ADP-heptose is a pathogen-associated molecular pattern of *Shigella flexneri* infection

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ABSTRACT

During an infection, the detection of pathogens is mediated through interactions between pathogen-associated molecular patterns (PAMPs) and pathogen recognition receptors. β -Heptose 1,7-bisphosphate (β HBP), a metabolite of the lipopolysaccharide (LPS) biosynthesis pathway, was recently identified as a PAMP of gram-negative bacteria. It was reported that β HBP sensing leads within minutes to oligomerization of the protein TIFA, a mechanism controlling NF- κ B activation and pro-inflammatory gene expression. Here, we compared the ability of chemically synthesized β HBP and *Shigella flexneri* lysate to induce TIFA oligomerization in epithelial cells. In contrast to lysate, we found that β HBP fails to trigger rapid oligomerization of TIFA. β HBP only induces delayed signaling, suggesting that it has to be processed intracellularly to induce inflammation. By dissecting the LPS biosynthesis pathway with deletion mutants and functional complementation experiments, we show that ADP-D-*glycero*- β -D-*manno*-heptose and ADP-L-*glycero*- β -D-*manno*-heptose are the bacterial metabolites responsible for rapid TIFA oligomerization, and that they strongly induce interleukin-8 expression during *S. flexneri* infection. Altogether, our results rule out a major role of β HBP in *S. flexneri* infection and identify ADP-heptose as a new PAMP.

INTRODUCTION

During an infection, an immune response is triggered by the detection of pathogens. This is mainly achieved by cells of the innate immune system including macrophages and neutrophils. Epithelial and endothelial cells are also involved in this process. They act as sentinels of the immune system and contribute, via the secretion of inflammatory mediators, to elicit an integrated immune response. At the molecular level, pathogen recognition is mediated by the interactions between pathogen-associated molecular patterns (PAMPs) and pathogen recognition receptors. Well characterized bacterial PAMPs include among others lipopolysaccharide (LPS), flagellin, peptidoglycan, lipoteichoic acid and DNA. β -Heptose 1,7-bisphosphate (β HBP) was recently identified as a PAMP of *Neisseria meningitidis* [1]. β HBP is a soluble metabolite of the LPS biosynthesis pathway. In *N. meningitidis*, it is produced by the D- β -D-heptose 7-phosphate kinase activity of the HldA protein that phosphorylates D-*glycero*- β -D-*manno*-heptose 7-phosphate into β HBP [2]. Gaudet et al. showed that it can be secreted by *N. meningitidis* and sensed in the cytosol of eukaryotic cells after being internalized by endocytosis [1]. Our laboratory reported that β HBP sensing was involved in the inflammatory response of epithelial cells to *Shigella flexneri* and *Salmonella typhimurium* infection [3]. This finding was based on the analysis of the *hlde* and *gmhb* deletion mutants of the LPS biosynthesis pathway. In these bacterial species closely related to *Escherichia coli*, *hlde* encodes a bifunctional protein that harbors both D- β -D-heptose 7-phosphate kinase and D- β -D-heptose 1-phosphate adenylyltransferase activities, responsible for the production of β HBP and ADP-D-*glycero*- β -D-*manno*-heptose (ADP-D- β -D-heptose), respectively [4]. The dephosphorylation of β HBP into D-*glycero*- β -D-*manno* heptose 1-phosphate is

provided by the phosphatase activity of GmhB [5]. We showed that infection with an *hlde* deletion ($\Delta hlde$) mutant failed to induce the expression of the inflammatory cytokine interleukin-8 (IL-8) whereas this latter was restored after infection with a mutant deleted for *gmhb* ($\Delta gmhb$) [3]. *S. flexneri* infection leads to oligomerization of the proteins TRAF-interacting protein with FHA domain-containing protein A (TIFA) and TNF receptor-associated factor 6 (TRAF6), and activation of the transcription factor NF- κ B. TIFA is a 20-kDa protein that was first identified as a TRAF6-interacting protein in a yeast two-hybrid screen [6]. It contains a forkhead-associated (FHA) domain, known to bind phosphothreonines and phosphoserines [7], and a consensus TRAF6-binding motif. TIFA oligomerization is dependent on the phosphorylation of a threonine at position 9 and on the FHA domain [8]. Unphosphorylated TIFA is thought to exist as an intrinsic dimer constitutively linked to TRAF6. When T9 is phosphorylated, this is recognized by the FHA domain of other TIFA dimers leading to its oligomerization and thus the oligomerization of TRAF6. This enhances the E3 ubiquitin ligase activity of TRAF6 and leads to the activation of the NF- κ B pathway [9]. We recently revealed the role of ALPK1 in bacterial infections and showed that this atypical kinase of the α -kinase family regulates the oligomerization of TIFA [3]. More recently, the role of β HBP sensing and the contribution of the ALPK1/TIFA pathway were also observed in *Helicobacter pylori* infection [10–12], reinforcing the broad implication of this newly emerging pathway in innate immunity. In addition to the use of LPS biosynthesis mutants, the pro-inflammatory activity of β HBP was directly confirmed. It was first synthesized *in vitro* from the sedoheptulose 7-phosphate precursor by the sequential activities of purified GmhA and HldE [1]. More recently, β HBP was also chemically synthesized [13–16]. Its pro-inflammatory activity was validated on the basis that it induced NF- κ B activation after several hours of treatment

[14,15]. In this report, we compared the ability of chemically synthesized β HBP and *S. flexneri* lysates to induce the oligomerization of TIFA and the production of IL-8 in human epithelial cells. In contrast to lysate from wild-type bacteria, we found that β HBP was unable to induce rapid oligomerization of TIFA. Oligomerization was only observed after more than one hour, indicating that cellular processing of β HBP was necessary to trigger this mechanism. It also suggested that another PAMP was therefore responsible for the early mechanism of TIFA oligomerization occurring during infection [3]. By combining deletion mutants and functional complementation experiments, we identified ADP-heptose as a new PAMP of *S. flexneri* infection.

