The Type VI Secretion System in *Escherichia coli* and Related Species

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ABSTRACT

The Type VI secretion system (T6SS) is a multi-protein complex widespread in *Proteobacteria* and dedicated to the delivery of toxins into both prokaryotic and eukaryotic cells. It thus participates to inter-bacterial competition as well as pathogenesis. The T6SS is a contractile weapon, related to the injection apparatus of contractile tailed bacteriophages. Basically, it assembles an inner tube wrapped by a sheath-like structure and anchored to the cell envelope *via* a membrane complex. The energy released by the contraction of the sheath propels the inner tube through the membrane channel and towards the target cell. Although the assembly and the mechanism of action are conserved across species, the repertoire of secreted toxins, and the diversity of the regulatory mechanisms and of target cells make the T6SS a highly versatile secretion system. The T6SS is particularly represented in *Escherichia coli* pathotypes and *Salmonella* serotypes. In this chapter we summarize the current knowledge regarding the prevalence, the assembly, the regulation and the roles of the T6SS in *E. coli*, *Salmonella* and related species.

INTRODUCTION

The adaptation of bacterial species in their ecological niche relies on specific regulatory circuits to adapt the metabolism and the growth to the extracellular conditions, but also on the release of molecules – siderophores, exopolysaccharides and/or protein toxins – in the milieu. To facilitate their transport through the physical barriers that the membranes represent, protein toxins are specifically selected and secreted by dedicated machineries named “secretion systems”. Depending on the nature of the machine itself and on the mechanism of transport of the toxins, these secretory pathways are numbered I-IX. Most of these pathways, including Type I (T1SS), Type II (T2SS), Type III (T3SS), Type IV (T4SS or conjugation), Type V (autotransporters, intimin/invasin and two-partner pathways including contact-dependent growth inhibition systems) and curli/fimbriae/chaperone-usher pathways are assembled and active in *Escherichia coli* and related species such as *Salmonella*, *Shigella*, *Enterobacter* and *Citrobacter*, and therefore details regarding their architecture, assembly, mechanism of transport as well as the effectors they deliver are described in the corresponding chapters in *EcoSalPlus*. We will describe here one of the most recently identified secretion pathways, the Type VI secretion system (T6SS).

The T6SS is a multi-protein machine, widespread in Gram-negative *Proteobacteria*, with an over-representation in γ -*Proteobacteria* (1-4). However, T6SS-like machines have been identified and characterized in other phylum such as in Bacteroidetes (5). The assembly of this secretion system requires 13 different subunits, which are usually encoded within a single genetic locus on the chromosome (2, 6) (Fig. 1A). Basically, the T6SS can be viewed as a syringe-like structure anchored to the cell membrane by a trans-envelope complex (3, 4, 7-9) (Fig. 1B). The T6SS syringe is evolutionarily, structurally and functionally related to the puncturing tails of contractile tailed bacteriophages (Fig. 1C), although it is not known whether the genes encoding this structure emerged from co-option of bacteriophage genes (9, 10). A large diversity of toxin effectors has been identified in the recent years, from effectors promoting actin modification and disabling eukaryotic cells to peptidoglycan hydrolases targeting competing bacteria. Therefore the T6SS is a versatile weapon targeting cells ranging from bacteria to mammalian hosts. An example of T6SS-mediated inter-bacterial killing between entero-aggregative *E. coli* (EAEC) and *E. coli* K-12 is shown in Fig. 2.

MECHANISM OF ACTION OF THE T6SS

In the recent years, several aspects of the mechanism of action of the T6SS have been described. Genetic, biochemical and structural characterization of the different T6SS subunits or protein complexes have defined the overall architecture of this secretion apparatus. Time-lapse fluorescence microscopy recordings have provided a dynamic view on how the system works. These data, combined with the knowledge on similar contractile structures such as tail bacteriophages or pyocins, allows to design a model for the mechanism of action (3, 4, 8, 9) (Fig. 3). T6SS biogenesis starts with the assembly of the membrane complex (MC) and the tail assembly platform – or baseplate complex (BC) (Fig. 3a). The syringe, composed of the inner tube and capped by the needle spike and wrapped by a contractile structure, the sheath, then polymerizes to form a several hundreds of nm-long tubular structure (Fig. 3b-c). During T6SS assembly, toxin effectors are loaded into the tube or associate with the spike trimer. Contraction of the sheath propels the inner tube/spike, allowing perforation of the target cell membrane and delivery of the effectors (Fig. 3d). The contracted sheath is then disassembled and recycled by a dedicated ATPase named ClpV (Fig. 3e-f).

We will describe in the next sections the genetic organization, the prevalence of T6SS gene clusters, the regulatory mechanisms underlying their expression, as well as the structure, assembly and roles of this secretion machine, emphasizing the current knowledge on the T6SS in *E. coli*, *Salmonella* and related species.

GENETIC ORGANIZATION AND PREVALENCE OF T6SS GENE CLUSTERS IN *E. COLI*, *SALMONELLA* AND RELATED SPECIES

Type VI secretion system genes are distributed in Gram-negative *Proteobacteria* with an over-representation in γ -*proteobacteria* (1). Therefore, T6SS genes are found in most *E. coli* and *Salmonella* species except the *E. coli* B and K-12 laboratory strains. The genes encoding components and toxins of the Type VI secretion system are usually clustered and grouped into genetic islands (1, 6). The GC content of these regions is generally different from that of the core genome suggesting that they have been acquired by horizontal gene transfer (1, 2, 6). These gene clusters encode the 13 core components of the T6SS, *i.e.*, all the subunits required to assemble a functional apparatus. Additional genes present in these clusters encode toxins and anti-toxins, adaptor proteins that binds both machine components and toxins, as well as auxilliary proteins required for the assembly of the apparatus or genes of unknown function that might be required for the assembly of the T6SS, the recruitment and proper delivery of the toxin (3, 4, 9). According to the gene organization and to homologies/similarities, the *E. coli* T6SS gene clusters categorize in three distinct phylogenetic groups: T6SS-1-3 (Fig. 4-5). This observation suggests that these clusters were present in common ancestors or that genetic exchanges occurred between all these strains that may share the same environment. However, the former hypothesis is likely as (i) each of the phylogenetic group is found in both intestinal (AIEC, EAEC, EHEC, EPEC,...) and non-intestinal (UPEC, APEC, MNEC,...) strains and (ii) these groups are not found in bacteria that share similar environments such as *Salmonella* or *Enterobacter* species (see below). Among these three phylogenetic groups, the T6SS-1 (Fig. 5A) and T6SS-2 (Fig. 5B) gene clusters are the most commonly found in *E. coli* chromosomes. For example, the prevalence of T6SS-1, T6SS-2 and T6SS-3 in APEC genomes is 14.6, 2.4 and 0.8% respectively (11, 12). Interestingly, 85% of the T6SS⁺ APEC strains belong to the virulent phylogenetic groups (11, 12). It is worthy to note that, in

