Aeromonas hydrophila Flagella Glycosylation: Involvement of a Lipid Carrier

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Abstract

Polar flagellin proteins from Aeromonas hydrophila strain AH-3 (serotype O34) were found to be O-glycosylated with a heterogeneous glycan. Mutants unable to produce WecP or Gne enzymes showed altered motility, and the study of their polar flagellin glycosylation showed that the patterns of glycosylation differed from that observed with wild type polar flagellin. This suggested the involvement of a lipid carrier in glycosylation. A gene coding for an enzyme linking sugar to a lipid carrier was identified in strain AH-3 (WecX) and subsequent mutation abolished completely motility, flagella formation and glycosylation. This is the first report of a lipid carrier involved in flagella O-glycosylation. A molecular model has been proposed. The results obtained suggested that the N-acetylgalactosamines are N-acetylated, linked through the 376 Da sugar to the protein backbone, also in -linkage. Furthermore, deletion of several genes within the putative N-acetylhexosamines is rare. We recently demonstrated that polar and lateral flagellin proteins from Aeromonas hydrophila strain AH-3 (serotype O34) are glycosylated with different carbohydrate moieties [9]. The lateral flagellin is modified at three sites in O-linkage, with a single monosaccharide shown to be a pseudaminic acid derivative of 376 Da. The polar flagellin is modified with a heterogeneous glycan, comprised of a heptasaccharide of, the 376 Da pseudaminic acid type sugar, two hexoses, three N-acetylhexosamines, and an unknown glycan of 102 Da. This glycan chain is linked through the 376 Da sugar to the protein backbone, also in O-linkage. Furthermore, deletion of several genes within the putative N-acetylhexosamines, resulted in abolition of polar and lateral flagella formation by posttranscriptional regulation of the flagellin. In our prior work we were not able to determine the mechanism by which the heptasaccharide chain was formed, or the biological role of flagellin glycosylation. No flagellin until now has been shown to be modified by a glycan built up through Und-P, or showing a complicated glycan structure like the one described previously on A. hydrophila AH-3 [9].

Introduction

Mesophilic Aeromonas are ubiquitous water-borne bacteria, considered opportunistic pathogens of both aquatic and terrestrial animals, some species being associated with gastrointestinal and extraintestinal human diseases [1]. These bacteria constitutively express a single polar flagellum, although about 60% of strains most commonly associated with diarrhoea [2] are also able to express many lateral flagella when grown in viscous environments or on surfaces [3]. Investigations have revealed that both the polar and lateral flagella systems of the mesophilic Aeromonas are involved in adherence to both biotic and abiotic surfaces, as well as in biofilm formation [4]. Flagella motility in Aeromonas, like in other motile bacteria, represents an important advantage in moving towards favourable conditions or in avoiding detrimental environments and it allows the bacterium to successfully compete with other microorganisms [5]. A. hydrophila flagella morphogenesis, as happens in other motile Gram-negative bacteria, is a complex cascade of events that requires coordinated expression of more than 50 genes encoding structural subunits, regulatory proteins and chemo-sensor machinery.

Glycosylation, either N or O linked, is increasingly being observed in bacteria reviewed in [6–8], with the most commonly reported bacterial glycoproteins being flagellins and pili. Campylobacter jejuni is a bacterium with a well-characterized N-glycosylation general pathway with a heptasaccharide built through a lipid carrier undecaprenyl phosphate (Und-P) [7]. In addition to this general protein glycosylation pathway C. jejuni also shows O-linked glycans modifying their flagellins, however with more simple glycans never built through Und-P [7]. A similar situation is observed in Helicobacter pylori [7]. Frequently, the linkage of the glycan to the proteins is an O-linkage of the sugar moiety to the hydroxyl oxygen of serine (Ser) or threonine (Thr) residues of proteins [6]. This is usually catalysed through glycosyltransferases and the need for a lipid carrier for O-glycosylation of proteins is rare. We recently demonstrated that polar and lateral flagellin proteins from Aeromonas hydrophila strain AH-3 (serotype O34) are glycosylated with different carbohydrate moieties [9]. The lateral flagellin is modified at three sites in O-linkage, with a single monosaccharide shown to be a pseudaminic acid derivative of 376 Da. The polar flagellin is modified with a heterogeneous glycan, comprised of a heptasaccharide of, the 376 Da pseudaminic acid type sugar, two hexoses, three N-acetylhexosamines, and an unknown glycan of 102 Da. This glycan chain is linked through the 376 Da sugar to the protein backbone, also in O-linkage. Furthermore, deletion of several genes within the putative pseudaminic acid biosynthetic locus, resulted in abolition of polar and lateral flagella formation by posttranscriptional regulation of the flagellin. In our prior work we were not able to determine the mechanism by which the heptasaccharide chain was formed, or the biological role of flagellin glycosylation. No flagellin until now has been shown to be modified by a glycan built up through Und-P, or showing a complicated glycan structure like the one described previously on A. hydrophila AH-3 [9].
involvement of a lipid carrier in bacterial protein glycosylation has been only described in fimbriae glycosylation from Neisseria and Pseudomonas [10–12].