RESULTS AND DISCUSSION

β HBP fails to induce rapid oligomerization of TIFA

In order to characterize the role of β HBP in innate immunity, this bacterial metabolite was chemically synthesized. A synthetic route was implemented, that was inspired, for the most part, from the work of Vincent and coworkers [17] (Figure 1 and Materials and Methods). Briefly, starting from thiophenyl α -D-mannoside (**2**) [18], easily synthesized in three steps from D-mannose (**1**), a sequence of protection/deprotection steps afforded the production of 2,3,4-tri-O-benzylated derivative **3** on a large scale (55 mmol) and in good yield. Homologation of this thiomannoside intermediate by means of a Parikh-Doering oxidation/Wittig sequence gave the terminal olefin-containing heptoside **4** [19]. The latter was subsequently reacted with catalytic osmium tetroxide to obtain a 7:3 diastereomeric mixture of diols, from which the major isomer was isolated by chromatography. A subsequent four step protection/deprotection procedure was applied to give hemiacetal **5**. The hemiacetal was submitted to a Mitsunobu-mediated *bis*-phosphorylation at OH-1 and OH-7 by use of a combination of dibenzyl phosphate, DIAD, P(CIPh)₃, and triethylamine in tetrahydrofuran [13]. Chromatography of the obtained α/β mixture of the perbenzylated diphosphorylated monosaccharide gave the α -anomer and β -anomer in 30% and 41% yield, respectively. The purified β -heptose was subjected to a palladium-catalyzed hydrogenation followed by H⁺/Na⁺ exchange to give the expected β HBP [13]. The pro-inflammatory activity of chemically synthesized β HBP was first tested by measuring the production of IL-8 after transient permeabilization of cell membranes with digitonin. In this assay, the cellular process of β HBP internalization is bypassed and the mechanisms of PAMP recognition and inflammatory signaling can thus be directly

assessed. HeLa cells were treated for 30 minutes with digitonin and synthetic β HBP at indicated concentrations (Figure 2A and 2B). As a control, cells were treated with digitonin alone or digitonin and wild-type (wt) *S. flexneri* bacterial lysate. Cells were then washed, treated with monensin to trap IL-8 intracellularly, and IL-8 was monitored in single cells after 6 hours with automated microscopy (see Materials and Methods). Data showed that synthetic β HBP induced the production of IL-8 in a dose-dependent manner (Figure 2A and 2B). Interestingly, the β HBP concentration required to induce IL-8 expression to the level of *S. flexneri* lysate was unexpectedly high (Figure 2A and 2B). Since a millimolar-range concentration of β HBP is probably not reached in our bacterial lysate [20], this observation raised the possibility that additional bacterial factors may contribute to elicit the IL-8 response. β HBP-induced IL-8 expression was previously shown to depend on the oligomerization of TIFA proteins, a process controlling NF- κ B activation [1,3]. We therefore analyzed the effect of chemically synthesized β HBP on TIFA oligomerization. HeLa cells expressing GFP-tagged TIFA protein (TIFA-GFP) were treated as described above and TIFA-GFP oligomerization was monitored after 6 hours. It was quantified in single cells by automated image analysis as described in Materials and Methods. In line with IL-8 data, β HBP induced the formation of large TIFA-GFP oligomers in a dose-dependent manner (Figure 2C and 2D). This result was in agreement with previous reports showing that synthetic β HBP induces NF- κ B activation after several hours of treatment [14,15]. Given that TIFA oligomerization is observed within minutes of *S. flexneri*, *S. typhimurium* or *H. pylori* infections [3,10], we also monitored TIFA oligomerization after 30 minutes of treatment. Strikingly, whereas cells treated with *S. flexneri* lysate induced massive oligomerization of TIFA-GFP within this time period, synthetic β HBP failed to do so (Figure 2E). At least one hour was required to observe significant oligomerization of

TIFA-GFP in a small fraction of cells treated with 1 mM synthetic β HBP (Figure 2F). For this latter concentration, the level of TIFA-GFP oligomerization observed in cells treated with *S. flexneri* lysate was only reached after 2 hours of β HBP treatment (Figure 2F). Such delay in the response to β HBP suggested that this bacterial metabolite has to be processed intracellularly to trigger inflammation signaling, and that the sensing of another bacterial PAMP is likely responsible for the rapid mechanism of TIFA oligomerization observed upon *S. flexneri* lysate treatment.

Bacterial ADP-heptose synthesis is required for rapid oligomerization of TIFA.

Our results showing that β HBP was unable to trigger rapid oligomerization of TIFA-GFP (Figure 2E and 2F) opened a new avenue for the identification of new bacterial PAMPs inducing inflammatory signaling upon *S. flexneri* infection. Given that we previously reported that $\Delta hldE$ *S. flexneri* bacteria completely failed to trigger TIFA oligomerization and inflammatory gene expression [3], we further analyzed the role of the bifunctional enzyme HldE. More specifically, we analyzed the contribution of its D- β -D-heptose 7-phosphate kinase and D- β -D-heptose 1-phosphate adenylyltransferase activities that synthesize β HBP and ADP-D- β -D-heptose, respectively (Figure 3A). For this, we took advantage of the fact that in *N. meningitidis*, these enzymatic activities are harbored by two distinct proteins encoded by the *hlda* and *hldc* genes [2], and examined the ability of lysates from $\Delta hldE$ *S. flexneri* complemented with *hldc*, *hlda*, *hldc* or *hlda* and *hldc* to induce TIFA oligomerization in digitonin-permeabilized cells. Whereas treatment with wt *S. flexneri* lysate induced massive oligomerization of TIFA-GFP at 30 minutes, lysate from $\Delta hldE$ bacteria failed to do so (Figure 3B). As expected, complementation of $\Delta hldE$ bacteria with a plasmid encoding HldE (pHldE) fully restored

