



Manipulation of host membranes by the bacterial pathogens *Listeria*, *Francisella*, *Shigella* and *Yersinia*.

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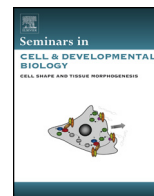
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Review

Manipulation of host membranes by the bacterial pathogens *Listeria*, *Francisella*, *Shigella* and *Yersinia*



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ABSTRACT

Bacterial pathogens display an impressive arsenal of molecular mechanisms that allow survival in diverse host niches. Subversion of plasma membrane and cytoskeletal functions are common themes associated to infection by both extracellular and intracellular pathogens. Moreover, intracellular pathogens modify the structure/stability of their membrane-bound compartments and escape degradation from phagocytic or autophagic pathways. Here, we review the manipulation of host membranes by *Listeria monocytogenes*, *Francisella tularensis*, *Shigella flexneri* and *Yersinia* spp. These four bacterial model pathogens exemplify generalized strategies as well as specific features observed during bacterial infection processes.

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Abbreviations: ECM, extracellular matrix; FPI, *Francisella* pathogenicity island; LLO, listeriolysin O; OMP, outer membrane protein; PMN, polymorphonuclear neutrophils; ROS, reactive oxygen species; Subsp, subspecies; T3SS, type 3 secretion system; TLR, toll-like receptor; YCV, *Yersinia*-containing vacuole.

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1. Introduction

The eukaryotic cell is a complex environment. Vital functions such as internalization of external solutes, synthesis of novel proteins or genetic information storage are compartmentalized in membrane-bound structures. From the perspective of a pathogen, the eukaryotic cell might be considered as a rich source of nutrients as well as a protected environment in which microbial replication may take place, avoiding contact with extracellular host defenses such as the complement or antibodies. Cells have adapted molecular strategies (e.g. phagocytosis or autophagy) to counteract and destroy intracellular pathogens, which in turn have developed strategies to cope with host cell defenses.

Four major bacterial pathogenesis paradigms are *Listeria monocytogenes*, *Francisella tularensis*, *Shigella flexneri* and *Yersinia* spp. *Listeria monocytogenes* is responsible for a food-borne disease associated with meningitis and abortions. *Francisella tularensis* is the agent of tularemia, responsible for ulcero-glandular and pneumonic severe infections. *Shigella flexneri* is the etiological agent of human bacillary dysentery. *Yersinia pseudotuberculosis* and *Y. enterocolitica* are responsible for enteritis, ileitis, and mesenteric lymphadenitis in humans, and are closely related to *Y. pestis*, the plague agent. Besides their clinical importance, these bacteria are useful paradigms to understand the diverse strategies used by microbial pathogens to subvert host cell functions. While *Listeria* and *Yersinia* invade cultured epithelial cells by interacting with host cell plasma membrane receptors via bacterial surface proteins, *Shigella* injects bacterial effectors within the host cell cytoplasm to promote internalization and *Francisella* induces spacious pseudopods to invade macrophages. Once inside epithelial cells, *Yersinia* proliferates in a membrane-bound compartment while *Listeria*, *Shigella* and *Francisella* disrupt their internalization vacuole and escape into the host cell cytoplasm. *In vivo*, *Yersinia* is mainly extracellular. We will review here the major interactions that occur between these important bacterial pathogens and host membranes.

2. *Listeria monocytogenes*

Listeria monocytogenes is a Gram-positive bacterial pathogen responsible for listeriosis, a disease causing meningitis in the new born and abortions in pregnant women. *Listeria* is now a classical model in the study of bacterial intracellular parasitism [1]. The pathogenic potential of *Listeria* is intimately associated with its capacity to traverse the intestinal, the foeto/placental and the blood/brain barriers, which requires invasion of host cells [2]. Using

tissue cultured cells, it has been observed that upon internalization *Listeria* is able to disrupt its membrane-bound compartment and to escape to the host cytoplasm, where it proliferates [3]. *Listeria* is then able to avoid the autophagosomal machinery by polymerizing host cell actin, which drives bacterial cytoplasmic movement and cell-to-cell spread [4] (Fig. 1). Exceptions to this canonical intracellular cycle model have been observed *in vivo*, particularly during the traversal of the intestinal barrier. Indeed, in goblet cells *Listeria* does not escape from the internalization vacuole and is transcytosed to the lamina propria [5]. It may also persist in spacious phagosomes [6].

2.1. Cell invasion mediated by InlA and InlB

Listeria induces its internalization within mammalian cells by using surface invasion molecules that activate host plasma membrane receptors. These proteins belong to the internalin family, whose members are characterized by the presence of N-terminal leucine-rich repeats which drive protein–protein interactions [7]. The prototype internalin (InlA) is a covalently-anchored bacterial cell wall protein that binds the cellular adherens junction molecule E-cadherin [8] and triggers *Listeria* internalization in polarized epithelial cells and tissues, particularly during traversal of the intestinal and the foeto/placental barriers [9,10]. InlB is a second bacterial surface molecule loosely attached to lipoteichoic acids which can interact with the host molecule Met [11], the receptor for the hepatocyte growth factor. *In vitro*, InlB promotes *Listeria* internalization in a wide variety of epithelial cells; *in vivo*, InlB has been shown to cooperate with InlA in crossing the foeto/placental barrier [2] (Fig. 1A).

2.1.1. InlA-invasion pathway

Lipid rafts are cholesterol-rich signaling platforms that allow the clustering of signaling proteins or lipids, and which play a central role in the infectious process of many bacterial pathogens. Interaction between InlA and E-cadherin leads to E-cadherin clustering, a process which is alleviated if lipid rafts are disorganized [12]. This event is followed by E-cadherin cytoplasmic tail phosphorylation and ubiquitylation by Src and Hakai, respectively [13], allowing clathrin recruitment via its adaptor Dab2 [14]. Clathrin is a major membrane coat implicated in the endocytosis of many surface receptors, and clathrin-mediated endocytosis is involved in the infectious process of several important viruses [15]. The study of *Listeria* internalization demonstrated that during bacterial entry, clathrin does not function as a classical endocytic coat but instead