general, the T6SS-2 gene cluster is over-represented in pathogenic strains with high virulence traits. This is particularly clear for entero-aggregative *E. coli* (EAEC), in which T6SS-2⁺ strains (e.g., 042) cause diarrhoea whereas T6SS-2⁻ strains (e.g., 17-2, 34b) fail to elicit diarrhoea in human volunteers (13). However, these prevalence information should be taken with care as this does not necessarily mean that T6SSs are directly involved in pathogenesis, but they may prepare the ground for virulence factors by clearing the niche of potential bacterial competitors. Indeed, T6SS-1 and T6SS-2 gene clusters are also found in non pathogenic strains of *E. coli* such as *E. coli* W (14; Fig. 5B); however, in this strain, the T6SS-1 cluster is inactivated by insertion of a mobile element (14) and it is not known whether it is functional.

With the exception of *Shigella sonnei* and *S. enterica* Gallinarum, which carry T6SS-2-like gene clusters (Fig. 5B), the T6SS genetic organization in *Salmonella*, *Citrobacter* or *Enterobacter* species is distinct from the *E. coli* T6SS-1-3 loci. In *Salmonella enterica* serotypes, T6SSs belong to five phylogenetically distinct families, named as the *Salmonella* pathogenicity island (SPI) they are encoded on. T6SSs are found in SPI-6 in *S. enterica* Typhimurium, SPI-19 (similar to T6SS-2) in *S. enterica* Dublin, Gallinarum or Enteridis, SPI-20 and SPI-21 in *S. enterica* Arizonae and SPI-22 in *S. bongori* (15, 16) (Fig. 4 and 6). However, SPI-6 T6SS remnant genes are found in serotypes Enteridis and Gallinarum suggesting these genes have been lost during evolution. These T6SS loci are characterized by the presence of non-core genes inserted in between the core elements (17). In *S. enterica* Typhimurium LT2, the organization of the SPI-6 core genes is different from the *E. coli* species but rather shares synteny with distantly strains such as *Burkholderia mallei* and *Ralstonia eutropha* (17). Therefore, *Salmonella* T6SSs derive from non-*E. coli* clusters, and have evolved from the original cluster(s) by the acquisition of non-core modules. These modules might have been transferred between strains as a *hcp-tae4-tai4* module encoding an Hcp protein and an amidase/immunity pair is found in both *S. enterica* Typhimurium and *Enterobacter cloacae* although the core genes differ (18). The addition of these distinct modules during evolution may confer specialized functions to these T6SSs.

A remarkable difference between the T6SSs in *Escherichia* and *Salmonella* strains is the nature of the toxin effectors. T6SS-1-like clusters generally encode effector belonging to phospholipases. T6SS-2 clusters have recombinant hot spots (Rhs) elements bearing putative activities whereas *S. enterica* SPI-6 and *Enterobacter cloacae* T6SS gene clusters encode amidases and Rhs-linked anti-bacterial activities (Table 1, Fig. 5B and 6).

In addition to the main T6SS gene clusters, additional islands encoding Hcp, VgrG, PAAR and putative toxins could be found disseminated on the genome. As Hcp and VgrG have been shown to be carriers for the transport of the effectors, the existence of Hcp/VgrG islands suggests that they correspond to additional modules that adapt to the core machine for the delivery of specific toxins. In several instances, these small islands are inserted within the core gene cluster. As described above, this is particularly visible in the case of the *Salmonella enterica* sp. Typhimurium SPI-6 gene cluster, in which additional islands are inserted within core genes (17, 19) (Fig. 6). It is also very clear when comparing T6SS gene clusters from distinct *E. coli* species, e.g., a *vgrG-tle-tli-paar* fragment is found in the T6SS-1-like T6SS operons in EAEC 042, AIEC LF82 and UPEC UT189, but with differences in the *vgrG* gene and in the effector-immunity pair (Tle effectors of families 1, 3 and 4 respectively) (Fig. 5A and Table 1).

Most of the strains contain several copies of T6SS gene clusters, this multiplicity likely corresponding to various lifestyles, and thus might be reflected by the regulatory mechanisms and the target cells (20). However, little information are available regarding whether these clusters are differently regulated, have distinct functions or similar functions in different conditions. In EAEC 17-2, two T6SSs are encoded within the *pheU* pathogenicity island (21). Both T6SSs have anti-bacterial activities (22) but are expressed in different conditions: while the T6SS-1 cluster is under the control of the Fur repressor and hence induced during iron starvation (23), the T6SS-2 cluster is under the control of AggR, the aggregation master regulator and is expressed in host cells or in synthetic media mimicking the macrophages environment (21). The regulatory mechanisms and function of the *E. coli* and *Salmonella* T6SSs are described in more details in the corresponding paragraphs below.

BIOGENESIS AND ARCHITECTURE OF THE T6SS

The T6SS proteins assemble two modules with different evolution history. As described above, the cytoplasmic syringe-like structure derives from the bacteriophage contractile tail or co-evolved with it from a common ancestor. The assembly of both structures follow a similar sketch : the Hcp tail tube protein polymerizes to form the inner tube, and is tipped by the VgrG membrane-penetrating needle. A sheath-like structure, constituted of the TssBC proteins, polymerizes in an extended, metastable conformation around this inner tube. This two-layered tubular edifice, usually hundreds of nanometers long, is assembled on a platform called the baseplate complex (10). The BC is tethered to the cell envelope via contacts with the second module, the membrane complex (24, 25). This MC is composed of three subunits distributed into the inner and outer membranes. Two of these subunits share sequence homologies with two components of the Type IVb secretion system found in *Legionella pneumophila* or *Coxiella burnetii* (1, 6).

