JSZ) (21) and in a structurally equivalent position in the unbound form of the α-toxin from C. novyi (PDB 2VK9) (31). As for the other toxins, W334 is a pivotal residue in the opening and closure of the catalytic cleft (40). In the ligand bound SseK3, its Nε1 forms a hydrogen bond with the UDP Pβ oxygen whilst the equivalent tryptophan residue (W519) in the structure of toxin A hydrogen bonds the glycosidic oxygen of the intact UDP-Glc (Figure S2B) (31). W334 is therefore in a position to stabilize the emerging negative charge of the former glycosidic oxygen transferred onto the UDP. Once the enzymatic reaction has occurred, the opening of the catalytic site and the release of the UDP reinitiate the cascade; indeed mutation of W334 ablates the catalytic and functional activity of SseK3.

The presence of a hydrolysed UDP-GlcNAc in the protein active site suggests that SseK3 can function as a hydrolase. A process observed also in the case of toxin A (28) and toxin B (19). The hydrolysis of UDP-GlcNAc, in the absence of target substrate, is likely to be a proxy of its physiological glycosyltransferase activity and the observed release by NMR of a α-anomeric form of GlcNAc, and presence of the same molecule in the enzyme active site, indicates that SseK3 is likely to be a retaining N-glycosylating enzyme. Reaction mechanisms of retaining glycosyltransferases have been controversial but it is generally accepted that they operate through the stabilization of an oxocarbenium-like transition state (41). Depending on the presence of a nucleophile in the active site correctly positioned on the β-face of the donor substrate, the reaction could proceed either via a front face or double inversion mechanism (42). For the clostridial O-glycosylating toxins, in the absence of other available bases in the active site, the deprotonation of the acceptor nucleophile is carried out by the nucleotide Pβ oxygen. Following the formation of a transient oxocarbenium state, the departure of the leaving sugar O1 and the formation of the new glycosidic bond occur on the same side of the anomeric carbon, retaining its stereochemistry (31). In the double displacement reaction, the presence of a nucleophile in close proximity to the anomeric carbon, directly under the sugar substrate β-face, stabilizes an oxocarbenium-like enzyme-ligand adduct. The reaction is then followed by a subsequent direct substitution by the target nucleophile at the opposite site of the anomeric C1 with a double inversion mechanism that retains the original carbon stereochemistry. This is the case for
lysozyme (43) and GT6 family member α3GalT (44). For these enzymes, a well-located nucleophile within the active site - D52 for lysozyme and E317 for α3GalT - is a necessary element for the creation of a stereochemically inverted intermediate set for nucleophilic substitution. Despite their structural similarity, the presence of a glutamate side chain in the active site of SseK3, 3.2 Å from the anomeric carbon, suggests that this enzyme could operate as a retaining GT more similarly to α3GalT than to the clostridial toxins. The E258 side chain is important for SseK3 catalytic activity and is a conserved active site residue across the SseK effectors family and in NleB1, substituted by uncharged residues in structurally related toxins (Figure 6). Mutation into an alanine residue, similarly to the mutation of the equivalent residue in NleB1 (E253A), impairs the arginine-GlcNAcylation ability of SseK3 and NleB1 without, for the latter, abrogating binding to the target protein (22). To date the only available structure of an arginine-glycosylation enzyme is that of the *Pseudomonas aeruginosa* EarP that catalyses the transfer of a rhamnose molecule onto the bacterial translation elongation factor EF-P (16). EarP is an inverting glycosyltransferase belonging to the GT-B family and structurally unrelated to SseK3. Interestingly, EarP has three negatively charged residues in the active site (D13, D17 and E273), in close proximity to the sugar, that have been shown to be important in the catalytic activity and play a crucial role in the stabilization of the positive charge of the acceptor guanidino group. Mutation of those residues to alanine, as for the E258A mutation in SseK3, severely impairs EarP enzymatic activity without disrupting substrate binding (16).