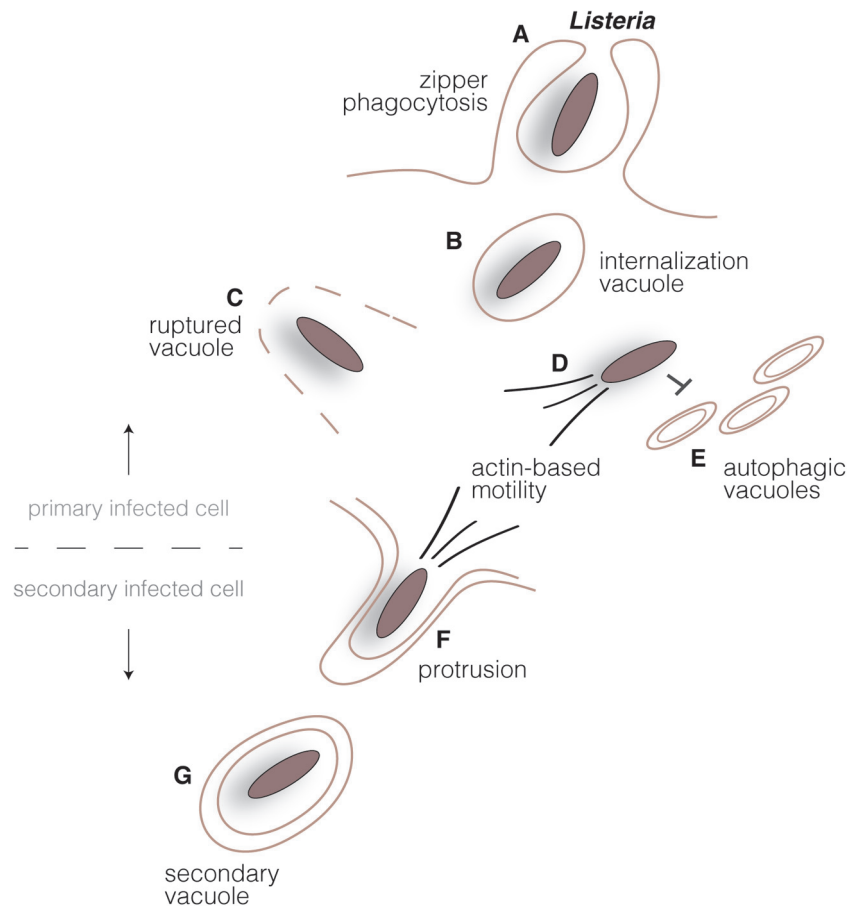


Fig. 1. *Listeria* intracellular cycle, *Listeria* surface invasion proteins interact with host cell receptors in epithelial cells triggering bacterial engulfment (A). *Listeria* transiently resides in an internalization vacuole (B), but bacterial secreted toxins lyse the vacuole (C) and bacteria are released in the host cell cytoplasm. Cytoplasmic bacteria multiply and polymerize host cell actin, promoting an actin-based motility system (D) which propels the bacteria and allows escape from the autophagic pathway (E). Upon reaching the host cell plasma membrane, *Listeria* induces the formation of protrusions (F) and are found in double-membrane vacuoles in secondary cells (G). Disruption of this compartment by bacterial secreted toxins allows the start of new infection cycle.

as a molecular hub that triggers actin reorganization at bacterial entry sites [16–18]. Indeed, clathrin coats are stabilized by phosphorylation of the clathrin heavy chain, followed by the sequential recruitment of Hip1R, actin and myosin VI [14]. PI 3-kinase activity, constitutively present in intestinal goblet cells, is also required for the InlA-dependent bacterial crossing of the intestinal barrier following the InlA/E-cadherin interaction [19] (see below for discussion on phosphoinositide metabolism).

2.1.2. InlB-invasion pathway

The second *Listeria* invasion protein InlB activates the receptor tyrosine kinase Met and triggers recruitment of several protein adaptors [20,21]. Cbl is a ubiquitin ligase which post-translationally modifies Met, favoring clathrin recruitment and actin reorganization during cell entry [14,22,23]. Cbl, together with the protein adaptors Shc, Gab-1 and CrkII, also contributes to the infection process by recruiting PI 3-kinase to the *Listeria* entry foci [24]. PI 3-kinase is a central enzyme in many signaling cascades which locally produces the phosphoinositide PI(3,4,5)P₃ and promotes further recruitment of cellular effectors that link receptor activation with actin cytoskeleton rearrangements. In the InlB-invasion pathway, cholesterol-rich lipid rafts are not required for the initial Met clustering but instead, for optimal PI(3,4,5)P₃ distribution within the inner leaflet of the plasma membrane, and for subsequent Rac1 recruitment [12,25]. Rac1 is a member of the Rho family of small GTPases which modulates cortical actin polymerization during *Listeria* entry by activating WASP-related complexes WAVE-1/-2 [26].

The Arp2/3 complex is the final component of this signaling cascade which directly promotes nucleation of actin polymerization at bacterial entry foci [27]. The Arp2/3 complex has been traditionally considered as a single molecular entity composed of 7 subunits: ARPC1/p41, ARPC2/p34, ARPC3/p21, ARPC4/p20 and ARPC5/p16 together with the Arp2 and Arp3 subunits [28]. A genome-wide siRNA screen recently determined that the subunits ARPC4/p20 and ARPC5/p16 are not strictly required to drive *Listeria* internalization within host cells, revealing the presence of alternative Arp2/3 complexes in mammalian cells [29]. Another cytoskeletal element implicated in *Listeria* infection are septins, a family of small GTPases which form non-polar filaments [30]. The septin cytoskeleton contributes to the anchorage of Met to the actin cytoskeleton [31] and different members of the septin family have been shown to modulate host cell invasion by *Listeria* [32–34].

As mentioned above, PI 3-kinase activity is constitutive in intestinal cells but not in the placenta, and therefore InlB plays a critical role in the traversal of the foeto/placental barrier by promoting the PI 3-kinase-dependent production of PI(3,4,5)P₃ [19]. Additionally, in epithelial cells PI 4-kinase activity is necessary for efficient *Listeria* internalization [35–37]. PI4P production takes place downstream of the tetraspanin CD81 [38], a plasma membrane molecular hub which interacts with integrins and has been also implicated in cellular entry of *Plasmodium falciparum* [39] and the hepatitis C virus [40]. *In vivo*, it has been recently demonstrated that CD81-associated signaling inhibits T-cell anti-*Listeria* responses [41].

Host molecules may also counterbalance plasma membrane signaling triggered by *Listeria* and down-regulate infection: for example, the host phosphatidylinositol 5-phosphatase Oculocerebrorenal Syndrome of Lowe Protein (OCRL) cleaves PI(4,5)P₂ and PI(3,4,5)P₃, inhibiting actin polymerization and therefore bacterial entry [42].