fast oligomerization of TIFA, confirming the critical role of HldE in this process. Interestingly, lysate from $\Delta hldE$ bacteria complemented with plasmid-encoded HldA (pHldA) had no effect at 30 minutes (Figure 3B). This result was in line with synthetic β HBP data showing that this metabolite was indeed unable to trigger rapid oligomerization of TIFA-GFP (Figure 2E and 2F). By showing that bacterial D- β -D-heptose 1-phosphate adenylyltransferase activity was required, this new finding indicated that the PAMP eliciting rapid oligomerization was produced downstream of the adenylyltransferase enzymatic reaction within the LPS biosynthesis pathway (Figure 3A). Complementation of $\Delta hldE$ *S. flexneri* with plasmid-encoded HldC (pHldC) failed also to trigger TIFA oligomerization at 30 minutes, showing that bacterial synthesis of β HBP, although not sufficient, was required to trigger early signaling. As expected, this process was restored with $\Delta hldE$ bacteria complemented with both pHldA and pHldC (Figure 3B). The observation that double complementation only led to partial restoration may be explained by a low expression of the HldA and HldC proteins encoded from two different plasmids, or by a difference in enzymatic efficacy between HldE of one part and HldA and HldC on the other hand. Together, these results showed that D- β -D-heptose 7-phosphate kinase and D- β -D-heptose 1-phosphate adenylyltransferase activities are both required to stimulate rapid pro-inflammatory signaling, indicating that bacterial synthesis of β HBP and ADP-D- β -D-heptose is necessary. In order to further characterize the PAMP involved in rapid inflammatory signaling, we analyzed the *hldD* and *waaC* deletion mutants of the LPS biosynthesis pathway, named $\Delta hldD$ and $\Delta waaC$ respectively. HldD harbors an epimerase activity catalyzing the interconversion between ADP-D- β -D-heptose and ADP-L-*glycero*- β -D-*manno*-heptose (ADP-L- β -D-heptose) [21] (Figure 3A). Directly downstream of HldD, WaaC, which has an ADP-heptose-lipopolysaccharide

heptosyltransferase activity, allows heptose transfer to the LPS core [21]. Data showed that $\Delta hldD$ and $\Delta waaC$ lysates were both very effective at triggering TIFA-GFP oligomerization at 30 minutes, indicating that the PAMP controlling this process was produced upstream of HldD or WaaC respectively, and was accumulated in these mutants. Altogether, by showing that the PAMP of interest was dependent on β HBP, produced downstream of the D- β -D-heptose 1-phosphate adenylyltransferase enzymatic reaction, and upstream of HldD or WaaC (Figure 3B), our results point to the role of ADP-D- β -D-heptose and ADP-L- β -D-heptose (ADP-D/L- β -D-heptose, also named ADP-heptose for simplification reason) in triggering rapid oligomerization of TIFA upon treatment with *S. flexneri* lysate. TIFA-GFP oligomerization and IL-8 production were also analyzed at 6 hours. Based on synthetic β HBP data (Figure 2A, 2B, 2C, 2D and 2F), we hypothesized that at this time point β HBP and ADP-D/L- β -D-heptose contribute to inflammatory signaling. Our data confirmed a critical role for bacterial expression of HldE in lysate-induced TIFA-GFP oligomerization and IL-8 production at 6 hours (Figure 3C and 3D). In line with synthetic β HBP data (Figure 2A, 2B, 2C and 2D), pHldA-mediated complementation partially restored TIFA-GFP oligomerization and IL-8 production (Figure 3C and 3D), showing that bacterial synthesis of β HBP is sufficient to trigger delayed inflammatory responses. Complementation of $\Delta hldE$ *S. flexneri* lysate with pHldC failed to trigger TIFA-GFP oligomerization (Figure 3C) as well as IL-8 production (Figure 3D), showing that bacterial synthesis of β HBP is strictly required for both processes. Complementation with pHldA and pHldC partially restored both oligomerization and IL-8 expression (Figure 3C and 3D). The observation that expressing HldA or both HldA and HldC in $\Delta hldE$ bacteria had a rather similar effect may be explained by a higher expression of HldA in single complementation experiments and enhanced β HBP concentration in

bacteria lacking D-β-D-heptose 1-phosphate adenylyltransferase activity. As expected, lysates from $\Delta hldD$ and $\Delta waaC$ mutants induced massive TIFA-GFP oligomerization and IL-8 production, confirming the role of ADP-D/L-β-D-heptose in inflammation (Figure 3C and 3D). Altogether, our data showed that βHBP is only able to trigger delayed inflammatory signaling, suggesting that it is not directly sensed by the innate immune system but that it first has to be processed in cells. In addition, our results reveal that the bacterial factors ADP-D/L-β-D-heptose trigger early oligomerization of TIFA-GFP and significantly contribute to the inflammatory response observed upon treatment with *S. flexneri* lysate.

ADP-heptose is a PAMP of *S. flexneri* infection

The use of digitonin-permeabilized cells is a powerful method to directly address PAMP recognition and inflammatory signaling. However, it does not provide information on the actual involvement of a given PAMP during infection. The respective contribution of βHBP and ADP-D/L-β-D-heptose was therefore tested in *S. flexneri* infection experiments. For this purpose, HeLa cells were infected with wt *S. flexneri* or with the different mutants of the LPS biosynthesis pathway described above, and TIFA-GFP oligomerization was visualized 30 minutes post infection. Representative images are shown in Figure 4A. As previously reported [3], wt bacteria induced rapid oligomerization of TIFA-GFP in infected cells. In contrast, the $\Delta hldE$ mutant failed to do so whereas pHldE-complementation fully restored oligomerization. Together, these results confirmed the critical implication of HldE in the early host response to *S. flexneri* infection. In line with data obtained with bacterial lysates (Figure 3B), complementing the $\Delta hldE$ mutant with pHldA or pHldC failed to trigger TIFA-GFP oligomerization at 30