Surface bacterial polysaccharides, such as O-antigen lipopolysaccharides, capsules or cell wall peptidoglycans, are synthesised in the cytoplasm and subsequently translocated to their location on the bacterial surface. Despite the heterogeneity in the different surface carbohydrates, biosynthesis is commonly initiated with an integral membrane protein that catalyzes the transfer of glucose (Glc)/galactose (Gal)-1-phosphate (WbaP) or N-acetylhexosamine (GlcNAc or GalNAc)-1-phosphate (WecA or WecP, respectively) onto the Und-P [13,14]. These enzymes have multiple transmembrane helices, and the cytosolic loops contribute to form a catalytic site [15]. The predicted topology of these enzymes is variable and linked to their substrate specificity in different cases, like WhaP, WecA or WecP [16–18].

In the current study we demonstrated that a lipid carrier is involved in the heptasaccharide O-glycosylation of polar flagellins in strain AH-3. We showed that the HexNAc from the polar flagellin glycan is N-acetyl galactosamine (GalNAc) or a derivative of it. In addition we begin to shed light on the biological role of flagellin glycosylation with data linking flagellin glycosylation to motility.

Results

Our previous work has shown A. hydrophila AH-3 flagellins to be post-translationally modified with glycan moieties [9]. The lateral flagellin was observed to be glycosylated with a unique 376 Da sugar residue, a putative pseudaminic acid derivative. In contrast, the polar flagellin was modified with a heptasaccharide glycan, comprised of three N-acetylhexosamines additionally modified by variable numbers of phosphate groups and methyl groups, two hexoses and two unknown glycans of 376 Da (putative pseudaminic acid derivative) and 102 Da. This gave a monosaccharide mass sequence of 376-162-162-203-296-376-102 Da.

The aim of the current study is to understand the mechanisms of the differential glycosylation of polar and lateral flagellins using two previously isolated motility deficient mutants from this strain: a wecP mutant (AH-3ΔWecP) and a gne mutant (AH-2767). WecP is the enzyme codified by a gene in the O34-antigen LPS cluster linking UDP-GalNAc to the Und-P [6], while Gne is the enzyme able to 4-epimerize UDP-GlcNAc to UDP-GalNAc [19] codified outside the O34-antigen LPS cluster by a gene alone between non related genes codifying for a ferredoxin oxidoreductase and a protein-disulfide isomerase. We have previously shown that both mutants are devoid of the O34-antigen LPS [18,19].

A. hydrophila AH-3 wecP Mutant (AH-3ΔWecP)

This mutant is able to produce both polar and lateral flagella. No differences were observed between the wild type strain and the AH-3ΔWecP mutant in lateral flagellin production grown on solid or semisolid media when examined by EM. However, the AH-3ΔWecP mutant showed a reduction in motility compared to the wild type strain when assayed on swim agar plates (Figure 1). This motility reduction was confirmed by light microscopy observations in liquid media. This fact prompted us to purify the polar flagellum and compare it to the wild type. A reduction in the molecular weight of AH-3ΔWecP polar flagellins, was observed by SDS-PAGE, as shown in Figure 2A. The amount of polar flagellin obtained from 11 culture of the mutant versus the same amount of wild type growth is reduced in approximately 50% judged by protein concentration analysis.