We, therefore, propose a catalytic mechanism, in which the glutamate side chain of E258 acts as the intramolecular nucleophile in the first step of a double displacement mechanism, forming an SseK3-GlcNAc intermediate state primed for the C1 nucleophilic attack to the sugar α-side by the guanidino group on the target arginine. E258 could aid in the selection of the arginine as the N-GlcNAcylation site and, similarly to the phosphate role in the clostridial toxins, increase the nucleophilic character of the acceptor by deprotonation of its side chain. A double displacement mechanism, where the C1-O1 bond is broken before the nucleophilic attack, could also be favoured by the presence of the guanidium group of R335 tucked in SseK3 active site. R335 sits above the sugar O1, obstructing the access to the sugar α-side, sterically hindering a direct front-face reaction.

In SseK3, the well-located R335 side chain hints to a potential intramolecular GlcNAcylation process. However, *in vitro* auto-GlcNAcylation of SseK3 is unlikely as NMR and mass spectrometry experiments only detected unbound glucosamine following UDP-GlcNAc hydrolysis by SseK3. Nevertheless, a specific role of this residue in SseK3 function cannot be excluded. In the *E. coli* MurG, an O-GlcNAc GT-B family glycosyltransferase, the presence of a positively charged arginine residue (R261) in the active pocket provides extra stability for the negatively charged UDP emerging from the catalytic reaction (45). In the structure of the UDP and GlcNAc bound GtfA, an O-GlcNAc GT-B, an active site arginine residue (R328), crucial for catalytic activity, forms a strong hydrogen bond with the Pβ oxygen, stabilizing the negative charge generated from the processing of the substrate (33). The active site of EarP has an arginine residue (R271), whose guanidino Nε1 is 2.5 Å from the phosphate oxygen that participates in the stabilization of the enzymatic products whose mutation into an alanine reduces EarP glycosylation activity (16). In SseK3, R335 is tucked in the active site at 2.62 Å from the sugar O1 and 3.24 Å from the oxygen beta phosphate and its mutation R335A induced a severely reduced arginine-GlcNAcylation pattern compared to wild-type SseK3 (Figure 5). As for EarP, this could reflect a decrease in enzymatic activity due to a decreased stabilization of the enzymatic products. Alternatively, as this residue is only conserved in SseK2, which like SseK3 localises to the Golgi network during infection (8), but is substituted by an alanine in SseK1 and...
absent in NleB1, it could be indirectly involved in substrate specificity via reduced Golgi network association.

During infection of macrophages, similarly to their *E. coli* ortholog, SseK1 and SseK3 both function to mediate inhibition of NF-kB activation as well as inhibition of an inflammatory necrototic host cell death (8). The structure of SseK3 bound to its donor substrate hydrolysis products revealed which amino acids are necessary for ligand binding and hence arginine-GlcNAcylation of TRADD and SseK3-mediated inhibition of NF-kB activity. In addition to the ion-coordinating DXD motif, previous mutational analysis of NleB1, identified residue Y219 as required for function (22). This amino acid is conserved in SseK proteins (Y224) and is important for the coordination of the uracil of the UDP and therefore necessary for both SseK3 catalytic activity and function.