minutes (Figure 4A), showing that bacterial D- β -D-heptose 7-phosphate kinase and D- β -D-heptose 1-phosphate adenylyltransferase activities are both necessary to trigger rapid oligomerization of TIFA-GFP. This mechanism is indeed restored when cells were infected with pHldA and pHldC-complemented $\Delta hldE$ bacteria (Figure 4A). Finally, oligomerization was observed in cells infected with $\Delta hldD$ and $\Delta waaC$ mutants, showing that the PAMP responsible for this process was produced upstream of HldD or WaaC, respectively. In *E. coli*, $\Delta hldE$ and $\Delta waaC$ mutants have a phenotype of increased bacterial auto-aggregation and stronger cell surface hydrophobicity compared to wt bacteria [22]. In *S. flexneri*, we previously observed that these mutants are at least hundred times more invasive than wt [3], making the comparison of TIFA oligomerization between all strains inappropriate. For this reason, only mutants that had the same range of infectivity (Figure 4B) were considered for quantification. Data confirmed that none of the single-complemented mutants induced oligomerization in infected cells (Figure 4C). In contrast, although infectivity of the double-complemented mutant was slightly reduced compared to $\Delta hldE$ bacteria (Figure 4B), it significantly led to TIFA-GFP oligomerization in infected cells (Figure 4C). The weak impact of HldA and HldC expression on bacterial infectivity compared to the hundred fold change that was previously observed between $\Delta hldE$ and wt [3] suggested that the LPS of these bacteria was still significantly altered and that complementation was therefore only partial. This observation was correlated with the rather modest restoration of inflammatory signaling observed with the double-complemented mutant (Figure 3B, 3C and 3D), and may be explained by low expression or weak enzymatic activities of HldA and HldC in these bacteria. In line with lysate data (Figure 3B), infection with $\Delta hldD$ and $\Delta waaC$ mutants induced massive oligomerization at 30 minutes (Figure 4C). Altogether, these results showed that β HBP is not sufficient to trigger early signaling

in *S. flexneri* infected cells and revealed the critical role of ADP-D/L- β -D-heptose in this process. In order to better understand the dynamics of the inflammatory response during infection, TIFA-GFP oligomerization and IL-8 production were also analyzed 4 hours post infection. In agreement with a role of β HBP in late signaling, oligomerization was visible in a small fraction of cells infected with pHIdA-complemented bacteria (Figure 4D). However, no IL-8 was observed in response to this infection (Figure 4E). We hypothesized that the level of TIFA-GFP oligomerization was too low to elicit IL-8 production within 4 hours. This time point was chosen to prevent cell toxicity due to bacterial replication. As expected, infection with pHIdC-complemented bacteria failed to trigger TIFA-GFP oligomerization (Figure 4D) and IL-8 production (Figure 4E), confirming the strict requirement of β HBP bacterial synthesis. Infection with the double-complemented mutant induced significant oligomerization of TIFA and IL-8 production, showing that the presence of both enzymes allows IL-8 expression. This latter is more likely induced by ADP-D/L- β -D-heptose than β HBP since, within 4 hours of infection, the single pHIdA-complemented mutant was unable to produce IL-8 (Figure 4E). Finally, infection with $\Delta hldD$ and $\Delta waaC$ mutants showed massive oligomerization and IL-8 production, confirming the ability of ADP-D/L- β -D-heptose to trigger inflammation during *S. flexneri* infection.

In conclusion, our data showed that β HBP can only trigger delayed inflammatory signaling, and that it has a marginal contribution in controlling the IL-8 response during *S. flexneri* infection. The delay in the response to β HBP suggests that it has to be intracellularly processed to induce inflammation. β HBP is therefore likely not a true PAMP, which is, by definition, directly sensed via the interaction with a cognate pathogen recognition receptor. Thus, the role of ADP-D/L- β -D-heptose in *S. flexneri*

infection challenges our previous results implicating β HBP [3]. The ability of the $\Delta gmhB$ mutant to induce TIFA oligomerization was the rational basis to conclude that the PAMP inducing TIFA-dependent innate immunity was indeed β HBP. However, this argument is weakened by the possibility that the $\Delta gmhB$ mutant might only have a partial mutant phenotype. Indeed, the observation that, in contrast to $\Delta hldE$, $\Delta hldD$ and $\Delta waaC$, the $\Delta gmhB$ mutant is not hyper-invasive [3], indicated that its LPS was only partially altered and that ADP-D/L- β -D-heptose synthesis was therefore still occurring in these bacteria. A partial phenotype of the $\Delta gmhB$ mutant was also observed in *E. coli*, and the existence of a yet-unidentified phosphatase activity that compensates for the absence of GmhB was proposed [22]. Given that the $\Delta gmhB$ -based argument was also used in other studies [11,12], the direct contribution of β HBP has been likely more widely overestimated. The identification of ADP-heptose as a PAMP of *S. flexneri* infection opens a new avenue to understand how inflammation is regulated at the level of the intestinal epithelium in shigellosis. The mechanism of ADP-heptose sensing will have to be thoroughly characterized in space and time during infection. At the time of finalizing this manuscript, Zhou et al. elegantly reported that the kinase ALPK1 is the cytosolic immune receptor for ADP-heptose [23]. More work is required to identify the respective contribution of ADP-D- β -D-heptose and ADP-L- β -D-heptose, to understand how these metabolites are delivered in cells during infection, and in particular, whether they are injected by the type III secretion apparatus of *S. flexneri*. Given that ADP-D/L- β -D-heptose are broadly present in bacteria, our study opens also new perspectives to elucidate the molecular mechanisms regulating innate immunity in infections by other severe human pathogens, including *H. pylori*, *S. typhimurium* and *Pseudomonas aeruginosa*. Beyond infections, it will also contribute to characterize new molecular

cross-talks between bacteria of the microbiota, the intestinal epithelium and the immune system that regulate intestinal homeostasis.

MATERIALS AND METHODS

Cells

HeLa cells (American Type Culture Collection) were cultured in Dulbecco's modified Eagle's (DMEM) medium supplemented with 10% FCS and 2 mM Glutamax-1 (complete growth medium). HeLa cells stably expressing TIFA-GFP were obtained after transfection of pEGFP-C1 plasmid encoding TIFA-GFP and geneticin selection at 1 mg.mL⁻¹.