Figure 1. Motility phenotypes exhibited in swim (0.25%) agar by A. hydrophila AH-3 (A), AH-2767 gne mutant (B), AH-3ΔWecP mutant (C), AH-2767+ pACYC-Gne (D), and AH-3ΔWecP pBAD-WecP (E).

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A full recovery of all the reduced characteristics observed in the AH-3ΔWecP mutant versus the wild type (motility or molecular weight reduction and variation of polar flagellins) were achieved when we introduced pBAD33-WecPAh plasmid [18] (Figure 1 and 2). No changes were observed when we introduced the plasmid vector alone in the same mutant.

Mass Spectrometry Analysis of Polar Flagellin from an A. hydrophila AH-3 WecP Mutant

Protein mass analysis was carried out by using electrospray ionisation mass spectrometry on polar flagellins isolated from the AH-3ΔWecP mutant. Although poor ionisation prevented spectral deconvolution, and the observed protein masses were not determined.

MS/MS analyses of tryptic peptides from polar flagellins isolated from the AH-3ΔWecP mutant showed the same peptides to be modified with glycan as the wild type strain (Table 1), however differences in the glycan modification were observed. To highlight the observed differences in glycopeptide profiles observed between AH-3ΔWecP mutant and wild type, Figure 3 shows alignment of LC-MS spectra from tryptic digests of polar flagellin isolated from wild type (Figure 3a) and AH-3ΔWecP mutant (Figure 3b) at 13.1–13.8 minutes (elution time for the 94–109 TLAQQSANGSN NTDDR glycopeptide). In our previous study, we reported the glycopeptides to be observed in LC-MS spectra as clusters of multiple charged ions. These clusters of ions corresponded to the glycan heterogeneity. In Figure 3a, these clusters are denoted 1–4 and correspond to different amounts of
We purified the polar flagellum from the mutant and compared it with the flagellum of the wild type strain. The amount of polar flagellin obtained from 1l culture of the mutant versus the same amount of wild type growth is reduced in approximately 70% judged by densitometry of the gel bands. Figure 4 shows that there is a large variability in the molecular weight of flagellins in the mutant, with bands corresponding to polar flagellin migrating to approximate molecular weights between 44 and 50 kDa. This is in comparison with polar flagellin from wild type, which migrates to three discrete bands between 43 and 45 kDa. A full recovery of the wild type polar flagellin profile and protein abundance was observed when the mutant was complemented by pACYC-GNE plasmid [19] and wild type polar flagellin restored, as could be observed in Figure 4.

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**Mass Spectrometry Analyses of Polar Flagellin from *A. hydrophila AH-3 gne Mutant***