SseK effector proteins share a high sequence identity but most of the conserved residues are those in the catalytic domain (Figure 1 and S5). Sequence variability is mainly concentrated around residues in the α-helical insertion suggesting that this region might direct substrate specificity. Despite its similarity to the other effectors, SseK2 does not show activity in cell-based assays (11). However, purified SseK2 can arginine-GlcNAcylate FADD *in vitro*, without forming a stable interaction, suggesting that this enzyme is functional (11). As all residues tested that are required for enzymatic activity in SseK3 are conserved in SseK2 it is not immediately clear why SseK2 is not active after translocation into host cells. In the absence of structural information for the other members of the SseK family, we created homology models of SseK1, SseK2 and NleB1 based on the SseK3 crystal structure (Figure 6). The distribution of surface charges shows that both SseK1 and NleB1 have the active site surrounded by acidic residues whilst the surfaces of SseK2 and SseK3 appear to be more similar, with 72% sequence identity, having the ligand site surrounded by larger positively charged areas compared to SseK1. Also, as the C-terminal residue in both SseK2 and SseK3 is a positively charged residue (R348 and R334), substituted by a smaller side chain in SseK1 (A332) and absent in NleB1, the active site in SseK1 and NleB1 appears to be more accessible than in SseK2 and SseK3. Recently, it has been reported that NleB1 preferentially GlcNAcylates the death domain (DD) of FADD at residue R117 (23), although it can interact with and modify the DD of other proteins including TRADD and RIPK1 (9,10). Interestingly, SseK1 is the only SseK family member that can modify FADD (also at R117) after bacterial delivery into host cells (8). The structure of FADD-DD shows an extensive basic surface patch whose residues have been shown to be important in the interaction with Fas receptor (CD95) (46). In particular, mutation of R117, part of the FADD-DD extended positively charged surface, has been shown to abolish interaction with the cytoplasmic death domain of CD95 (47). Therefore, the contiguous acidic surfaces of SseK1 and NleB1 could act as a complementary surface, binding the death domain of FADD and allowing the side chain of residue R117 to position itself in the active site for sugar transfer. The lack of activity in cell-based assays of SseK3 towards FADD, and the absence of a phenotype in *Salmonella* for SseK2, could then be the result of the interplay between a less accessible active site and altered substrate specificity, with SseK3 still able to modify TRADD.

In summary, we have solved the first X-ray crystal structure of a GT-A arginine-glycosyltransferase, identifying core-conserved residues that are required for catalytic activity and virulence function of SseK3 allowing us to propose a potential enzymatic mechanism for N-arginine glycosylation.

**Experimental procedures**

**Protein cloning, expression and purification**

Wild type SseK314-333 and SseK314-335 constructs (Uniprot: A0A0H3NMP8) were cloned in pGEX-6P1 (GE Healthcare) vector by standard Gibson Assembly protocol (48) and expressed with a cleavable N-terminal GST tag. The proteins were expressed in *E. coli* BL21 (DE3) Gold in LB supplemented with 100 µg/ml ampicillin. Cells were grown at 30 °C until OD600 ~ 0.6-0.8, induced with 0.5 mM
SseK3 is a retaining type-A glycosyltransferase

isopropyl β-D-1-thiogalactopyranoside (IPTG) and incubated at 18 °C for overnight expression. Proteins were purified by GST-Sepharose affinity chromatography followed by cleavage of the GST N-terminal tag by 3C-protease and size-exclusion chromatography in 25 mM HEPES buffer pH 7.5, 150 mM NaCl and 0.5 mM TCEP. The 3C-cleavage leaves amino acids GPLGS preceding the first amino acid of SseK3. SseK3 mutant plasmid construction

SseK3 mutant variants were created by overlap mutagenesis PCR from the m4pGFP-SseK3 wild-type plasmid (8) and ligated into the mammalian expression vector m4pGFP to create plasmids encoding SseK3 mutants with an N-terminal GFP-tag. All plasmids were checked by sequencing.

Cell culture

293ET cells (gift from Felix Randow) were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Sigma) supplemented with 10% foetal calf serum (Sigma) at 37°C in 5% CO₂.

Isothermal titration calorimetry

ITC experiments were performed at 293 K using a Microcal iTC200 calorimeter (Malvern). The protein solutions were prepared in buffer containing 25 mM HEPES buffer pH 7.5, 150 mM NaCl, 0.5 mM TCEP and either 5 mM MgCl₂ or MnCl₂. All experiments were performed by placing the solution containing SseK3 in the cell at concentrations of 50 µM and the solution containing the UDP-glucose derivatives in the syringe at 500 µM. The concentration of UDP-derivatives were estimated using the UDP extinction coefficient at λ = 262 nm of 9800 M⁻¹ cm⁻¹. For each titration 20 injections of 2 µL were performed. Integrated data, corrected for heats of dilution, were fitted using a nonlinear least-squares algorithm to a 1:1 binding model, using the MicroCal Origin 7.0 software package. The fitting parameters are ΔH (reaction enthalpy change in kcal·mol⁻¹), K_d (equilibrium binding constant in M⁻¹), and n (number of binding sites). The entropic contribution values (defined as -TΔS) were calculated from the values of ΔH and K_d. Each experiment was repeated at least twice and average values are reported in Table 1.