2.2. Phagosomal membrane disruption by LLO, PlcA and PlcB

Upon cellular invasion, *Listeria* is located in a phagosomal compartment (Fig. 1B) that is disrupted via the lytic action of the pore-forming toxin listeriolysin O (LLO), a member of the cholesterol-dependent cytolysin (CDC) family. CDCs can mediate hemolysis in their reduced state, and the cellular enzyme γ -interferon-inducible lysosomal thiol reductase (GILT) present on lysosomes contributes to activate LLO within phagosomes [43]. An acidic pH also potentiates LLO lytic activity [44]. Insertion of LLO in the *Listeria* phagosome allows Ca²⁺ leakage which inhibits phagosomal maturation and blocks interaction with lysosomal compartments [45,46]. Rupture of the phagosomal membrane is enhanced by the two bacterial phospholipases PlcA and PlcB which display phosphatidylinositol-specific and broad-range phospholipase C activities, respectively [47] (Fig. 1C). It is important to mention that besides its critical function in disrupting phagosomal compartments, LLO released by extracellular *Listeria* also contributes to infection by inducing Ca²⁺ influx [48] that promotes bacterial internalization [49], while triggering the targeting of late endosomal compartments to the bacterial entry sites (Kühbacher, Cossart and Pizarro-Cerdá, unpublished observations) as has been observed for the internalization process of parasites like *Leishmania* or *Trypanosoma* [50,51]. LLO also favors *Listeria* infection by influencing host protein sumoylation [52], histone post-translational modifications [53] and mitochondrial dynamics [54].

2.3. Autophagy escape mediated by ActA and InlK

After vacuolar escape, cytosolic *Listeria* replicate and polymerize host actin via the recruitment to bacterial poles of the Arp2/3 complex by the *Listeria* surface protein ActA [55,56]. ActA mimics WASP and activates actin filament barbed-end branching by the Arp2/3 complex [57]. Similar to what has been mentioned above for bacterial entry, it has been recently found that the Arp2/3 subunit ARPC5/p16 is not required for the *Listeria* actin-based motility, suggesting that other cellular molecules may associate with the Arp2/3 complex to modulate its function and/or localization [29]. A similar phenotype has been recently observed for the motility of the Vaccinia virus [58], indicating that the diversity of Arp2/3 complexes is more widespread than initially appreciated. The actin comet tails formed by *Listeria* contain more than 20 different molecules which stabilize the actin filaments or modulate force generation [59] (Fig. 1D). Actin-based motility is important not only to favor invasion of host neighboring cells (see below) but it is also critical to escape from the autophagy machinery [60–62] (Fig. 1E). Indeed, the autophagic pathway has been shown to limit *Listeria* cytoplasmic growth [63]. Besides ActA, the surface molecule InlK also protects *Listeria* from autophagic degradation by recruiting to the bacterial surface the major vault protein and therefore inhibiting autophagic recognition [64]. In addition, it has been recently shown that the ubiquitin-like protein ISG15 is induced upon *Listeria* infection, modifies a subset of host proteins involved in cytokine secretion and restricts *Listeria* proliferation both *in vitro* and *in vivo*. Interestingly, ISG15 leads to increased cytokine secretion following over-expression or infection, a phenomenon that could explain the *in vivo* phenotype. How ISG15 leads to fewer bacteria *in vitro* is still an open question [65].

2.4. Cell-to-cell spread

Initial studies suggested that actin-based motility was sufficient to induce the formation of host membrane protrusions and uptake by neighboring cells during *Listeria* cell-to-cell spread [66–68] (Fig. 1F). Attachment of actin tails to plasma membrane by the phosphorylated form of the actin-binding molecule ezrin has been also proposed to promote the formation of *Listeria* protrusions [69]. Interestingly, formins seem to catalyze the generation of unbranched actin filaments that assist the formation of *Listeria* protrusions initially triggered by the propulsive force generated by the Arp2/3 complex [70]. A siRNA screen has revealed a role for the CNSK1A1 kinase in actin-tail formation and for the CNSK2B in favoring cell-to-cell spread, but their precise mechanism of action remains to be identified [71].

Interestingly, the secreted molecule InlC has been shown to contribute to cell-to-cell spread by directly interacting with the host cell molecule Tuba [72]. Tuba normally controls the structure of cortical actin structures by recruiting N-WASP and by interacting with COPII components: by sequestering Tuba, InlC induces loose membrane junctions that are more permissive to *Listeria* protrusion formation [72,73]. More recently, it has been proposed that *Listeria* cell-to-cell spread exploits efferocytosis, which is a process by which dying or dead cells are removed by phagocytosis [74]. *Listeria* uptake in secondary-infected cells leads to bacterial sequestration within a double-membrane compartment (Fig. 1G). The phospholipases PlcA and PlcB are required for disruption of the inner membrane while LLO is involved in lysis of the outer membrane, allowing initiation of a new *Listeria* intracellular cycle [75].

3. *Francisella tularensis*

Francisella tularensis is a Gram-negative bacterium and the causative agent of the zoonotic disease tularemia. This highly infectious bacterial pathogen can infect a broad variety of animal species, from mammals to arthropods. Tularemia can be transmitted to humans in numerous ways [76] and the clinical outcome of the disease depends on the route of infection and bacterial subspecies (subsp) [77]. Four major *F. tularensis* subsps exist that differ in virulence and geographic distribution. They are designated subsps *tularensis*, *holarctica*, *mediasiatica* and *novicida*. The most virulent subsp *tularensis*, capable of causing a life-threatening disease in humans, is considered a potential bioterrorism agent [78] by the U.S. Centers for Disease Control and Prevention (CDC). *Francisella* is an intracellular pathogen which upon adhesion and invasion of host cells resides transiently in a phagosomal compartment; disruption of this phagosome is followed by active cytosolic multiplication. Dissemination to adjacent cells does not involve actin polymerization and relies instead in cell lysis and bacterial dissemination.