Protein mass analysis was carried out on polar flagellin isolated from the AH-2767 mutant (AH-3 gne mutant). The MS profile showed a complex ion envelope with poor ionization, which could not be resolved to give observed intact protein masses. However, the protein MS spectrum did show an intense ion at m/z 377 likely corresponding the glycan oxonium ion of the 376 Da pseudaminic acid derivative (data not shown). MS/MS analyses of tryptic peptides from polar flagellin isolated from the mutant showed four of the five glycopeptides previously observed in the wild type [9]. However, the predominant modification of these peptides in the mutant strain was observed to be a single 376 Da moiety, compared with the wild type flagellin where the major modification was a complex and variable heptasaccharide glycan chain. The MS/MS spectrum of the 94–109 TLAQQSANGSNNTDDR glycopeptide (sequence present in both FlaA and FlaB), with a 376 Da glycan modification is shown in Figure 5. Figure 3a and c, show the LC-MS spectra of flagellin tryptic digests from wild type and AH-2767 mutant at 13.1–13.8 minutes (elution time for the 94–109 TLAQQSANGSNNTDDR glycopeptide). Comparison of
Figure 3. nLC-MS alignment of the AH-3 polar flagellin FlaA/FlaB glycopeptide. The 1691 Da FlaA/FlaB peptide TLAQQSANGSNNTDDR (eluting between 13.1 and 13.8 minutes) was modified by a heptasaccharide glycan with HexNAc residues additionally variably modified by 0–2 phosphate and 0–2 methyl groups [9]. These additional variable modifications resulted in four major glycopeptide ion cluster regions, as indicated below each spectrum. The cluster number corresponds to the number of additional phosphate groups modifying the heptasaccharide, and peaks within that cluster possess a variable number of additional methyl groups. Only the most prominent peak within each cluster is annotated. a) Wild type polar flagellin. Glycoforms of the peptide modified with the heptasaccharide were more abundant, compared with the peptide modified with only a single 376 Da (this form of the glycopeptide is indicated in the Figure with an arrow at m/z 1034.4). The glycopeptide peak intensity increased with the addition of each phosphate group. b) AH-3ΔwecP mutant polar flagellin. In contrast to wild type, the predominant modification in AH-3ΔwecP was a single 376 Da sugar. The glycopeptide clusters resulting from the heptasaccharide glycan and additional variable modification were observed, but with decreased intensity as compared to the monosaccharide modification. c) AH-2767 (gne mutant) polar flagellin. A decrease in the complexity of glycan modification was observed. The most abundant modification observed was the single 376 Da glycan. Glycopeptide clusters were not observed in the same manner when compared to wild type and the longer chain modifying glycans were not observed. In addition, the unmodified peptide was also observed at m/z 846.4. This was not detectable in wild type, Figure legend: solid pentagon = 376 Da sugar; ● = Hexose; □ = HexNAc; ▲ = 102 Da unknown moiety; ●● = phosphorylation; ●●● = methylation; * = truncated peptide. doi:10.1371/journal.pone.0089630.g003
clusters for the polar flagellin from the mutant, compared with
respectively). Table 1 shows the relative intensities of the ion
m/z 1196 and 1115 (376 monosaccharide with two or one hexoses,
m/z charged ions at 

AH-2767 mutant showed the most intense glycopeptide peak of
the wild type data, the relative intensity of the peaks it can be observed that
intermediate glycan chain lengths [Pse-162 or Pse-162–162
MTA FTISGIASSTK, which is unique to FlaB. As
previously noted, the wild type polar flagellin (Figure 6a) shows a complex pattern of glycosylation, with glycan isoforms with varying degrees of phosphorylation and methylation, denoted in Figure 6a as clusters 1–4. Again, reduced complexity of glycosylation was observed in the polar flagellin AH-2767 mutant (Figure 6b), evidenced by the absence of the glycopeptide ion clusters. It is interesting to note the in the AH-2767 mutant, the unmodified form of the FlaB peptide (m/z 751.5) and the peptide modified only by the 376 Da moiety (m/z 939.4), were not observed at high levels, as shown in Figure 6 and Table 1. As previously indicated, AH-3ΔWecP polar flagellin showed reduced
levels of full length sugar chains in comparison with the same flagellin from the wild type (Figure 6c).

Isolated polar flagellins from AH-2767 mutant were able to react with specific antiserum against GlcNAc while no reaction was observed with isolated polar flagellins from wild type strain or AH-3ΔWecP (Figure 4D).

A. hydrophila AH-3 Mutants Lacking Only O34-antigen LPS
Both mutants (AH-2767 and AH-3ΔwecP) lack the O34-antigen LPS characteristic of this strain [20], and others have suggested the possible involvement of O-antigen LPS in flagella glycosylation in other bacteria [21]. We therefore investigated the possible role of O34-antigen LPS in A. hydrophila AH-3 flagella glycosylation.

Previously isolated mutants (AH-3ΔwaaL) [22] and AH-3ΔManC constructed in this study were also devoid of the O34-antigen LPS. AH-3ΔwaaL is able to produce O34-antigen LPS but lacks the O-antigen ligase (WaaL) that links it to the LPS-core. We have previously published a study using an insertional mutant in manC with possible polar effects in wecP of the cluster who34 [23]. Then, we generated in this study an in frame manC mutant in order to avoid polarity effects. AH-3ΔManC lacks the mannose-1-phosphate guanylyltransferase, required to produce GDP-Man, a biosynthetic precursor for polysaccharides. The AH-3ΔManC mutant is unable to produce O34-antigen LPS according to their LPS gel profile in SDS-PAGE (Figure 7A). No relevant differences in motility were observed in both mutants versus the wild type strain or by EM both mutants showed a similar polar flagellum as the wild type strain (Figure 7). Purified polar flagellins from both mutants showed the same molecular weight profile as wild type polar flagellins (Figure 8).
A Second Lipid Carrier on Strain AH-3