Crystallization

Initial screens for SseK3 were set up by sitting drop method at two protein concentrations (20 and 10 mg/mL) by combining 0.1 µL of protein solution with 0.1 µL of reservoir. Commercially available crystallization screens were dispensed by using an automated Mosquito machine (TTP Labtech). The most promising initial crystal hits grew from drops set up with 10 mg/mL protein solution in 0.1 M Tris pH 8.5, 0.2 M NaCl and 25% PEG3350. No single crystals were obtained from the initial screening and multiple rounds of seeding were necessary to improve crystal quality. The seed stock was prepared by harvesting the small crystals, obtained in 0.1 M Tris pH 8.5, 0.2 M NaCl and 25% PEG3350, in 50 µL solution of the same condition. The stock was then homogenised by vortexing for 30 sec using a Hampton seed bead. Sitting drops were dispensed by mixing 0.2 µL of protein solution at 9 mg/ml, 0.18 µL of reservoir solution (0.1 M Tris pH 8.5, 0.2 M NaCl and 25% PEG3350) and 0.02 µL of seed stock. Single crystals of better diffraction quality could be grown with the Seleno-methionine SseK3 protein sample in the same condition but using a total protein concentration of 7 mg/mL. For X-ray data acquisition crystals were cryo-protected
with mother liquor containing 0.1 M TRIS pH 8.5, 0.2 M NaCl, 25% PEG3350 and 20% ethylene glycol. To crystallise SseK3$_{14-335}$ in complex with UDP-GlcNAc a stock solution was prepared by mixing the protein at 10 mg/mL with a three-fold molar excess of UDP-GlcNAc (900 µM) and a final concentration of MnCl$_2$ of 5 mM. Small single crystals were obtained in 0.1 M TRIS pH 8.0, 0.2 M NaI and 20% PEG3350 and improved by multiple rounds of seeding. For X-ray data acquisition crystals were cryo-protected with mother liquor containing 0.1 M TRIS pH 8.0, 0.2 M NaI, 20% PEG3350 and 30% trehalose. All the crystallization experiments were done at 20 °C.

Data collection, phasing and refinement
Despite keeping crystallisation and cryo-protection protocols the same, in total four polymorphs were observed for SeMet-labelled SseK3$_{14-333}$ crystals. Data from a monoclinic crystal were collected at the long-wavelength MX beamline I23 at Diamond Light Source (29) for experimental SAD phasing using a wavelength of $\lambda = 2.7751 \text{ Å}$ and processed with XDS (51). The anomalous substructure of 20 selenium and 20 sulphur atoms (corresponding to 4 molecules in the asymmetric unit) was located by SHELXD (52) at 3.5 Å resolution. The substructure solution was confirmed by the presence of a 4-fold non-crystallographic symmetry (NCS). Initial experimental electron density maps were of poor quality possibly due to the presence of a pseudo-translation and the anisotropy of the data. Since automatic building programs failed to provide a starting model, a careful inspection of the density map revealed the position of two helices that were placed as poly-alanine helices using Coot (53). The 4-fold NCS symmetry was applied to locate the corresponding helices in the asymmetric unit. The anomalous substructure and the initial model of the asymmetric unit comprising 8 helices were fed to CRANK2 (54) to produce improved maps benefiting from 4-fold averaging and an almost complete model.