Bacterial strains and generation of mutants

Wild-type (wt), $\Delta hldE$ and $\Delta waaC$ M90T *S. flexneri* strains have been previously described [3]. The $\Delta hldD$ mutant was generated by allelic exchange using a modified protocol of lambda red-mediated gene deletion [24]. Briefly, the kanamycin cassette of the pkD4 plasmid was amplified by PCR with the primers listed in Table 1. The purified PCR product was electroporated into the wt strain expressing the genes for lambda red recombination from the pKM208 plasmid. Recombinants were selected on TSB plates containing 50 µg.mL⁻¹ of kanamycin. Single colonies were screened by PCR. *S. flexneri* M90T expressing the *hldA* gene from *N. meningitidis* was previously described [3]. *S. flexneri* M90T expressing the *hldC* gene from *N. meningitidis* was generated as follows. *hldC* was amplified by PCR from a bacterial lysate with the primers listed in Table S1. After gel purification (Macherey-Nagel), the PCR product was digested with EcoRI and HindIII, and ligated into an EcoRI/HindIII-digested pHSG396 plasmid, a derivative of pUC-type of plasmid harboring a chloramphenicol resistance cassette (Clontech). The ligation product was used to transform Top10 *E. coli*. pHSG396-HldC was purified and used to electroporate $\Delta hldE$ and $\Delta hldE$ expressing *hldA* strains of *S.*

flexneri M90T. All strains were transformed with pCLR7 and constitutively expressed dsRed protein (kind gift from D. Bumann, Biozentrum, Basel, Switzerland).

Preparation of bacterial lysates

Bacteria from overnight cultures were washed in PBS and resuspended at 10^{10} bacteria.mL⁻¹. Once transferred into cryotubes, they underwent through two freeze-thaw cycles in liquid nitrogen. Bacterial debris were removed by centrifugation at 13 000 rpm for 20 min. Lysates were stored at -80 °C.

Cell treatments

Cells were seeded in 96-well plates the day before the experiment. For treatment, they were washed in permeabilization buffer containing 100 mM KCl, 3 mM MgCl₂, 50 mM Hepes, 0.1 mM DTT, 85 mM sucrose, 0.2% BSA and 0.1 mM ATP. They were then incubated with digitonin (5 µg.mL⁻¹) alone or a mixture of digitonin and indicated lysates for 30 min in this same buffer. They were then either fixed with 4% PFA or washed to remove digitonin and incubated for 5.5 h in DMEM supplemented with 1% FCS. For IL-8 measurements, cells were treated with 50 µM monensin at 3 hours.

Infections

For infection experiments with *S. flexneri*, bacteria were used in exponential growth phase. Cells, seeded in 96-well plates, were infected at indicated MOIs in DMEM supplemented with 10 mM Hepes, 2 mM glutamax-1 and 1% FCS. After adding bacteria, plates were centrifuged for 3 min at 1200 rpm and placed at 37 °C for the indicated time periods. Extracellular bacteria were killed by gentamicin (100 µg/mL) and infection was stopped by adding 4% PFA.

IL-8 measurements

The production of IL-8 was measured by immunofluorescence 6 h after cell treatments or 4 h post infection. In order to trap IL-8 intracellularly, monensin (50 μ M) was added after 3 h. After fixation in 4% PFA, cells were stained with an anti-human IL-8 antibody in 0.2% saponin in PBS (BD Pharmingen, San Jose, USA) and an Alexa 647-conjugated secondary antibody (Invitrogen, Carlsbad, USA). IL-8 was quantified by automated image analysis (see below). In parallel, cells were stained with Hoechst (Invitrogen) to visualize nuclei.

Automated microscopy and image analysis

Images were acquired with an ImageXpress Micro (Molecular devices, Sunnyvale, USA). Each data point correspond to triplicate wells and more than 6 images were taken per well. Image analysis was performed using the custom module editor (CME) of MetaXpress. For IL-8, measurements were performed as previously described [3]. Briefly, cell nuclei were identified by the "Autofind blobs" function. Each nucleus was extended by 6 pixels to define the cell mask in which IL-8 signals were quantified. They were detected with the "Keep marked object" function of CME based on minimal/maximal size requirements and intensity threshold. For the oligomerization of TIFA, foci were detected by applying the "Find blobs" function based on size and intensity above background parameters. Cell nuclei were identified with the "Find round objects" module. Cells were defined by extending each nucleus by 10 pixels with the "Grow objects without touching" function and TIFA punctates were quantified within this mask. For infection measurements, nuclei and cells were segmented as above. Bacteria were detected by the "Find blobs" function based on size and intensity above

background parameters. For IL-8, TIFA oligomerization and infection measurements, the fraction of cells respectively containing IL-8, TIFA oligomers or bacteria was extracted, and used for quantification.

Chemical synthesis of β HBP

General Information

Thin Layer Chromatography (TLC) was performed on precoated slides of Silica Gel 60 F₂₅₄ (Merck). Detection was effected when applicable, with UV light, and/or by charring with orcinol (35 mM) in 4 N aqueous (aq.) sulfuric acid and ethanol (95:5). Flash chromatography was performed by elution from columns of Silica Gel 60 (particle size 0.040-0.063 mm). Analytical reverse phase high performance liquid chromatography (RP-HPLC) was performed by elution from a C18 X-Bridge Waters Amide column (4.6x250 mm), using CH₃CN in 5 mM aq. ammonium formate (0→3%) at 0.5 mL.min⁻¹ and detecting at 230 nm. Nuclear Magnetic Resonance (NMR) spectra were recorded at 30 °C for solutions in CDCl₃ or D₂O (400 MHz for ¹H, 100 MHz for ¹³C). Residual CHCl₃ (7.28 ppm for ¹H and 77.0 ppm for ¹³C), and HOD (4.79 ppm) were used as internal references for solutions in CDCl₃ and D₂O, respectively. Proton-signal assignments were made by first-order analysis of the spectra and analysis of 2D ¹H-¹H correlation maps (COSY). Of the two magnetically non-equivalent geminal protons at C-7, the one resonating at lower field is denoted H-7a, and the one at higher field is denoted H-7b. Carbon-signal assignments were supported by 2D ¹³C-¹H correlations maps (HSQC) and analysis of the DEPT and ¹³C spectra. Interchangeable assignments are marked with an asterisk. Electrospray Ionisation-Time Of Flight (ESI-TOF) mass spectra were recorded in the positive-ion mode using a 1:1 CH₃CN/water containing 0.1% formic acid ESI-TOF spectrometer-solution. Anhydrous (anhyd.)

dichloromethane (DCM), 1,2-dichloroethane (DCE), tetrahydrofuran (THF), *N,N*-dimethylformamide (DMF) and toluene (Tol), sold on molecular sieves (MS) were used as such. Reactions requiring anhyd. conditions were performed under an Argon atmosphere.