The use of several primers able to amplify different Und-P domains in different Aeromonas strains allowed us to identify a second enzyme able to link sugars to a lipid carrier (named WecX: GenBank KF734980) in strain AH-3 (see Materials and Methods). This identification allowed us to obtain defined wecX insertion mutants devoid of it. The gene codifying for the enzyme WecX is located between a gene codifying for a signal transduction histidine kinase upstream and a two downstream genes codifying for glycosyltransferases which does not seem to be involved in LPS but maybe in exopolysaccharide production. Unfortunately, the complete genome of strain AH-3 is not currently available.

The single mutant AH-3::WecX was non motile (Figure 9) and unable to produce polar and lateral flagella by EM (Figure 10). The similar situation was observed for the double mutant AH-3::WecP::WecX. No changes could be observed in their LPS profiles in comparison with their respective wild type in SDS-PAGE [18] (Figure 9). The complementation of the double mutant AH-3::WecP::WecX with pBAD-WecX was unable to change their LPS profile in gels (Figure 9). Because some unattached portions of flagella could be observed in the EM background of the mutants, we tried to purify the polar flagellin from the mutants by cesium chloride. A minor protein band was visualized on 1D-PAGE, obtained using 10 times more growth volume for flagellin purification from the mutants than the amount used for the wild type strain. Tandem mass spectrometry analyses of tryptic digests of the band showed it harbor low amounts of polar flagellin protein, with no detectable glycopeptides. Mass spectrometry analyses of the tryptic digests of this mutant gave 20 percent sequence coverage. De novo sequencing showed no evidence of previously observed glycopeptides, and no evidence of post translational modification of any peptides. Polar flagellin is
synthesized in the cytoplasm fraction by these _wecX_ mutants as the wild type, but very few if any are properly assembled in the membrane fraction when compared to the wild type (Figure 10). When we introduced the pBAD-WecX into the single mutant AH-3:WecX or the double mutant AH-3ΔWecP:WecX, and we grow the strains under inducing conditions (+0.2% arabinose), they recover the full motility (Figure 9) and the flagella production by EM was the same as observed with the wild type strain (Figure 10). The complemented strains showed flagellin in the membrane fraction and in purified flagella (Figure 10) with the same MW as the wild type. No changes in motility or flagella production by EM could be observed when we introduced the plasmid vector alone (pBAD33) and the mutant strains grow under inducing conditions.

The mutants that lack O34-antigen LPS showed a reduced virulence when tested in several animals by intraperitoneal injection [19,23]. When we tested the AH-3ΔWecX or mutants unable to produce flagella from the same strain no such reduction of the virulence could be observed in the same animal models.

**Discussion**

The list of bacterial species that modify their flagellin proteins with diverse glycans continues to grow and was recently reviewed [6,24], however an understanding of the biological role of glycosylation remains limited. In the current study, we show that flagellin glycosylation impacts the ability of _A. hydrophila_ AH-3 to form full flagellar filaments and motility.

_WecP_ is the integral membrane protein that catalyzes the transfer of GalNAc-1-phosphate onto Und-P and the major modification for AH-3ΔWecP mutant was the loss of O34-antigen LPS [18]. Several fully sequenced strains of _Aeromonas_ encode genes for more than enzyme linking sugars to a lipid carrier. For example, strain ATCC7966T [30] constitutively expresses polar flagellin and lacks an inducible lateral flagella [25], and showed several putative enzymes able to link sugars to lipid carriers in addition to _WecP_.