3.1. Adhesion and invasion

Francisella is able to infect and multiply *in vitro* within numerous cell types including hepatocytes, endothelial cells and fibroblasts, to which bacterial attachment is mediated by subsps-specific outer membrane proteins and the type IV pilus [79]. However, *in vivo* the *Francisella* main cellular niche is the macrophage, used for bacterial dissemination [80]. Different macrophage receptors involved in *Francisella* uptake have been identified over the past ten years, including the complement receptor CR3 (CD11b/CD18), the mannose receptor (MR), the scavenger receptor A (SRA), Fc γ Rs and surface-exposed nucleolin [81]. The subversion of phagocytic or scavenger receptors to invade macrophages is not specific to *Francisella* and is common to a subset of bacterial intracellular

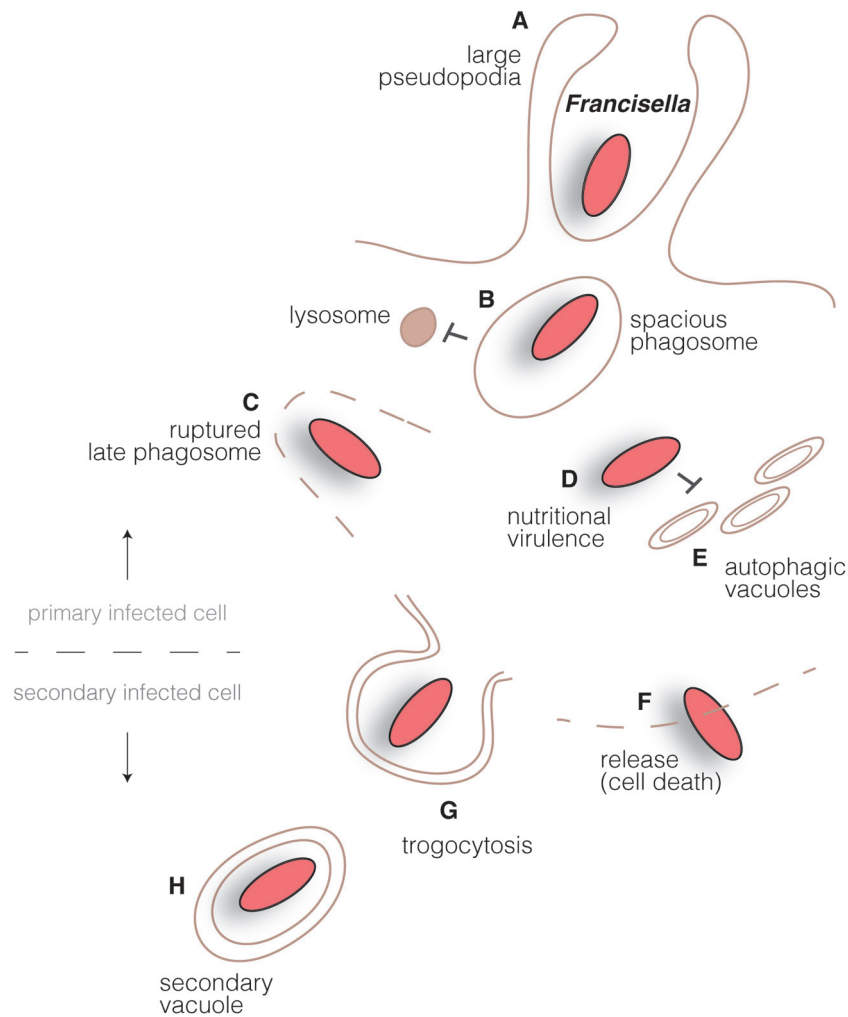


Fig. 2. *Francisella* intracellular cycle. *Francisella* interact with innate immune receptors at the surface of macrophages in order to get internalized by large pseudopodia (A). Inside cells, bacteria reside in a large spacious phagosome that matures into a late phagocytic compartment but avoids fusion with lysosomes (B). A type VI secretion system is involved in the rupture of the bacteria-containing compartment (C) and *Francisella* is found in the host cell cytoplasm, where it takes advantage of available resources ('nutritional virulence') (D) and avoids interaction with autophagic vacuoles (E). Bacteria can be released from dying cells (F) or can be trapped by 'trogocytosis' in double membrane compartments that allows infection of neighboring cells (H).

pathogens including *Mycobacterium tuberculosis* [82], which do not actively promote surface cytoskeletal rearrangements and instead rely on the internalization machineries of host cells. As observed for the vast majority of bacterial intracellular pathogens, *Francisella* internalization is dependent on actin polymerization; remarkably, *Francisella* is captured by macrophages via a unique engulfment mechanism which involves the initial formation of large loops or pseudopodia [83] (Fig. 2A). As in the case of *Listeria*, lipid rafts are required for efficient *Francisella* internalization [84]. In order to avoid cellular recognition by the host immune system, *Francisella* harbors a peculiar lipopolysaccharide (LPS) which does not activate host Toll-like receptor 4 (TLR4); nevertheless, the bacterial lipoproteins (BLPs) do activate TLR2, the primary TLR involved in the inflammatory response to *Francisella* infection.

3.2. Phagosomal stage

3.2.1. Spacious phagosome maturation

After engulfment by phagocytic cells, *Francisella* transiently resides in a spacious phagosome (Fig. 2B). This bacterial-containing compartment sequentially displays membrane markers of early (EEA1) and late endosomes/lysosomes (LAMP-1 and -2) but fails to acquire the hydrolase cathepsin D or lysosomal tracers [85].

Depending on the uptake route, the phagosomal stage may last 30 min for non-opsonized bacteria to 2–4 h for opsonized *Francisella*. While the importance of phagosomal acidification has been shown to be an absolute prerequisite for *Listeria* to escape from the phagosome as mentioned above [44], controversy exists on the importance of this process on the ability of *Francisella* to access to the host cytosol. Indeed, some studies indicate that compartments containing *Francisella* become acidified prior to phagosomal disruption [86,87] while others report that *Francisella* phagosomes resist acidification and acquire limited amounts of vacuolar ATPases, while still capable of accessing the cellular cytoplasm [88].

3.2.2. Inhibition of NADPH oxidase

During this transient time spent in the phagosome, *Francisella* must actively evade several host antimicrobial defenses, including anti-microbial peptides and reactive oxygen species (ROS) produced by the NADPH oxidase. *Francisella* is precisely one of the few bacteria capable of surviving within human polymorphonuclear neutrophils (PMNs), which are potent ROS producers, due to its ability to block either the assembly of the NADPH oxidase integral membrane gp91^{phox}/p22^{phox} components, or the phosphorylation and recruitment of the its cytosolic p47^{phox}/p40^{phox} subunits [89]. *Francisella*-infected monocytes or macrophages also

fail to trigger a robust oxidative burst [90]. *Francisella* is additionally equipped with a series of enzymes that include superoxide dismutase, catalase and acid phosphatases that allow bacterial survival in the hostile phagosomal environment [91].