Synthetic protocols and analytical data.

Phenyl 2,3,4-tri-*O*-benzyl-1-thio- α -D-mannopyranoside (3). To a solution of thioglycoside **2** [18] (15.0 g, 55.1 mmol, 1 equiv.) and imidazole (11.3 g, 165.2 mmol, 3 equiv.) in anhyd. THF (110 mL) was added triisopropylsilyl chloride (TIPSCI) (12.4 mL, 58.8 mmol, 1.05 equiv.) dropwise for 15 min at 0 °C. The reaction mixture was stirred at 0 °C for 20 min before warming up to room temperature for 2.5 h, at which time TLC showed the disappearance of the starting material (DCM/MeOH 8:2, R_f = 0.71) and the presence of a less polar compound (Tol/EtOAc 3:7, R_f = 0.6). An additional amount of TIPSCI (0.08 mL, 0.40 mmol, 0.05 equiv.) was added and after 5 h, the reaction was then quenched with aq. saturated (sat.) NH_4Cl . The phases were separated, the aq. layer was extracted with EtOAc and the organic phases were washed with brine. The organic phase was dried over Na_2SO_4 , filtered, evaporated under reduced pressure and the crude was used as such in the next step.

To a solution of the crude triol (23.6 g, 55.1 mmol, 1 equiv.) and benzyl bromide (BnBr, 23.6 mL, 198.4 mmol, 3.6 equiv.) in anhyd. DMF (306 mL) was added NaH (60% in mineral oil, 13.2 g, 330.6 mmol, 6 equiv.) portionwise for 15 min at 0 °C. The reaction mixture was allowed to warm up to rt overnight, at which time TLC showed the disappearance of the starting material (Tol/EtOAc 3:7, R_f = 0.6) and the presence of a less polar compound (Cyclohexane/EtOAc 9:1, R_f = 0.75). The reaction was then quenched with MeOH at 0 °C and volatiles were evaporated under reduced pressure.

EtOAc and H₂O were added and the aq. layer was extracted with EtOAc. The organic phases were washed successively with aq. sat. NH₄Cl and brine, dried over Na₂SO₄, filtered, evaporated under reduced pressure and the crude was used as such in the next step.

To a solution of the silyl ether (38.5 g, 55.1 mmol, 1 equiv.) in anhyd. THF (306 mL), was added tetrabutylammonium fluoride (TBAF, 110 mL, 1 M in THF, 110 mmol, 2 equiv.) at rt. The reaction mixture was stirred for 3 h at which time TLC showed the disappearance of the starting material (Tol/EtOAc 7:3, R_f = 0.9) and the presence of a more polar compound (Tol/EtOAc 7:3, R_f = 0.6). The reaction was quenched with aq. sat. NH₄Cl. The phases were separated, the aq. layer was extracted with EtOAc and the organic phases were washed successively with aq. sat. NH₄Cl and brine, dried over Na₂SO₄, filtered and evaporated under reduced pressure. The crude was purified by column chromatography on silica gel using Cyclohexane/EtOAc (100:0 to 80:20) to afford alcohol **3** as a yellow oil (28.0 g, 93% over 3 steps). (ESI⁺)-HRMS: Calcd for C₃₃H₃₈NO₅S [M+NH₄]⁺ *m/z*: 560.2471, Found: 560.2485. Other analytical data were similar to those published [13].

Phenyl 2,3,4-tri-O-benzyl-6,7-dideoxy-1-thio- α -D-manno-hept-6-enopyranoside (4) [19]. To a solution of alcohol **3** (15.0 g, 276.4 mmol, 1 equiv.) in anhyd. DCM (2.0 mL) stirred at 0 °C were added successively DMSO (9.8 mL, 138.2 mmol, 5 equiv.), (DIPEA) (14.5 mL, 82.9 mmol, 3 equiv.) and SO₃·Py (13.2 g, 82.9 mmol, 3 equiv.). The reaction mixture was stirred at 0 °C for 15 min and was then allowed to warm up to rt for 4 h. At this time, the reaction was quenched with aq. sat. NH₄Cl. The phases were separated, the aq. layer was extracted with DCM and the organic phases were washed

successively with aq. sat. NH_4Cl and brine, dried over Na_2SO_4 , filtered and evaporated under reduced pressure. The crude was used as such in the next step.

To a suspension of Ph_3PMeBr (52.3 g, 146.5 mmol, 5.3 equiv.) in anhyd. THF (420 mL) stirred at 0 °C was added *n*-BuLi (33.2 mL, 82.9 mmol, 3 equiv.) dropwise. The yellow solution was warmed up to rt for 1 h. Then, a solution of the obtained aldehyde (14.9 g, 276 mmol, 1 equiv.) in anhyd. THF (10 mL) was added dropwise at -78 °C. The reaction mixture was allowed to warm up to rt overnight, at which time TLC showed the disappearance of the starting material (Tol/EtOAc 8:2, R_f = 0.65) and the presence of a less polar compound (Cyclohexane/EtOAc 8:2, R_f = 0.9). The reaction was quenched with aq. sat. NH_4Cl . The phases were separated, the aq. layer was extracted with EtOAc and the organic phases were washed successively with aq. sat. NH_4Cl and brine, dried over Na_2SO_4 , filtered and evaporated under reduced pressure. The crude was purified by column chromatography on silica gel using Tol/EtOAc (100:0 to 95:5) to afford heptoside **4** as a yellow oil (13.6 g, 85%). (ESI⁺)-HRMS: Calcd for $\text{C}_{34}\text{H}_{38}\text{NO}_4\text{S}$ $[\text{M}+\text{NH}_4]^+$ m/z : 556.2521, Found: 556.2527. Other analytical data were similar to those published [13].