Complete high-resolution data from for SeMet-labelled crystals of SseK3$_{14-333}$ were collected at Swiss Light Source (Villigen, Switzerland) and processed using DIALS (http://dials.diamond.ac.uk/). The structure of SeMet-labelled SseK3$_{14-333}$ was solved at a single wavelength of 0.97640 Å using the structural model obtained by long-wavelength S-SAD as template in a molecular replacement with MOLREP (55). Data for SseK3$_{14-335}$ bound to UDP-GlcNAc and Mn$^{2+}$ were collected at Diamond Light Source, I04 (Oxford, UK) and processed using XDS (51). The structure of SseK3$_{14-335}$ co-crystallized in the presence of UDP-GlcNAc and Mn$^{2+}$ was solved using the apo-structure as template for molecular replacement with MOLREP (55). Models were iteratively improved by manual building in Coot (53) and refined using REFMAC5 (56) and Phenix (57). The parameters for the ligand stereochemistry were obtained from the standard Coot library. All structural figures were prepared in Pymol (Schrödinger, LLC) and Chimera (58). Further details on data collection and refinement statistics are summarised in Table 2.

NF-κB reporter assays
1 × 10$^5$ 293ET cells (seeded the day before use) were transfected with a mixture of 50 ng p4kB:Luc, 20 ng pRLTK and 500 ng m4pGFP-SseK3 variants (or m6pPAC-FLAG-GFP control) using Lipofectamine 2000 (Invitrogen). After 24 h of transfection, the cells were stimulated with 50 ng/ml human TNFα (Sigma) for 17 h and luciferase activity was measured using the Dual Luciferase reporter assay system (Promega) and a Tecan Infinite200 PRO plate reader. NF-κB regulated luciferase activity was first normalised to Renilla luciferase activity and then the fold activation relative to unstimulated conditions of each SseK3 variant was calculated.

TRADD-modification by SseK3 variants
4 × 10$^5$ 293ET cells were transfected for 40 h with 1 µg m6pPAC-FLAG-TRADD and 1 µg m4pGFP-SseK3 variant (or 1 µg m6pPAC-FLAG-GFP control) plasmid DNA using Lipofectamine 2000 (Invitrogen). Subsequently the cells were harvested and lysed for 30 min on ice in lysis buffer (150 mM NaCl, 0.3% (v/v) Triton X-100, 20 mM Tris-Cl (pH 7.4), 5%
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glycerol, 5 mM EDTA, 1 mM PMSF, 1 mM Benzamidine, 2 µg/ml Aprotinin, 5 µg/ml Leupeptin, 1 mM DTT) and cytoplasmic proteins isolated by 30 min centrifugation at 16,000 \( \times g \) at 4°C. FLAG-TRADD was then immuno-precipitated using anti-FLAG M2 Affinity Gel (Sigma) for 2 h at 4°C, washed and analyzed by SDS-PAGE and immuno-blotting using rabbit anti-FLAG (Sigma), rabbit anti-GFP (Invitogen), rabbit anti-Arginine-GlcNAc (EPR18251, Abcam), rabbit anti-tubulin (EPR16774, Abcam) antibodies and HRP-conjugated anti-rabbit secondary antibody (Dako) for detection.

**Nuclear Magnetic Resonance**

All spectra were recorded at 25 °C on a Bruker AVANCE spectrometer operating at 700 MHz, equipped with a cryogenically cooled quadruple resonance \((^1H, ^15N, ^13C\) and \(^{31}P\)) probe including Z-axis pulse field gradients. Data were acquired and processed with Topspin (version 3.5, Bruker). Reaction solutions of 10 µM of both SseK3 constructs (Santa Cruz Biotechnology, weight >98%) were prepared in 20 mM Tris pH 7.5, 100 mM NaCl, 0.5 mM TCEP and 5 mM MgCl₂. 500 µM UDP-GlcNAc added to the proteins solutions and hydrolysis was allowed to proceed at 30 °C under gentle stirring. The reaction was quenched at different time points by adding 10 mM EDTA (pH 7.5) and 5% D₂O required for the magnetic field lock. The same reaction conditions were used to monitor UDP-GlcNAc hydrolysis in the absence of enzyme. 1D \(^1H\) and \(^{31}P\) NMR spectra were recorded using the standard excitation sculpting pulse sequences zgesp and zgpg30 implemented in Topspin acquisition software.
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**Acknowledgements:** We thank the Mass Spectrometry Science Technology Platform, the Biomedical NMR Centre at the Francis Crick Institute, in particular Geoff Kelly, for access and advice, and the Diamond Light Source (Oxford, UK) and the Swiss Light Source (Villingen, Switzerland) for synchrotron access and Andrew Purkiss from the Structural Biology Science Technology Platform at the Francis Crick Institute for help in data collection and interpretation. We would also like to thank Ramona Duman and Vitaliy Mykhaylyk for their support during the long-wavelength experiments.