3.2.3. Phagosomal disruption by a T6SS

Bacterial secretion systems are sophisticated nano-machines that allow protein export from the bacterial cytosol toward specific external environments. Type VI secretion systems (T6SS) have been recently discovered in Gram-negative bacteria and are critical to the virulence of many pathogens such as *Vibrio cholerae* and *Pseudomonas aeruginosa*. A *Francisella* pathogenicity island (FPI) has been presumed to encode a T6SS apparatus, and many genes expressed from this locus or regulating expression of this locus, including *IglC* and *mgIA* [92], *iglI* and *iglJ* [89] or *fevR* [93], were previously implicated in promoting *Francisella* escape from its phagosome and bacterial cytoplasmic proliferation (Fig. 2C). The atomic structure of the *Francisella* T6SS was recently solved by cryo-electron microscopy, demonstrating for the first time in this type of system how its two main components, *IglA* and *IglB*, are interdigitated into a single fold [94]. Moreover, it is the first T6SS which displays assembly upon KCl stimulus or intracellular residence, and its contribution to phagosomal escape and intra-macrophage replication has been confirmed [94]. However, the precise molecular contribution of the *Francisella* T6SS effectors to phagosomal membrane disruption, together with additional not FPI-encoded proteins also implicated in this process, remain to be identified.

3.3. Cytoplasmic life and interactions with the autophagic pathway

Once in the cytosol, *Francisella* adapts its nutritional needs to the available nutrient sources, a function that has been recently coined as ‘nutritional virulence’ [95] (Fig. 2D). Asparagine uptake has been identified as critical for efficient bacterial cytosolic multiplication [96]. It has been proposed that in mouse embryonic fibroblasts (MEFs), cytoplasmic *Francisella* stimulates an ATG5-independent autophagy process which further supply nutrients that supports bacterial growth [97]. An early report by Celli et al. indicated that, at late stages of infection in bone-marrow derived macrophages, a subset of intracellular *Francisella* could be found inside LC3-positive, double membrane-bound vacuoles [98]. This observation suggested that late residency within autophagic compartments constitutes an intrinsic part of the *Francisella* intracellular life cycle [98]. However, further studies in other cell types contradicted this notion by demonstrating that metabolically active *Francisella* in fact avoid autophagic capture [85] (Fig. 2E) whereas mutant bacteria displaying impaired cytosolic multiplication are captured and destroyed by autophagy. In addition, guanylate-binding proteins (GBPs) represent another cell system to attack cytosolic *Francisella* in order to activate the Absent in Melanoma 2 (AIM2) inflammasome, which controls *Francisella* replication [99].

3.4. Cell-to-cell spread

Unlike *Listeria* (mentioned above) or *Shigella* (see below), *Francisella* is not able of actin-based cytosolic movements. Hence, *Francisella* dissemination to adjacent cells was thought to occur exclusively after their release from lysed infected cells following apoptotic or pyroptotic cell death (Fig. 2F). A novel cell-to-cell dissemination mechanism was very recently described by which *Francisella* can also infect adjacent cells without passing through an extracellular stage: this unique process, designated trogocytosis, allows bacterial transfer from infected cells to uninfected macrophages through a transient, contact-dependent mechanism, leaving both donor and recipient cells intact and viable [100]

(Fig. 2G). Therefore, *Francisella* is able to exploit natural host cell cytosolic exchange mechanisms to directly transfer from infected to non-infected cells.

4. *Shigella flexneri*

The Gram-negative pathogenic bacteria *Shigella* spp. cause bacillary dysentery in humans, also known as shigellosis. *Shigella* transmission takes place generally through the fecal-oral route by the uptake of contaminated food or water, giving rise to a disease that results in colonic epithelium destruction and elevated intestinal inflammation. Annually, about 160 million cases of shigellosis with about 1 million deaths have been estimated to occur mainly in the developing world; hence, *Shigella* infections constitute an important health burden, particularly for young children [101]. The rise in antibiotic resistance and the absence of an efficient vaccine against *Shigella* underline the urgency for the development of novel antibacterial strategies against this important pathogen [102].

Four *Shigella* species have been described: *S. dysenteriae*, *S. flexneri*, *S. boydii* and *S. sonnei*. The molecular basis of infection has been studied using the *Shigella flexneri* model (referred from now on as *Shigella*). Primordial for *Shigella* infectivity is the presence of a large virulence plasmid that encompasses the mxi-spa pathogenicity island, encoding the structural genes for a type 3 secretion system (T3SS) and about 25 effector proteins that are injected into targeted host cells upon contact with the bacterium [103,104]. The T3SS is a major virulence determinant that has been described in many pathogenic bacteria including *E. coli*, *Salmonella* and *Yersinia* (see below). The *Shigella* T3SS is expressed and assembled at 37 °C together with a first wave of effector proteins, however secretion together with the expression of a second wave of effectors is only triggered upon contact with the host cell. The effectors are translocated directly from the bacterial cytoplasm into the host cytoplasm through the needle complex that connects with the bacterial translocon, made of the two proteins IpaB and IpaC spanning the host cell membrane. Subsequently, the translocated effector proteins subvert host cellular processes resulting in cytoskeletal rearrangements, bacterial invasion and the modulation of the host immune response. Injection of the bacterial effectors is a rapid process with half the pre-stored proteins translocated within 240 s [104,105].

4.1. T3SS effector injection triggers *Shigella* internalization

The cellular mechanisms that take place during the early contact between *Shigella* and the host cells have been investigated in detail over the last two decades. Recent data have highlighted the capture of the pathogen in the vicinity of host cells through cellular extensions that resemble filopodial-like structures involving a functional T3SS. The interactions between the pathogen and the filopodial tips induce their retraction to give rise to close contacts between the bacterium and the cellular surface [106]. Additionally, the bacterial translocator/effector IpaB, the bacterial factor IcsA and the host receptors CD44 and $\alpha 5 \beta 1$ -integrin have been proposed in the establishment of early bacterial-host contacts [107–109]. Also, the presence of cholesterol in lipid-raft-like domains is required for proper translocon assembly and bacterial internalization [110,111]. Together, these interactions promote *Shigella* entry through a site-specific invasion process [112].