The best characterized T6SS in *E. coli* strains and related species is the EAEC T6SS-1 (Sci-1) machinery. The biogenesis of the T6SS starts with the formation of the membrane complex. The MC serves as a conduit for the passage of the Hcp-VgrG needle during contraction. During T6SS biogenesis, it is the first structure to be assembled. Most cells assemble 1 to 3 MCs that remain static (26). Fluorescence microscopy experiments showed that its localization is not spatially restricted to the cell pole or to the septum but rather that it is randomly distributed in the cell envelope (26). The MC is composed of three proteins (Fig. 7A and 7B): the outer membrane TssJ lipoprotein and the inner membrane TssL and TssM proteins (27-29). The EAEC TssJ lipoprotein is tethered to the outer membrane via acyl chains but faces the periplasm (28). Its tri-dimensional structure has been solved (see Fig. 7C; 30). TssJ has a transthyretin fold, *i.e.*, a β -sandwich of two β -sheets. Two loops, notably that connecting β -strands 1 and 2, mediate contact with the TssM C-terminal domain (26, 30, 31). TssM is a 130-kDa large protein, composed of three trans-membrane segments in its N-terminal third, whereas the 2/3 of the protein extends through the periplasm, from the inner membrane to the outer membrane-anchored TssJ lipoprotein (29, 30). Recently, the crystal structure of the C-terminal portion of the periplasmic region of TssM in complex with TssJ has been solved (Fig. 7C). It is composed of two domains. The structure confirmed that contacts between the two partners are established by interactions of the TssM C-terminal β -domain with loops L1-2 and L3-4 of TssJ (26). Interestingly, this β -domain is followed by an α -helix (colored purple in Fig. 7C) that inserts into the outer membrane, the insertion being facilitated by the TssJ lipoprotein (26). The 300-amino-acid loop located between TM2 and TM3 shares an NTPase fold, although the presence of Walker A and B motifs is not conserved among TssM homologues. TssM contacts TssL, which is composed of a ~ 200-

amino-acid cytoplasmic domain anchored to the inner membrane by a unique C-terminal trans-membrane segment, categorizing TssL as a C-tail protein (32). The proper insertion of this C-tail protein requires the YidC protein and the contribution of the DnaK cytoplasmic general chaperone (32). The structure of the EAEC TssL cytoplasmic domain has been reported (33). It is composed of two bundles of three α -helices, with a general shape resembling a fish hook (Fig. 7D; 33). In several instances, the TssL C-terminus is fused to an additional domain of the OmpA/Pal/MotB family that mediates contact with the peptidoglycan (34). In the T6SS-1 of EAEC and other *E. coli* pathotypes, the TssL protein interacts directly with an additional component, TagL, which carries this motif. *In vivo* and *in vitro* studies have shown that this domain anchors the T6SS to the cell wall, and that mutations preventing TagL interaction with the peptidoglycan abolish T6SS function (27). The structure of the EAEC TssJLM complex has been recently solved at 11.6-Å resolution by negative stain electron microscopy (Fig. 7E; 26). The TssJLM complex has an overall rocket-shape structure with 5-fold symmetry. It is composed of 10 copies of each proteins, and its base is composed of the TssL and TssM cytoplasmic and intra-membrane domains. The TssM periplasmic domains form 10 arches starting from the base and converging to the tip of the structure in two layers of 5 pillars. The assembly of the MC starts from the outer membrane and progresses inward: the TssJ lipoprotein recruits TssM and then TssL is added to the TssJM complex (26). The interior of the membrane complex has a size sufficient to accommodate the VgrG protein. The tip complex closes the structure at the outer membrane and it has been proposed that large conformational changes occur – notably a re-organization of the pillars – to allow the passage of the needle upon sheath contraction (26). Indeed, conformational changes in the periplasmic portion of TssM have been reported in *Agrobacterium tumefaciens* (35).

Once assembled, the MC serves as a docking station for the BC, *i.e.* the assembly platform of the T6SS tail (Fig. 8A and 8B). The T6SS BC is constituted of the TssE, -F, -G, -K and VgrG subunits, which assembles independently of the MC (25). In bacteriophages, the minimal baseplate is composed of six wedges (constituted of the gp6, gp53 and gp25 proteins in bacteriophage T4) that assemble around the spike complex (36). TssE is the T6SS counterpart of the bacteriophage gp25 wedge protein, and the structure of the EAEC TssE can be modeled based on the structures of gp25 homologues (Fig. 8C; 1, 6, 37). This subunit has been suggested to be connected to the sheath. Although we still lack structural information on TssF and TssG, these proteins share limited homologies with the phage T4 gp6 and gp53 proteins, respectively (25). In agreement with phage baseplate structure, a complex comprising TssE, -F, -G and VgrG could be purified from EAEC (25). In addition, it has been shown that TssF, -G and -K form a stable complex in *S. marcescens* (38). Once assembled, the BC is recruited to the MC via multiple contacts including TssK-TssL, TssK-TssM and TssG-TssM interactions (24, 25). The hub of the bacteriophage – and probably of the T6SS – baseplate is the spike complex. The structure of the T6SS spike protein, VgrG, from uropathogenic *E. coli* CFT073 has been reported (Fig. 8D; 39). This membrane-penetrating needle is a trimer with a base that connects to the inner tube, followed by a region composed of repeats that form a highly stable three-stranded β -helix, or β -prism. The structure of the β -prism of the *E. coli* O157 VgrG protein is also known (Fig. 8D; 40). In most cases, an adaptor protein called PAAR interacts at the tip of VgrG and sharpens its extremity (41). The VgrG proteins sits on the Hcp inner tube. Hcp proteins assemble hexameric rings with an internal lumen of ~ 40 Å and an external diameter of ~ 110 Å. The structures of several Hcp proteins from diverse bacteria have been reported in the literature, including that of EAEC (Fig. 8E; 42). The Hcp rings stack on each other on a head-to-tail organization, and this assembly is strictly controlled *in vivo* by additional T6SS subunits (25, 43). The assembly of the inner tube is coordinated with that of the sheath but the rigid tube serves as template for sheath