2,3,4,6-Tetra-O-benzyl-D-glycero- α -D-manno-heptopyranose (5). To a solution of phenyl 2,3,4,6-tetra-O-benzyl-1-thio-D-glycero- α -D-manno-heptopyranoside [13] (1.0 g, 1.51 mmol, 1 equiv.) in a 9:1 mixture of acetone/ H_2O (15 mL) was added *N*-bromosuccinimide (NBS) (537 mg, 3.02 mmol, 2 equiv.) at 0 °C under Ar and in the dark. The orange mixture was stirred at 0 °C for 15 min then was warmed up to rt. After stirring for 1.5 h, an additional amount of NBS (1 equiv.) was added to the colourless solution at 0 °C. The mixture was stirred overnight at rt, at which time TLC showed the disappearance of the starting material (Tol/EtOAc 8:2, R_f = 0.7) and the presence of a

more polar compound (Tol/EtOAc 8:2, R_f = 0.2). The reaction was quenched with aq. sat. $\text{Na}_2\text{S}_2\text{O}_3$. The phases were separated, the aq. layer was extracted with EtOAc and the organic phases were washed successively with aq. sat. NaHCO_3 and brine, dried over Na_2SO_4 , filtered and evaporated under reduced pressure. The crude was purified by column chromatography on silica gel using Tol/EtOAc (100:0 to 50:50) to afford heptopyranose **5** as a white wax (822 mg, 95%). ^1H NMR (400 MHz, CDCl_3) δ 7.38 – 7.17 (m, 20H, H-Ph), 5.18 (brs, 1H, H-1), 4.92 (d, J = 11.1 Hz, 1H, H- CH_2Ph), 4.77 (d, J = 12.2 Hz, 1H, H- CH_2Ph), 4.71 (d, J = 12.0 Hz, 1H, H- CH_2Ph), 4.64 (s, 2H, H- CH_2Ph), 4.64 (d, J = 12.3 Hz, 1H, H- CH_2Ph), 4.59 (d, J = 11.1 Hz, 1H, H- CH_2Ph), 4.55 (d, J = 12.0 Hz, 1H, H- CH_2Ph), 4.13 (brd, J = 8.6 Hz, 1H, H-5), 4.02 – 3.95 (m, 2H, H-3, H-4), 3.80 (t_{app} , J = 2.2 Hz, 1H, H-2), 3.80 – 3.73 (m, 2H, H-6, H-7a), 3.72 – 3.65 (m, J = 14.3, 7.6 Hz, 1H, H-7b), 3.45 (brs, 1H, OH). ^{13}C NMR (100 MHz, CDCl_3) δ 138.3 (Cq-Ph), 138.3 (Cq-Ph), 138.2 (Cq-Ph), 138.1 (Cq-Ph), 128.6 – 127.6 (20C-Ph), 92.5 (C-1), 79.9 (C-3*), 78.1 (C-6), 75.1 (C-2), 74.7 (C-4*), 74.7 (C- CH_2Ph), 72.8 (C- CH_2Ph), 72.7 (C-5), 72.1 (C- CH_2Ph), 71.8 (C- CH_2Ph), 61.8 (C-7). (ESI⁺)-HRMS: Calcd for $\text{C}_{35}\text{H}_{42}\text{O}_7\text{N}$ $[\text{M}+\text{NH}_4]^+$ m/z : 588.2961, Found: 588.2979.

D-Glycero- β -D-manno-heptopyranose 1,7-bisphosphate (β HBP). To a solution of dibenzyl 2,3,4,6-tetra-O-benzyl-7-di(benzyloxy)phosphoryl-D-glycero- β -D-manno-heptopyranosyl phosphate (70 mg, 64 μmol , 1 equiv.) in a 3:1 mixture of 1,4-dioxane/ H_2O (4 mL) was added $\text{Pd}(\text{OH})_2/\text{C}$ (20%, 20 mg) at rt. The reaction mixture was stirred for 16 h under H_2 (1 atm). The reaction mixture was degassed, filtered on 0.2 μm filter and washed with H_2O . The filtrate was adjusted to pH = 7.0 with Na^+ Dowex 50 and stirred for 30 min. The resin was filtered off and the filtrate was lyophilized to give β HBP (sodium salt form). (ESI⁻)-HRMS: Calcd for $\text{C}_7\text{H}_{15}\text{O}_{13}\text{P}_2$ $[\text{M}-$

H]⁻ *m/z*: 368.9988, Found: 368.9974. *t_r* (RP-HPLC): 3.69 min. Other analytical data corresponded those published [13].

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AUTHOR CONTRIBUTIONS

CA, LAM, JC, DGW and ASD designed research. DGW, ASD, JC and LT performed research and analyzed data. HR performed research. CA, LAM and JC wrote the manuscript.

CONFLICT OF INTEREST

We have no conflict of interest to declare.

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FIGURE LEGENDS

Figure 1: Chemical synthesis of β HBP

Chemical synthesis of β HBP from D-mannose (1): overview.