**Conflict of interest:** The authors declare that they have no conflicts of interests with the contents of this article.

**Author Contributions:** DE, RG and LM performed experiments and analysed the data. AW and KEO collected data at the I23 beamline and calculated an initial structural model. DE, TT and KR designed the overall research and wrote the manuscript.
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References

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SseK3 is a retaining type-A glycosyltransferase

FOOTNOTES
The work was supported by The Francis Crick Institute (grant number FC001142), which receives its core funding from Cancer Research UK, the UK Medical Research Council and the Wellcome Trust. TT was funded on an Imperial College London Junior Research Fellowship (RSRO_P50016) and RG on an Imperial College London PhD studentship.

The atomic coordinates and structure factors (codes 6EYR and 6EYT) have been deposited in the Protein Data Bank (http://www.pdb.org).
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Table 1
Thermodynamic quantities for the interaction of SseK3 constructs and UDP-glucose derivatives

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<th>Protein/Ligands</th>
<th>Kd (µM)</th>
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<th>-ΔS (cal/mol deg)</th>
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<td>nbd</td>
<td>nbd</td>
<td>nbd</td>
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<tr>
<td>SseK314-335/UDP-GlcNAc/Mg^{2+}</td>
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<td>0.8 ± 0.1</td>
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<tr>
<td>SseK314-335/UDP-GlcNAc/Mn^{2+}</td>
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<td>8.1 ± 0.2</td>
<td>0.9 ± 0.1</td>
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<td>SseK314-335/UDP-Gal/Mn^{2+}</td>
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<td>0.8 ± 0.1</td>
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<tr>
<td>SseK314-335/UDP-Glu/Mn^{2+}</td>
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<td>0.9 ± 0.1</td>
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<td>0.8 ± 0.1</td>
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<td>SseK314-333/UDP-Gal/Mn^{2+}</td>
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nbd: no binding detected
Table 2
Crystallographic data collection and refinements statistics

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<th>SseK3&lt;sub&gt;UDP-GlcNAc&lt;/sub&gt;</th>
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**Data Collection statistics**

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<td>73.61 86.05 95.22</td>
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<td>90 90 90</td>
<td>90 90 90</td>
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**Refinement statistics**

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<tr>
<td>Ligands (Å²)</td>
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Highest-resolution shell values are given in parentheses.
SseK3 is a retaining type-A glycosyltransferase

Figure 1. Binding affinity of the ligand to SseK3. Isothermal Titration Calorimetry curves for the interaction of SseK3_{14-333} and SseK3_{14-335} with UDP-Galactose (UDP-Gal), UDP-Glucose (UDP-Glc) and UDP-Glucosamine (UDP-GlcNAc). The experiments were executed in the presence of 5 mM MnCl$_2$. The integrated heat signals for the interaction were integrated as a function of the molar ratio of titrant to protein in the cell. The data were fitted to a 1:1 binding model and the dissociation constants are reported for each experiment.
Figure 2. Structure of ligand-free and bound SseK3. Ribbon representation of the structure of the unbound SseK3\textsubscript{14-333} (A) and SseK3\textsubscript{14-335}/UDP-GlcNAc/Mn\textsuperscript{2+} complex (B). The figure shows the central GT-A catalytic domain that binds the ligand substrate and highlights the \(\alpha\)-helical insertion that shows variability in the two protomers in the asymmetric crystallographic unit.
SseK3 is a retaining type-A glycosyltransferase