polymerization (43). The sheath, composed of the TssB and TssC proteins, is the contractile structure that assembles in an extended, metastable conformation that stores the energy required to propulse the inner tube (44). By using time-lapse fluorescence microscopy, it has been shown that the assembly of the 600 nm-long T6SS tube/sheath is completed in ~ 20 seconds and could be maintained in the extended conformation for several hundreds of seconds (22, 44). However, how the extended sheath is stably maintained requires further investigations. The atomic structure of the T6SS sheath in its contracted form has been solved by cryo-electron microscopy (Fig. 8F; 45-47). The sheath is a helical structure composed of 6-TssB/TssC heterodimer strands. Contacts between the heterodimers involve the formation of a 4-stranded β -sheet handshake domain comprising two β -strands from TssC, one from the next TssC on the same strand, and one from TssB from the neighboring strand. This assembly mechanism therefore connects heterodimers of the same strand, as well as with that of the next strand (47). The contraction of the sheath is a fast event that occurs in less than 5 msec (44). Although the propulsion of the inner tube or the delivery of effectors has not been imaged yet, the contraction of the sheath coincides with the lysis of the prey cell (Fig. 9; 22). Once contracted, an N-terminal helix of the TssC subunits protrudes from the structure and recruits the ClpV ATPase that will recycle the sheath subunits (44, 48-50).

FUNCTION AND EFFECTORS

The Type VI secretion system garnered interests due to its ability to target both eukaryotic and prokaryotic cells, therefore delivering toxins with a broad range of activities including nucleases, phospholipases, peptidoglycan hydrolases, NAD(P)⁺ glycohydrolases or cytoskeleton remodeling enzymes (51-55). These toxins are independent polypeptides confined into the Hcp tail tube lumen or displayed at the tip of the VgrG needle via adaptor proteins, or covalently linked as additional C-terminal domains to VgrGs (41, 54, 56-62).

The Type VI secretion system recently emerged as one of the key players during bacterial pathogenesis. However, beside the fact that T6SS expression is usually co-regulated with virulence factors, the role of the T6SS in the infection process could be either direct or indirect. In a few instances, including in pathogenic strains of *E. coli* (see below), the T6SS has been shown to be directly involved in bacterial virulence such as mediating adhesion to host cells or participating to the survival into macrophages or to systemic proliferation, but the molecular details on how the apparatus – or specific secreted proteins – interferes with the host cells have not been defined. In *V. cholerae*, it has been shown that the T6SS disables macrophage by interfering with the dynamics of the cell cytoskeleton. This ability depends on the C-terminal domain carried by the VgrG1 spike, which possesses actin cross-linking activity, therefore curbing actin dynamics, preventing cell movement and inhibiting further phagocytosis of bacterial cells (63-65). The *Aeromonas hydrophila* VgrG1 C-terminal extension has been proposed to carry actin-targeting ADP-ribosyltransferase activity (66). Additional T6SS effectors with anti-host activities include phospholipases in *Pseudomonas aeruginosa*, and toxins that induce membrane fusion (such as the *B. thailandensis* and *B. pseudomallei* VgrG5 C-terminal extensions) or interfere with microtubule dynamics (67-70).

The direct role of the T6SS for virulence towards mammalian models of infection has been challenged by the discovery that the vast majority of T6SSs characterized so far are involved in bacterial growth inhibition. The T6SS is used to deliver anti-bacterial effectors with peptidoglycan hydrolase (amidase (Tae), glycoside hydrolase (Tge)) or phospholipase (Tle) activities into the periplasm of the target prey cell (71-74). These amidases, muramidases and phospholipases belongs to various families that hydrolyse bonds within the peptidic stems (Tae1-4 families) or glucosidic chains (Tge1-3 families) of the peptidoglycan or ester bonds

of phospholipids (Tle1-5 families), respectively (52-54, 72, 74). Toxins with nuclease (Tde) and NAD(P)⁺ glycohydrolase activities have been reported and therefore should reach the cytoplasm for their action (55, 75, 76). How these toxins are transported across the inner membrane is not clearly defined, but it has been proposed that the translational elongation factor Tu (EF-Tu) contributes to the translocation of the NAD(P)⁺ glycohydrolase in *P. aeruginosa* (55). Producing cells are protected from their own effectors or the effectors of their siblings by the co-production of specific protein inhibitors – or immunity proteins – that bind and inhibit the activity of the cognate toxins (52-54). T6SS⁺ bacteria, including closely related species, carries different and unique sets of anti-bacterial effectors, supporting a leading role for T6SS effectors in inter- and intra-bacterial competition and in reshaping bacterial communities (52).

Functions and Effectors in *E. coli*, *Salmonella* and related species

Phenotypes associated with T6SS in *E. coli* and *Salmonella*

Most of the *E. coli* and *Salmonella* T6SSs studied so far have been shown to participate to adherence to biotic and abiotic surfaces, to bacterial competition or to virulence towards corresponding models of infection (Table 1). Taken together, the available information on *E. coli* T6SSs point to a role of T6SS-1 and T6SS-3 for anti-bacterial activity, and T6SS-2 for pathogenesis.

T6SS-dependent bellicose behaviors towards neighboring bacteria have been evidenced for EAEC 17-2 (T6SS-1 and T6SS-3; 22), APEC TW-XM (T6SS-1; 77) and *Citrobacter rodentium* (78). *S. enterica* Typhimurium LT2 has also been recently reported to have antagonistic activities against *E. coli* and *Salmonella* species in a SPI-6 T6SS-dependent manner (19). However, in *S. enterica* Typhimurium, the T6SS is up-regulated in the late stages of infection, once the bacterium is internalized in eukaryotic phagocytic cells whereas no up-regulation could be observed in mixed bacterial cultures (17, 19, 79). These observations suggest that the anti-bacterial activity is activated inside the host. Interestingly, the T6SS is under a regulatory network that also regulates virulence factors including proliferation genes. Because the *S. enterica* Typhimurium population inside phagocytic cells is usually clonal, Brunet et al. proposed that the T6SS may serve to eliminate disabled *S. enterica* Typhimurium cells of the progeny, *i.e.*, the cells unable to produce the T6SS immunity and proliferation genes (19).