The translocon complex assembles within the host membrane and is connected with the T3SS, to allow the injection of the bacterial effectors into the host cytoplasm or in association with the host membrane complex. These effectors include IpaA, IpaB, IpaC, IpgB1, IpgB2, IpgD and VirA, all involved in the entry of *Shigella* through a trigger-like process [113] (Fig. 3A). IpaA has been found to inter-

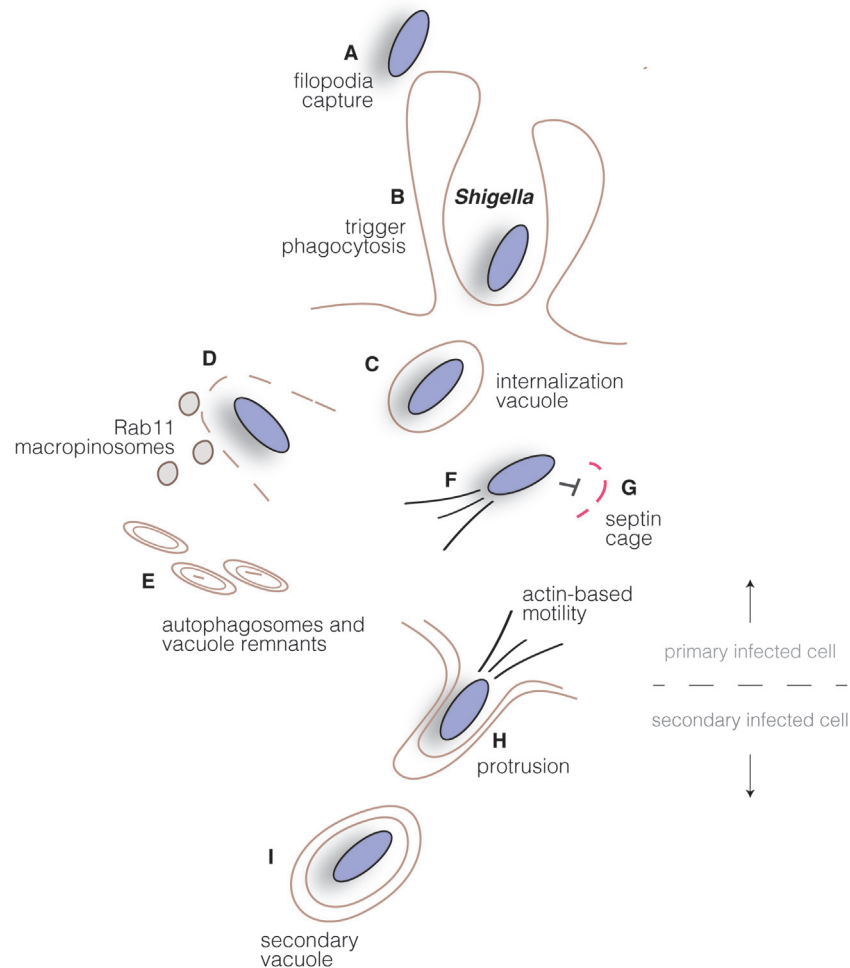


Fig. 3. *Shigella* intracellular cycle. *Shigella* promotes its uptake into host cells by triggering plasma membrane rearrangements through injection of type III secretion system effectors in the cytosol of host cells. Bacteria are transiently found in an internalization vacuole (B), which is disrupted upon recruitment of Rab11-positive macropinosomes (C), leading the uptake of vacuole remnants by autophagosomes (D). Cytoplasmic bacteria polymerize actin and move intracellularly (E), while septin cages can also function as bacterial traps (F). Upon reaching the host cell plasma membrane, *Shigella* induces the formation of protrusions (G) and subsequently localize in double-membrane vacuoles in neighboring cells (H). Disruption of this compartment allows the start of a new infection cycle.

act directly with human vinculin, which would allow a link with the actin cytoskeleton [114]. More precisely, IpaA promotes the capping of F-actin barbed ends via vinculin controlling the polymerization dynamics of actin at the bacterial contact sites [114]. Also, the targeting of $\beta 1$ -integrin by IpaA has been suggested to indirectly stimulate the GTPase activity of the small GTPase RhoA, which would lead in turn to the loss of actin stress fibers via the ROCK/myosin-II pathway [115]. Such stress fiber disassembly may facilitate the availability of actin for the formation of membrane ruffles at the bacterial entry site (Fig. 3A). IpaC is a constituent of the T3SS translocon complex and plays a direct role in the bacterial entry process. It recruits and activates the host tyrosine kinase Src to the site of bacterial contact where it phosphorylates the actin-binding protein cortactin that implicates the Arp2/3 complex and leads to actin remodeling and polymerization [116]. In addition, the activated form of cortactin interacts also with the Src-related tyrosine kinase Crk that further boosts actin polymerization at the *Shigella* entry site [117].

The two homologues effectors, IpgB1 and IpgB2, play a key role in *Shigella* entry. IpgB1 stimulates Rac and Cdc42 [118]. It binds ELMO, a protein involved in engulfment and cell motility, leading to the activation of Rac1 through the ELMO/Dock180 pathway. Consequently, it induces Arp2/3 complex-dependent membrane ruffling at the bacterial contact site [119], and it has also been revealed as

a pace maker for *Shigella* internalization [120]. In contrast, IpgB2 binds to mDia1 and to ROCK, which leads to actin nucleation and stress fiber formation, thus mimicking RhoA [121,122]. Even though IpgD, a PI(4,5)P₂ phosphatase that specifically depletes PtdIns(4,5)P₂ into PtdIns(5)P, does not affect *Shigella* entry in general, it may weaken the connection between cortical actin and the plasma membrane to facilitate membrane extensions [15]. Another effector that has been implicated during the early entry of *Shigella* is VirA, which correlates with local degradation of microtubules, which indirectly affects ruffle formation through a feedback loop via RhoA and Rac1 [123–125]. This illustrates, how different enzymatic activities of the bacterial effectors lead to finely tuned spatiotemporal modulations of the host cytoskeleton at the contact site allowing the internalization of *Shigella* in a vacuolar compartment.

4.2. Rupture of the *Shigella* containing vacuole

Shigella ruptures its endocytic vacuole rapidly upon internalization to reach the host cytoplasm (Fig. 3B). The T3SS in particular, through the action of the translocon complex proteins IpaB and IpaC, has been proposed to play a crucial role in the direct destabilization of the *Shigella* containing vacuole: the T3SS translocon proteins IpaB and IpaC could form a pore complex in cholesterol-