Figure 2: β HBP fails to induce rapid oligomerization of TIFA

A) β HBP-induced IL-8 production is dose-dependent. HeLa cells were treated for 30 min with digitonin alone, digitonin and wild-type *S. flexneri* lysate or digitonin and β HBP at indicated concentrations. IL-8 production was monitored by immunofluorescence microscopy after 6 h. Hoechst and IL-8 are shown in blue and red, respectively. Scale bar: 20 μ m. **B)** Quantification of IL-8 measurements. Cells were treated as in A and IL-8 was quantified by automated image analysis. Data represent the mean $\pm$ SD of 3 independent experiments. **C)** TIFA oligomerization is visible after 6 h of β HBP treatment. HeLa cells expressing TIFA-GFP were stimulated as in A, fixed and observed by epifluorescence microscopy 6 h post treatment. Scale bar, 20 μ m. **D)** Quantification of TIFA oligomerization as shown in C. Cells with TIFA-GFP oligomers were quantified by automated image analysis. Data correspond to the mean $\pm$ SD of 3 independent experiments. **E)** β HBP fails to induce rapid oligomerization of TIFA-GFP. HeLa cells expressing TIFA-GFP were stimulated as in A, fixed and observed by microscopy 30 min post treatment. Cells with TIFA-GFP oligomers were quantified by automated image analysis. Data correspond to the mean $\pm$ SD of 3 independent experiments. **F)** Kinetics of TIFA-GFP oligomer formation upon β HBP treatment. Cells were stimulated as in A and TIFA-GFP oligomerization was monitored at indicated time points. Cells with TIFA-GFP oligomers were quantified by automated image analysis.

Data correspond to the mean $\pm$ SEM of 4 independent experiments. For comparison between mock and treated conditions (B, D, E, F) or *S. flexneri* lysate and β HBP (F, dashed bar), statistical significance was assessed using unpaired, two-tailed Student's *t*-test, $p^* < 0.05$, $p^{**} < 0.005$, $p^{***} < 0.0005$, non-significant (ns).

Figure 3: Bacterial ADP-heptose synthesis is required for rapid oligomerization of TIFA

A) ADP-heptose biosynthetic pathway. HldA and HldC from *N. meningitidis* are shown in brackets. **B)** Quantification of TIFA-GFP oligomerization at 30 min. HeLa cells were treated with digitonin alone (Mock) or digitonin and a lysate from indicated *S. flexneri* strains for 30 min. Cells with TIFA-GFP oligomers were quantified by automated image analysis. Data were normalized to wt lysate treatment. Data correspond to the mean $\pm$ SD of 3 independent experiments. **C)** Quantification of TIFA-GFP oligomerization at 6 h. HeLa cells were treated with digitonin alone (Mock) or digitonin and a lysate from indicated *S. flexneri* strains for 30 min and washed. The fraction of cells with TIFA-GFP oligomers was quantified 6 h post treatment. Data were normalized to wt lysate treatment. Data correspond to the mean $\pm$ SD of 3 independent experiments. **D)** Quantification of IL-8 production at 6 h. HeLa cells were treated with digitonin alone (Mock) or digitonin and a lysate from indicated *S. flexneri* strains for 30 min and washed. The fraction of IL-8-positive cells was quantified 6 h post treatment. Data were normalized to wt lysate treatment. Data correspond to the mean $\pm$ SD of 3 independent experiments. For comparison between mock and treated conditions (B, C, D), statistical significance was assessed using unpaired, two-tailed Student's *t*-test, $p^* < 0.05$, $p^{**} < 0.005$, $p^{***} < 0.0005$, non-significant (ns).

Figure 4: ADP-heptose is a PAMP of *S. flexneri* infection

A) Representative images of the formation of TIFA oligomers in cells infected with different mutants of *S. flexneri*. HeLa cells expressing TIFA-GFP were left uninfected (mock) or infected for 30 min with indicated strains of *S. flexneri* at MOI 500. pHIdE indicates plasmid-encoded HIdE. Scale bar, 20 μ m. **B)** Infection rate of mutants of the ADP-heptose biosynthetic pathway. HeLa cells were infected at MOI 0.5 with indicated strains and treated with gentamicin. Oligomerization was observed at 4 h post infection with automated microscopy. The fraction of infected cells was analyzed with automated image analysis. Data correspond to the mean $\pm$ SD of 3 independent experiments. **C)** Quantification of TIFA-GFP oligomerization after 30 min of infection with the mutants of the ADP-heptose biosynthetic pathway shown in A. Data correspond to the mean $\pm$ SD of 3 independent experiments. **D)** Quantification of TIFA-GFP oligomerization after 4 h of infection with the *S. flexneri* mutants of the ADP-heptose biosynthetic pathway. Cells were infected for 30 min and treated with gentamicin. Oligomerization was observed at 4 h post infection with automated microscopy. The fraction of infected cells was analyzed with automated image analysis. Data correspond to the mean $\pm$ SD of 3 independent experiments. **E)** Quantification of IL-8 production 4 h post infection with the *S. flexneri* mutants of the ADP-heptose biosynthetic pathway. Cells were infected as in D and treated with monensin. IL-8 was detected by immunofluorescence and quantified by automated image analysis. Data correspond to the mean $\pm$ SD of 3 independent experiments. For comparison between mock and each infected condition (C, D, E) or Δ *hIdE* and double-complemented mutant (dash bar), statistical significance was assessed using unpaired, two-tailed Student's *t*-test, $p^* < 0.05$, $p^{**} < 0.005$, $p^{***} < 0.0005$, non-significant (ns).

Figure 1

Figure 2

A **B** **C** **D** **Figure 3**

Figure 4

Table 1: Table listing the primers used in this study.

Primer	Sequence (5'>3')
<i>S. flexneri</i> $\Delta hldD$ Forward	AGCATCCTTGACCTCTATGTAAAGGGCTGAAGGGATTGGATGTGATGGTATGATTACA GACATTCGTGTCTGAGATTGTCTCTGACTCCATAATTCGAAGGTTACAGTTATGATCTGTGT AGGCTGGAGCTGCTTCG
<i>S. flexneri</i> $\Delta hldD$ Reverse	ACATCGATTATCGCCTGGGGATAGCGCGCCTGGAGCGTGGGATAGAGACTTTGCGACATC ATCATGTCGCCAACCCAAGACGGGCCGATCACCAGTATTTTCATGCAGAGCTCTTATGCCA TATGAATATCCTCCTTAG
<i>N. meningitidis</i> <i>hldC</i> Forward	CCCAAGCTTAATGGTTGACGCTTGGTCTGTCCCT
<i>N. meningitidis</i> <i>hldC</i> Reverse	CCGGAATTCTCATTTTCCGCCCTCTGCCGCACG