A role of the T6SS for biofilm formation has been reported for EAEC 17-2 (T6SS-1; 28) and APEC SEPT362 (T6SS-2; 80). In APEC, the defect in biofilm is accompanied by decreased adherence to epithelial cells (80). However, it is likely that these phenotypes are caused by impacts on fimbriae gene regulation or by perturbations of the biofilm structure due to the anti-bacterial activity. Indeed, deletions of T6SS-2 genes in APEC SEPT362 affect the expression of type 1 fimbriae and flagella, two extracellular structures required for adhesion and biofilm formation (80, 81).

T6SS-dependent pathogenesis towards hosts has been tested for a number of strains (Table 1). From the available data, and in agreement with the over-representation of T6SS-2 in virulent strains, no defect in virulence have been noted for T6SS-1 or T6SS-3 mutants, whereas mutations in T6SS-2 gene clusters usually impact colonization, survival or invasion. The EAEC 17-2 T6SS-1 and T6SS-3 are not required for virulence towards *C. elegans* and intestinal survival within BALB/c mice respectively (21, 28) and mutants in the UPEC CFT073 T6SS-1 gene cluster do not present colonization defects in CBA/J mice bladders and kidneys (82). By contrast to T6SS-1 and T6SS-3, the APEC DE719 and SEPT362 T6SS-2 display attenuated virulence and decreased systemic dissemination in chicks or ducks, and reduced intracellular survival in chicken macrophage cells (12, 80). In APEC strain TW-XM,

T6SS-2 is necessary for cerebral infection and penetration of the blood-brain barrier (11). Similarly, the MNEC K1 T6SS-2 is required for internalization in human brain microvascular endothelial cells (83). In MNEC K1, T6SS-2 carries two Hcp proteins. Analyses of the phenotypes of mutations in these two genes showed that Hcp1 is required for efficient binding to brain endothelial cells whereas Hcp2 induces stress fiber formation, cytoskeleton rearrangements, cytokine and chemokine release and cell apoptosis *via* activation of the caspase 8 pathway (83). Therefore the function of this apparatus is necessary at two different stages of the infection process, probably by the Hcp1- and Hcp2-specific transport of distinct effectors. In *Salmonella* Typhi, the SPI-6 genes are required for systemic infection in a humanized mouse model (84) whereas conflicting data have been reported for the *S. enterica* Typhimurium SPI-6 T6SS regarding replication, survival and proliferation in macrophages or in mice (17, 79, 85, 86). However, the most important effect on replication within phagocytic cells is observed with the *tai4* mutation, a gene that encodes the immunity to the Tae4 amidase, suggesting that this defect is indirect and due to self-intoxication caused by the loss of the immunity (17, 19). The *S. enterica* Gallinarum SPI-19 is required for survival and growth within chicken macrophages and for efficient colonization of the chick gastrointestinal tract and internal organs (87, 88). It is interesting to note that the phenotypes associated with the deletion of the SPI-6 T6SS gene cluster in *S. enterica* Typhimurium can be rescued by the expression of the *S. enterica* Gallinarum SPI-19 T6SS suggesting that both T6SSs perform similar functions despite their phylogenetic differences (89).

E. coli and *Salmonella* T6SS toxins

In *E. coli* and *Salmonella* strains, few effectors have been characterized in details, but their presence and putative activities can be easily predicted. They are organized in tandem with genes encoding small proteins that likely correspond to cognate immunity proteins. Furthermore, they usually co-occur with *hcp*, *vgrG* or *paar* genes.

E. coli T6SS-1 gene clusters encode putative phospholipases (Table 1, Fig. 5A). They localize upstream the *vgrG* genes suggesting that, as shown for *Vibrio cholerae* (90), they use the VgrG needle as carrier for their transport. Interestingly, close inspection of these phospholipase genes suggest they belong to different families : while the AIEC LF82 or UPEC CFT073 T6SS-1 clusters carry putative phospholipases of the Tle3 family, those present on the EAEC 042 and APEC TW-XM genomes are closely related to phospholipases of the Tle1 and Tle4 families respectively (Fig. 5A; 77).

E. coli T6SS-2 gene clusters, as well as the *S. enterica* Gallinarum SPI-19 T6SS-2-like T6SS, usually contain genes encoding Rhs elements (Table 1), but the activity carried by these Rhs proteins can not be inferred from *in silico* prediction algorithms. In addition to the main T6SS gene cluster, *S. enterica* Gallinarum possesses an *hcp* island that encode an Hcp protein, an amidase of the Tae3 family and its cognate Tai3 immunity (15, 73). No gene with putative toxin activity is found within *E. coli* T6SS-3 gene clusters.

The *S. enterica* Arizonae SPI-21 T6SS carries two pairs of S-type pyocins/immunity, as well as a specialized Hcp protein corresponding to a fusion between a traditional Hcp protein to an effector domain (15).

The *S. enterica* Typhimurium SPI-6 T6SS encodes Rhs elements as well as an amidase of the Tae4 family (73) (Table 1, Fig. 6). The structure of Tae4 is available, alone or in complex with its cognate Tai4 immunity protein (91, 92). Tae4 is a DL-endopeptidase with a typical NlpC/P60 domain. It hydrolyses the D-Glu/meso-diaminopimelic acid (mDAP) bond of peptidoglycan peptidic stems. A dimer of Tai4 binds to Tae4 with a K_D of 3×10^{-10} M and inhibits Tae4 activity by inserting the L4 protruding loop of one Tai4 monomer into the Tae4 catalytic pocket (91, 92). Contacts between Tae4 and Tai4 are stabilized by the α -helix of the second Tai4 subunit (91, 92). A similar Tae4/Tai4 pair is encoded within the *Enterobacter*

cloacae SPI-6-like T6SS (73). Indeed, cross-immunity between the *S. enterica* Typhimurium and *E. cloacae* Tae4/Tai4 pairs has been demonstrated (18). Finally, an original mechanism has been revealed in the case of the *S. enterica* Typhimurium Rhs (93). The gene encoding the full-length Rhs protein (Rhs^{main}) is followed by a non-translated region encoding an orphan C-terminal extension (Rhs^{orphan}). However, serial passages in broth or within the mouse induces a duplication of the region and a genetic chromosomal re-arrangement that results to the production of a chimera Rhs protein constituted of the Rhs^{main} core and the Rhs^{orphan} C-terminal extension with anti-bacterial activity. This elegant mechanism therefore provides a selective advantage to cells of the evolved bacterial lineage as it enables to maintain the immunity to Rhs^{main} and to deploy a new toxin that is active against ancestral cells (93). Finally, two genes present within the *Citrobacter freundii* and *E. cloacae* T6SS gene clusters encode proteins with MIX domains, an N-terminal sequence associated with several T6SS toxins (94).