rich membrane domains of the vacuole, causing its disruption and subsequent bacterial escape to the cytosol [126,127]. This was substantiated through experiments that demonstrated red blood cell hemolysis *via* IpaB and IpaC [128]. Recently, T3SS reconstitution of the *Shigella* injection device in *E. coli* resulted in the destabilization of the vacuole containing the *E. coli* expressing the T3SS in macrophages and hinted at the capacity of the T3SS to disrupt endocytic vacuoles [129]. Nevertheless, it has been puzzling that other T3SS containing bacteria, including *Salmonella* and *Yersinia* remain mainly within phagolysosomes without damage arguing for a vacuolar damage mechanism that requires additional factors. For example, a non characterized region on the *Shigella* virulence plasmid has been implicated in the uncoupling of *Shigella* entry step and the vacuolar escape step [130]. Novel approaches for the tracking of vacuolar damage in single cells at high temporal resolution reveals that this process occurs in less than 10 min after bacterial entry [120,131,132]. Using this method, a new mechanism of vacuolar membrane damage caused by *Shigella* has been discovered, which depends on the subversion of host membrane trafficking pathways through the injection of *Shigella* T3SS effectors. A high-content small interfering RNA (siRNA) screen identified several host proteins that modulate *S. flexneri* vacuolar membrane rupture [133]. They include a network of Rab GTPases, namely Rab5 and Rab11, a component of the host recycling pathway (Fig. 3C). Depletion of Rab5 and Rab11 delays vacuolar rupture, and both GTPases are recruited to the bacterial entry site through the effector IpgD that converts of PI(4,5)P2 to PI(5)P. Also, the *ipgD* mutant increases the time of vacuolar escape by *Shigella*. Using correlative light electron microscopy (CLEM) in three dimensions, it was further shown that Rab11 was only recruited to macropinosomes in the surrounding of the bacterial containing vacuole, and that *Shigella* was not taken up within such spacious compartments [134]. It was therefore identified that *Shigella* exclusively interacts with *in situ* induced macropinosomes having no contact with other endomembrane compartments. The interaction between the *Shigella* containing vacuole and surrounding macropinosomes through membrane contacts is necessary for vacuolar rupture through a mechanism that requires further investigation [134].

4.3. *Shigella* lifestyle after vacuolar rupture and cell-to-cell spread

Vacuolar rupture and bacterial escape into the cytoplasm results in the generation of small vesicles derived from vacuolar membrane remnants, which are associated with poly-ubiquitinated proteins. The autophagy marker LC3 and the adaptor p62 are recruited, as well as inflammasome components and caspase-1, and the membrane remnants are targeted to autophagic degradation [120,135] (Fig. 3D). Therefore, the vacuolar rupture process itself represents a site for the establishment of a signaling platform in the host cell. Within the cytoplasm, *Shigella* polymerizes actin at one of its poles with the help of the outer membrane protein IcsA (or VirG). IcsA interacts with and recruits N-WASP, which in turn activates the Arp2/3 complex to form an actin comet tail [136] (Fig. 3E). Intracellular bacterial motility can reach speeds of 3–26 $\mu\text{m}/\text{min}$ [136,137]. Structurally, as for *Listeria*, actin comet tails are composed of short, highly branched cross-linked actin filaments that can leave a long trail behind one of the bacterial poles. A subset of intracytoplasmic *Shigella* may be found surrounded by septin-cages that impede their mobility and interact with the host autophagy machinery [138] (Fig. 3F). As *Listeria*, intracellular moving *Shigella* make protrusions upon reaching the host cell plasma membrane (Fig. 3G) which allows them to invade neighboring cells, in which bacteria will be found in two-membrane secondary vacuoles (Fig. 3H).

Rupture of these secondary vacuoles by still non-identified mechanisms initiate a new intracellular infection cycle.

5. *Yersinia* spp.

Yersinia spp. are Gram-negative, facultative anaerobes that belong to the Enterobacteriaceae family, for which eighteen species have been reported, including three species that display pathogenic features in humans and animals [139]. *Y. pseudotuberculosis* and *Y. enterocolitica* are enteropathogenic bacteria responsible for enteritis, ileitis, and mesenteric lymphadenitis; *Y. pestis*, the etiological agent of bubonic plague, is still endemic in rodent populations on several continents and considered by the World Health Organization as a reemerging disease. *Y. pseudotuberculosis* and *Y. enterocolitica* (referred from now on as *Yersinia*) are mainly transmitted by ingestion: upon uptake of contaminated food or water, bacteria gain access to the gastrointestinal tract, reach the terminal ileum and translocate across the epithelial barrier by transcytosis through antigen-sampling M cells associated with Peyer's patches [140]. Thereafter, *Yersinia* predominantly survives extracellularly (Fig. 4A), inhibiting its internalization by PMNs and monocytes (Fig. 4B).

5.1. Facultative intracellular life

5.1.1. Cell invasion

In the gastrointestinal tract, *Yersinia* displays on its surface at high density the outer membrane protein (OMP) invasin [141]. Invasin is related to the intimins OMPs present in enteropathogenic and enterohemorrhagic *Escherichia coli*, and binds with high affinity to multiple members of the $\beta 1$ chain integrin family on a variety of cells, including M cells [142,143]. Binding of $\beta 1$ integrins by invasin promotes recruitment at bacterial entry sites of the focal adhesion kinase (FAK) and Src, which normally drive actin-dependent rearrangements required for cellular migration, but which are used by *Yersinia* to promote bacterial invasion [144,145]. Indeed, as in the case of *Listeria* infection process, the small GTPase Rac1 is subsequently recruited at *Yersinia* entry sites, where it induces actin polymerization *via* the Arp2/3 complex [146] and bacterial engulfment through zipper phagocytosis (Fig. 4C). The small GTPase Arf6 also participates in the bacterial invasion process and cooperates with Rac1 in the localized activation of PIP 5-kinase, which produces a focal increase of the phosphoinositide PI(4,5)P2 [147], probably required to recruit other actin-remodeling proteins required for *Yersinia* invasion. The final stages of *Yersinia* entry into host cells require PI 3-kinase-dependent fusion of Rab5-positive vesicles with bacterial-containing compartments that are still open to the extracellular space: the host lipid phosphatase OCRL is then recruited to these non-sealed vacuoles promoting hydrolysis of PI(4,5)P2 and vacuole scission from the plasma membrane [148] (Fig. 4D). Clathrin coats are, as for *Listeria*, critical for *Yersinia* internalization in host cells [23]. The lollipop-shaped multifunctional adhesin YadA also facilitates epithelial cell entry in the absence of invasin activity, through indirect binding of $\beta 1$ integrins *via* engagement of extra-cellular matrix molecules [149].