**Figure 3.** Close-up of the catalytic core of SseK3. (A) Sequence alignment of the SseK effectors family and their *E. coli* ortholog NleB. The conserved residues involved in the metal coordination, including the DXD motif, are reported as magenta dots on the sequence alignment. (B) Close-up of the residues that participate in the coordination of the Mn$^{2+}$ ion and (C) in the ligand interaction. (D) Downfield region of the proton NMR spectra of SseK3 showing that the addition of Mg$^{2+}$ and UDP-GlcNAc quashes the sharp resonances present in the spectrum of unbound SseK3. The asterisks show the resonances from a complex multiplet around 5ppm originated from the UDP-GlcNAc (green) and the resonance at 10ppm from the Hε1 of W334 (blue).
SseK3 is a retaining type-A glycosyltransferase

Figure 4. SseK3_{14-335} hydrolyses UDP-GlcNAc in UDP and free N-acetyl α-D-glucosamine. Phosphorous (A) and proton (B) NMR spectra of a 500 µM solution of UDP-GlcNAc in the presence of 10 µM SseK3_{14-335}. The spectra were recorded at different time points: 0 min (blue), 5 min (red), 15 min (green), 30 min (purple), 90 min (yellow) and 12 hours (orange). (C) UDP-GlcNAc hydrolysis reaction: the Pα and Pβ doublets and the H1 multiplets are assigned in the relative spectra and highlighted on the chemical structures of the ligand in its intact and hydrolysed form.
SseK3 is a retaining type-A glycosyltransferase

Figure 5. Screening SseK3 putative catalytic mutants for loss of Arg-GlcNAcylation towards TRADD. (A) 293ET cells co-transfected with FLAG-TRADD and the indicated GFP-tagged variants of SseK3 were lysed and inputs and anti-FLAG immunoprecipitates were analysed by immunoblot. Arg-GlcNAcylation of post-nuclear supernatants (input) and immunoprecipitated FLAG-TRADD was tested using anti-Arg-GlcNAc antibody. Expression of GFP-tagged SseK3 variants was tested using anti-GFP antibody in cell lysates. Antibodies to tubulin were used as a loading control. The shown immunoblots are representative of 4 independent experiments. (B) 293ET cells were co-transfected with an NF-κB-dependent luciferase reporter plasmid, pTK-Renilla luciferase and the indicated GFP-tagged SseK3 mutants for 24 hours prior to overnight stimulation with 50 ng/ml TNFα. Luciferase activity was measured in cell lysates and results are presented as the fold activation relative to unstimulated controls expressing each SseK3 variant. Data are the mean +/- s.e.m of 5 independent experiments. Cell lysates from (B) were analysed by immunoblot in supplementary Figure S4. * P < 0.05. DXD corresponds to the SseK3 D226A/D228A mutant.
SseK3 is a retaining type-A glycosyltransferase.

Figure 6. Glycosyltransferase structural alignment and SseK1, SseK2 and NleB1 homology structural models. Upper panel: Structural alignment of SseK3 with C. sordelli LT (PDB 2VKD, 19% sequence identity), C. difficile toxin A (3SRZ, 17%) and toxin B (2BVL, 15%), L. pneumophila Lgt1 (3JSZ, 11%) and P. asymbiotica toxin (4MIX, 19%). The catalytic important E258 residue is marked with an asterisk in the alignment. Lower panel: Solvent-accessible surface representation coloured according to the electrostatic potential (positive blue, negative red) of the structure of SseK3$^{14-335}$ bound to its UDP-GlcNAc ligand and homology structural models of SseK1 (55% sequence identity with SseK3), Ssek2 (72%) and NleB1 (53%). Ligand bound in the active site is highlighted.