REGULATORY MECHANISMS

T6SS gene clusters are tightly regulated to adapt their expression to the environmental conditions. In agreement with the broad diversity of T6SS targets and activities, T6SS gene clusters are not submitted to a unique regulation but rather have hijacked most of the regulatory mechanisms known in bacteria, including that at the level of transcription or post-transcription (20, 95): two-component systems, transcriptional activators and repressors, histone-like nucleoid associated proteins, quorum sensing, alternative sigma factors, small regulatory RNAs, etc. In addition, a post-translational phosphorylation-dependent pathway has been identified and characterized in Pseudomonads. This signal transmission involves sensing of specific stimuli and activation of a trans-envelope transducing cascade comprising the TagFQRST proteins and leading to the PpkA-dependent phosphorylation of the forkhead-associated FHA protein (96-100). The reversibility of the activation is secured by the dephosphorylation of FHA by the PppA phosphatase (96).

Regulation of T6SS gene clusters in *E. coli*, *Salmonella* and related species

Transcriptional regulation

T6SSs have been studied in details in *Pseudomonas*, *Agrobacterium* and *Vibrio* species. Hence, we have a comprehensive picture of the regulatory mechanisms underlying expression of the T6SS gene clusters in these bacteria. By contrast, except for the entero-aggregative *E. coli* and *S. enterica* Typhimurium T6SSs, only very few is known on the regulation of T6SSs in other *E. coli* and *Salmonella* serotypes.

Enteroaggregative *E. coli*. The EAEC 17-2 strain genome encodes two complete sets of T6SS genes (families T6SS-1 and T6SS-3). The T6SS-1 family cluster (also called *sci-1*) is under the control of the Fur repressor (23, Fig. 10). Fur – for Ferric uptake regulator – is the main regulator that couples iron homeostasis to gene expression (101). In presence of iron, Fur binds to two Fur boxes present in tandem in the promoter sequence of this cluster, overlapping with the transcriptional -10 (Fur⁻¹⁰) and -35 (Fur⁻³⁵), preventing RNA polymerase recruitment and therefore turning OFF the T6SS-1 genes (23). This very simple mechanism is complexified by an epigenetic circuit depending on the action of the Dam adenosine methyltransferase, which couples T6SS expression to the replication state. The Fur⁻¹⁰ box contains a GATC motif that is recognized and methylated by Dam. When Fur is bound to the Fur⁻¹⁰ box, the site is not methylated. However, under low iron conditions, Fur is dislodged, the T6SS-1 genes are turned ON. If cells replicate, the Fur⁻¹⁰ box is methylated after the first

replication, preventing Fur to bind back, therefore turning the T6SS genes under a constitutive ON state. Fur therefore controls the passage between the OFF and ON states whereas Dam is a sensor of replication and controls the passage between the reversible and constitutive ON states (23) (Fig. 10). In agreement with these data, the EAEC T6SS-1 is activated in minimal media or in iron depletion conditions (24). However, this mechanism is unlikely to be conserved between the T6SS-1 clusters shared by pathogenic *E. coli* as no Fur box could be readily identified in their promoter regions.

The EAEC T6SS-3 gene cluster (also called *sci-2* or *aai*) is activated in synthetic media mimicking the macrophage environment such as Eagle's medium (21, 22). DNA microarrays and quantitative RT-PCR have demonstrated that this activation depends on an AraC-like transcriptional regulator called AggR (21, 102). Although no consensus binding site has been defined for AggR, this activator regulates other EAEC virulence factors such as the Aaf fimbriae, the dispersin and the dispersin transporter (102).

No data are available yet for the T6SS-2 gene cluster found in the EAEC 042 strain. Although it remains to be experimentally tested, it has been proposed that regulation of the T6SS-2 *EC042_0229* gene involves the synergistical action of the cyclic AMP receptor protein (CRP) and the nucleoid-associated protein Fis by an original mechanism requiring Fis-dependent compensation of the non-optimal spacing between the CRP and RNA polymerase binding sites (103).

Salmonella enterica. In *S. enterica* Typhimurium LT2, the SPI-6 T6SS gene cluster is controlled by the SsrA/B Two-component system, one of the major regulatory pathways of *Salmonella* virulence (79). The SsrB response regulator binds to and positively regulates most SPI-2 promoters including those controlling expression of the T3SS genes (104). By contrast, SsrA/B exerts a negative control on the SPI-6 T6SS gene cluster (17, 79), probably by direct SsrB binding on distinct promoter regions (105). The expression of T6SS genes encoded within the SPI-6 pathogenicity island are not detected under laboratory *in vitro* conditions (79, 106); however, promoter reporter and transcriptional profiling studies showed that the expression of these genes is activated in the late stages of macrophage and epithelial cells infection (17, 107). In addition to SsrB, the SPI-6 T6SS genes are silenced by the histone-like nucleoid structuring H-NS protein (19, 108, 109). H-NS binds to A/T-rich motifs and polymerizes to spread and silence the genes by preventing access to the RNA polymerase or activators. H-NS is thus a xenogenic silencer that usually represses horizontally-acquired genes and islands (110). Because T6SS genes are clustered in these islands, H-NS is probably involved in the regulation of many T6SS gene clusters in pathogenic *E. coli* but this needs to be addressed.