5.1.2. Autophagy subversion

Yersinia actively inhibits its own internalization in PMNs and macrophages (see below). Interestingly, while *Y. enterocolitica* upon internalization in phagocytic cells is degraded by autophagy [150], *Y. pestis* replicates within intracellular compartments positive for the autophagy marker protein LC3 and which do not acidify [151]. Recently, it has been suggested that the small GTPase Rab1 plays a critical role in the inhibition of acidification by directly inhibiting the maturation of the *Y. pestis*-containing compartment [152]. *Y. pseudotuberculosis* has been also found to proliferate in LC3-positive

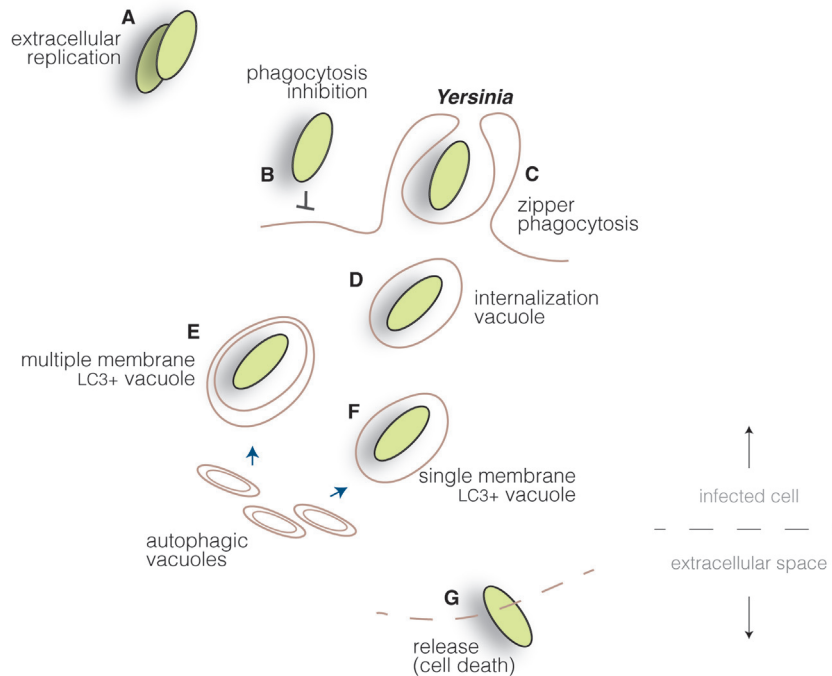


Fig. 4. *Yersinia* cycle. *Yersinia* proliferate mainly as extracellular pathogens (A), as they actively inhibit cellular uptake by injecting type III secretion system effectors into host cells, blocking actin cytoskeleton rearrangements (B). However, through the interaction of surface invasion proteins and host cell receptors, *Yersinia* can induce its internalization in specific cell lines (C). Bacteria are found initially within internalization vacuoles (D) which, through interaction with autophagic vacuoles, can mature into multiple (E) or single (F) membrane compartments, depending on the cell type. *Yersinia* are probably released upon cell death in cells permissive for bacterial replication (G).

vacuoles: in bone marrow-derived macrophages, the bacteria are detected in classical autophagosomes characterized by multiple limiting membranes which also avoid acidification [153] (Fig. 4E). Surprisingly, in epithelial HeLa cells, *Y. pseudotuberculosis* resides in single-membrane compartments also labelled by LC3, suggesting an autophagosomal origin [154] (Fig. 4F). The vesicle-associated membrane protein 7 (VAMP7), which belongs to the SNARE family of mediators of membrane docking and fusion, was found to regulate the association of LC3 to both single- and multiple-membrane compartments supporting *Y. pseudotuberculosis* replication; however, VAMP3 was shown to play a critical role in the commitment of bacteria towards single-membrane LC3-positive compartments in HeLa cells, suggesting that this SNARE functions as a molecular checkpoint for bacterial trafficking to single- or double-membrane autophagosomes [154]. The potential functional differences of these two types of autophagosomes are unknown. Interestingly, vacuolar escape for *Y. enterocolitica* has been described [155] but the relevance of this observation requires further investigation.

5.2. Extracellular life

As mentioned above, *Yersinia* spp. has developed molecular adaptations to survive within macrophages if uptake takes place. However, the first bacterial defense strategy to avoid degradation is to actively inhibit internalization within professional phagocytes. The T3SS of *Yersinia*, which is one of the first to have been thoroughly characterized, delivers into the cytoplasm of host cells, molecules which interfere with actin cytoskeleton rearrangements in order to inhibit phagocytosis [156]. The effector YopH is a tyrosine phosphatase that dephosphorylates critical cytoskeletal-related proteins such as FAK, p130Cas and paxillin [157], inhibiting Rac1 activation required for bacterial internalization, as mentioned above [145]. Direct manipulation of Rho GTPases is performed by YopE, which functions as a GTPase-activating protein (GAP) for Rho, Rac1 and Cdc42, completely inhibiting the rearrangement

of the actin cytoskeleton [158,159]. The effector YopO/YpkA is a multidomain effector which displays a GDP-dissociation inhibitor (GDI)-like domain which sequesters the GDI-free pool of Rac1 localized at the plasma membrane [160] and which also displays a phosphatase domain which targets the vasodilator-stimulated phosphoprotein (VASP), another critical regulator of actin assembly [161]. Finally, the *Y. enterocolitica*-specific YopT is a cysteine protease that cleaves the carboxy-terminal prenyl groups of Rho, Rac1 and Cdc42, disrupting their membrane localization [162].

6. Conclusions

In the course of infection, bacterial pathogens manipulate host cell membranes in exquisite ways. During internalization, the subversion of small Rho GTPases and phosphoinositide metabolism has been extensively characterized as mechanisms to control actin cytoskeleton rearrangements and vesicular traffic. Several recent findings have now refined our current view of bacterial infection strategies to harness host membranes, and clathrin stands as a novel critical regulator of actin polymerization for zippering pathogens such as *Listeria* and *Yersinia* [23]. Bacterial transcytosis has also emerged as a novel invasion mechanism for *Listeria* that allows translocation across the epithelial barrier [5]. It would be interesting to determine whether *Yersinia*, which also uses a zipper invasion mechanism based on the activation of surface host cell receptors, exploits a similar strategy during the initial steps of host infection. Another important finding concerns the co-existence of diverse Arp2/3 complexes in cells which differentially participate in *Listeria* entry or actin comet tail formation [29]. Similar observations have been made for viral infection [58] and focal adhesion formation [163], highlighting that cellular functions implicating the Arp2/3 complex may need to be re-investigated.

The rupture process of bacterial-containing compartments has been also recently enriched by the characterization of the *Francisella* T6SS, that appears essential for bacterial escape to the host

cytosol [94] and by the identification of small GTPases of the Rab family which are required for *Shigella* cytosolic translocation [133]. In the future, it will be important to determine whether other host molecules can directly influence vacuolar membrane stability and what are the specific molecular contributions of T6SS effectors for *Francisella* phagosomal escape. Concerning bacterial replication within host cells, the finding that *Yersinia* may reside in single- or multiple-membrane compartments [154] highlights the versatility of bacterial pathogens to adapt their molecular survival strategies to their specific local infection environment. This is also exemplified by the *Francisella* nutritional virulence phenotype [95].

Infectious diseases are still a major public health problem in the 21st century. Profound knowledge of how host cellular components are subverted by bacterial pathogens should allow identification of host targets to be used in novel therapeutic strategies.

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