In *S. enterica* Typhi Ty2, several regulators have been identified such as RcsB, PmrA and Hfq, but their contribution for the activation of the T6SS genes is relatively weak (111). In *Enterobacter cloacae*, several genes of the T6SS cluster are under the control of a LuxR/acylhomoserine lactone-dependent quorum sensing mechanism and therefore respond to the population density (112).

In addition to the EAEC T6SS-3 cluster, shown to be induced in media mimicking the macrophage environment, most of the *E. coli* and *Salmonella* T6SS genes are induced in *in vivo* conditions. This has been reported for the *S. enterica* Typhimurium SPI-6 T6SS genes, which are upregulated during macrophage infection (79, 107) and for the *S. enterica* Gallinarum SPI-19 T6SS genes which are upregulated after internalization by murine or avian macrophages (87). These data suggest that these T6SS gene clusters might play a direct role during infection or that host-mediated activation of the anti-bacterial activity might help to clear the niche and to enable efficient colonization.

Transcriptional frameshifting and post-translational activation

In *Citrobacter rodentium*, the *tssM* gene is subjected to transcriptional frameshifting. The *tssM* gene is interrupted by a premature stop codon, but a poly-A slippery tract located upstream the stop codon induces incorporation of additional adenosines in the RNA and hence the synthesis of TssM length variants (113). Yet, whether the frameshifting efficiency is influenced by environmental cues or by regulatory factors is unknown.

Finally, it is interesting to note that genes encoding the phosphorylation-dependent post-translational pathway found in *Pseudomonas* and *Agrobacterium* species are present in none of the *E. coli*, *Salmonella*, *Citrobacter* or *Enterobacter* strain genomes sequenced so far, except for those encoding FHA proteins which are found associated with *E. coli* T6SS-2-like and *Salmonella* SPI-19 clusters (Fig. 5B and Fig. 6). However, the contribution of FHA for the assembly or the activation of the system has not been reported in these strains.

CLOSING REMARKS AND FUTURE DIRECTIONS

This chapter summarizes the current knowledge on the T6SSs present in *E. coli* and related species. It is clear that the recent years have provided a detailed view on the architecture and mechanism of assembly of this apparatus. However, the regulatory mechanisms underlying the expression of these gene clusters, the effectors delivered by this machinery and the function of the T6SS during host infection remain enigmatic for most enterobacterial pathogens. Further studies will provide a better understanding of the T6SS contribution in the ecological niche of these strains or for pathogenesis. Similarly, although a number of anti-bacterial effectors with amidase, peptidoglycan hydrolase, phospholipase and DNase activities have been identified or predicted, it remains to determine whether phospholipases and DNases might be targeted into eukaryotic host cells and to identify anti-eukaryotic-specific effectors. These toxins would be therefore interesting targets for the development of drugs that will interfere with these toxins, not only for human health, but also in the cases in which the bacterial pathogen targets poultry or cattle.

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902

LEGEND TO FIGURES

Figure 1. Genetic organization and general architecture of the T6SS. (A) Schematic representation of the T6SS core genes. Genes are specified by a letter corresponding to the Tss nomenclature (“A” corresponding to “TssA”) or by their vernacular, usual names (Hcp, VgrG, PAAR and ClpV). The color code is shared with panels B and C. (B) Architecture of the T6SS. The membrane complex, composed of the TssJ lipoprotein (orange) and the TssM (blue) and TssL (red) inner membrane proteins, is indicated (OM, outer membrane; PG, cell wall; IM, inner membrane). The different regions of the tail (spike, tube, sheath & baseplate) are shown. (C) Architecture of a contractile tailed bacteriophage. Components that are shared with the T6SS (spike, tube, sheath & baseplate) are depicted with the same color code (LTF, long tail fibers).

Figure 2. Interbacterial competition between *E. coli* cells. Time-lapse fluorescence microscopy recordings of GFP-labeled EAEC T6SS⁺ cells (green) in presence of mCherry-labeled T6SS⁻ prey bacterial cells (red) in T6SS-3 inducing conditions (one image every 7.5 min). Prey cells that are killed and not present in the next frame are indicated by white arrows. Scale bar is 5 μ m.

Figure 3. Mechanism of action of the T6SS. The biogenesis of the T6SS starts with the assembly of the TssJLM membrane complex (MC) and recruitment of the baseplate complex (BC) (a), which serves as platform for polymerization of the tail tube/sheath structure (b, c). During elongation of the tail structure, effectors (red balls) can be loaded inside the inner tube lumen or attached to the VgrG spike. Following contact with a prey cell, the sheath contracts and propels the inner tube/spike towards the target, allowing penetration and delivery of the effectors (d). Once contracted, the ClpV AAA+ ATPase is recruited to the apparatus for recycling sheath subunits (e, f). The MC (and BC ?) might be re-used for a new round of assembly.

Figure 4. Phylogenetic tree of selected T6SS gene clusters. T6SS gene clusters categorize in 5 phylogenetic groups (A-E) (1, 2). The distribution of the *E. coli*-associated T6SSs (T6SS-1-3, red) and *Salmonella*-associated SPI T6SSs (green) is shown, as well as that of *E. cloacae* and *C. rodentium* (blue) and the model T6SSs from *P. aeruginosa*, *V. cholerae* and *F. tularensis* (black).

Figure 5. Organization of T6SS-1-3 gene clusters. Genes encoding the T6SS-1 (A), T6SS-2 (B) and T6SS-3 (C) in the indicated *E. coli* strains are shown schematically. Homologous genes are colored similarly (see box below). When predictable, putative phospholipase effector/immunity pairs (Tle1/Tli1, Tle3/Tli3 or Tle4/Tli4) or *rhs* genes are indicated. ORFs with unknown function are shown in white. Genes into brackets are not present or not identical in all the strains listed. Genes were identified using the SecReT6 database (114).

Figure 6. Organization of T6SS gene clusters in *Salmonella*, *Enterobacter* and *Citrobacter*. Genes encoding the T6SS in the indicated strains are shown schematically. Homologous genes are colored similarly (see box in Fig. 5). When predictable, *rhs* genes are indicated. The *rhs*^{main} and *rhs*^{orphan} ORF shown to undergoes re-arrangements (93) are indicated in the *S. enterica* Typhimurium SPI-6 gene cluster, as well as the Tae4/Tai4 effector/immunity pairs in *S. enterica* Typhimurium SPI-6 and *E. cloacae*. ORFs with unknown function are shown in white. Genes were identified using the SecReT6 database (114). Note that the transcription of the *C. rodentium* *tssM* gene, interrupted by an early stop codon, is rescued by frameshifting (113).

Figure 7. Architecture and structure of the T6SS membrane complex. (A) The *tssJ*, *tssL* and *tssM* genes which encode the components of the membrane complex. (B) Schematic representation of the TssJ, -L and -M proteins: TssJ is an outer membrane (OM)-tethered lipoprotein whereas TssL and TssM are inner membrane (IM)-embedded proteins. In T6SS-1, the membrane complex comprises an additional protein, TagL, which binds to the peptidoglycan (PG) layer (not depicted here) (27). (C) Crystal structure of the complex between the soluble fragment of TssJ (orange) and the two C-terminal domains of the TssM periplasmic segment (light and dark blue) including the C-terminal helix that

inserts into the outer membrane (in purple) from EAEC T6SS-1 (PDB: 4Y7O) (Reprinted from reference 26 with permissions). (D) Crystal structure of the cytoplasmic domain of TssL from EAEC T6SS-1 (PDB: 3U66) (33). (E) Negative stain electron microscopy structure reconstruction of the EAEC TssJLM complex (lower panel, EMDB: 2927) (adapted from reference 26 with permissions) (scale bar is 50 nm). The position of the outer (OM) and inner (IM) membranes are predicted based on the presence of detergent micelle and the putative location of the trans-membrane segments of TssM, respectively. In the upper panel is shown a top view of the TssJLM complex in which crystal structures of the TssJ-M complex (panel C) are docked, highlighting the presence of two concentric layers closing the channel at the outer membrane.

Figure 8. Architecture and structure of the T6SS tail complex. (A) The *tssA*, *tssB*, *tssC*, *tssE*, *tssF*, *tssG*, *tssK*, *hcp*, *vgrG* and *paar* genes which encode the components of the tail complex (blue, sheath subunits; black, inner tube subunit; green, spike subunits; pink, putative baseplate subunits). (B) Schematic representation of the T6SS tail complex (same color code as panel A). (C) Structural model of EAEC T6SS-1 TssE based on the bacteriophage gp25 crystal structure (PDB: 4HRZ). (D) Composite structure made with the crystal structures (from bottom to top) of the UPEC CTF073 VgrG1 protein (PDB : 2P57) (39), the *E. coli* O157 EDL933 β -helical prism (PDB: 3WIT) (40) and the *E. coli* O6 PAAR protein (PDB: 4JIW) (41). (E) Crystal structure of the EAEC T6SS-1 Hcp hexamer (left, top view; right, side view) (PDB: 4HKH) (42). (F) Cryo-electron micrograph of a contracted T6SS sheath from *V. cholerae* (left panel, scale bar is 100 nm) and atomic-resolution cryo-electron structure of the TssB-C complex (PDB: 3J9G) (47) (adapted from reference 47 with permissions).

Figure 9. T6SS sheath contraction coincides with target cell lysis. Time-lapse fluorescence microscopy recordings of EAEC producing fluorescently-labeled sheath subunits (TssB-sfGFP) in presence of mCherry-labeled T6SS⁺ *E. coli* K-12 prey cells (one image every 7.5 min). The time-lapse highlights the assembly and the contraction (white arrow) of the T6SS sheath, followed by the lysis of the target cell. Scale bar is 1 μ m. Adapted from reference 22 with permissions.

Figure 10. Regulation of the EAEC T6SS-1 gene cluster. (A) Schematic representation of the promoter organization of the EAEC T6SS-1 gene cluster. The location of the -10 and -35 transcriptional elements (blue), of the Fur-binding sequences (red) and of one of the GATC site (green) are shown. (B) Regulatory mechanism of the EAEC T6SS-1 gene cluster (23). In iron-replete conditions, a Fur dimer (red balls) represses the expression of the T6SS-1 gene cluster by binding to the Fur⁻¹⁰ box, which overlaps with the -10 element (OFF). When iron is limiting, the -10 element is available for the RNA polymerase allowing expression of the T6SS-1 genes (ON). Upon replication, the GATC site is methylated (CH₃) and by preventing Fur binding allows Fur-independent, constitutive expression of the T6SS-1 gene cluster.

T6SS	pathotype/ serotype	strain	activity	effector	references
T6SS-1	EAEC	17-2	biofilm	-	28
			anti-bacterial	Tle1 (putative)	-
	APEC	TW-XM	biofilm	-	77
			anti-bacterial	Tle4	77
T6SS-2	APEC	DE719	attenuated virulence in ducks, reduced intracellular in chicken macrophages	-	12
		SEPT362	attenuated virulence in chicks	-	80
		TW-XM	penetration of the blood-brain barrier during cerebral infection	-	77
	MNEC	K1	invasion of human brain microvascular endothelial cells	-	83
T6SS-3	EAEC	17-2	anti-bacterial	-	22
SPI-6	<i>S. Typhimurium</i>	LT2	affect replication in macrophage and systemic dissemination in mice and in chicks	-	17, 79, 85, 86, 89
			anti-bacterial	Tae4	19, 92
			anti-bacterial	Rhs ^{orphan}	93
	<i>S. Typhi</i>	Ty2	systemic infection in mice	-	84
	<i>E. cloacae</i>	ATCC13047	anti-bacterial (putative)	Tae4	18, 73
SPI-19	<i>S. Gallinarum</i>	287/91	colonization of the gastro-intestinal tract and systemic dissemination in chicks	-	88
CTS1	<i>C. rodentium</i>	IC68	anti-bacterial	-	78

Table 1. Phenotypes and effectors associated with T6SS in *E. coli*, *Salmonella* and related species.

A T6SS-1

B T6SS-2

C T6SS-3

A **B** **C** **TssE****D** **VgrG****E** **F**

100 nm

TssBC

assembly

contraction

prey lysis

A

B