A second enzyme able to link sugars to a lipid carrier on strain AH-3 was identified in this study, denoted WecX, that we show is responsible for the heptasaccharide glycosylation of the polar flagellin without modifying the O34-antigen LPS. This conclusion is in agreement with the data of no changes in the LPS and no production of polar flagellum by EM, no motility, or the non-glycosylated flagellin observed in the mutants (AH-3:WecX and AH-3ΔWecP:WecX). All these negative traits were recovered by the introduction of the AH-3 _wecX_. To our knowledge this is the first report of a lipid carrier involved in flagella O-glycosylation. It has been recently described that other external organelles like fimbrae showed O-glycosylation with the recruitment of a lipid carrier [10,26]. However, while fimbrae are inducible external organelles, the _A. hydrophila_ AH-3 polar flagellum is constitutively expressed [27]. WecX protein sequence showed clear undecaprenyl-phosphate glucose phosphotransferase domain. Computer programs, such as TMHMM 2.0 (Prediction of transmembrane helices in proteins from the Center for Biological Sequence Analysis, Denmark; http://www.cbs.dtu.dk/services/TMHMM/), have been used to predict transmembrane helices and membrane topology models for proteins involved in the transfer of sugar phosphate residues to undecaprenyl phosphate (Und-P) such as WecA [17], WecP [18] and WbaP [16]. The same computer program was used to model the _A. hydrophila_ AH-3 WecX molecular topology revealing that not feed in any of the known models, but two clear transmembrane regions could be observed at the beginning (first 40 amino acid residues) and the end of the protein (the last 25 amino acid residues). It is important to notice that none of these transferase-catalyzed reactions to the membrane-associated polyoliprenyl phosphate acceptor (Und-P) until now described [15] uses a sugar nucleotide related to _Pse_ or a _Pse_ derivative.

The results obtained in this current work allow us to suggest the hierarchy of polar flagellin O-glycosylation. While pseudaminic acid (Pse) biosynthetic mutants are unable to form flagella because they are unable to produce properly flagellin in the cytoplasm [9], the WecX mutant (AH-3:WecX) is able to do it. It seems unable to produce flagella probably because could not locate the heptasaccharide

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**Figure 5. nLC-MS/MS AH-2767 _one mutant polar flagellin glycopeptide spectrum.** nLC-MS/MS spectrum of the doubly charged ion at m/z 1034.4 eluting between 13.1 and 13.8 minutes. The peptide sequence was identified by y and b ion series as **“TLAQ55NSNNTDDR”** which corresponds to both the FlaA and FlaB sequences. The unmodified singly charged peptide ion can be seen at m/z 1691.7, giving a mass excess of 376 Da (pseudaminic acid derivative). The prominent peaks at m/z 377 and 359 correspond to the glycan oxonium ion and its dehydrated form, respectively.

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charide outside the membrane, absence of the lipid carrier. This second lipid carrier WecX is unable to restore the O:34 antigen LPS in the double mutant AH-3WecP:WecX, thus indicating that is used only in flagella glycosylation.

Thus, the first initial sugar linked to the flagellin is Pse-like, with the hexasaccharide [(162-162-203-297-376-102 Da), 2 Hex, 3HexNAc unsubstituted or substituted with phosphate and methyl groups, and unknown glycan 102 Da], being this hexasaccharide produced by the lipid carrier WecX. We proposed the following molecular model (Figure 11). We suggest that WecX is able to link CMP-Pse-like to Und-P, and several different glycosyltransferases are able to form sequentially the hexasaccharide linked to Und-P, being first the glycosyltransferases that add Hex, then the glycosyltransferases that add HexNAc, and finally a putative transferase for the unknown glycan of 102 Da. Then, one enzyme that we named OTase like, transfers in an O-glycosylation basis these heptasaccharide to the threonine/serine amino acids of the polar flagellin. Once the flagellins molecules are glycosylated could be transported to produce the polar flagellum.

The A. hydrophila AH-2767 gne mutant showed a great reduction in the phenotypic traits (motility in swimming plates, polar flagellum by EM or molecular weight size variation) compared with the wild type strain. This is in agreement with the protein glycosylation data, which show the polar flagellin in this mutant to be modified with Pse or truncated glycan chains with one or two hexoses major moieties and only a little of the full length heptasaccharide moiety. The AH-2767 gne mutant is unable to form UPDGalNAc from UDPGlcNAc [19], suggesting that in the wild type, the HexNAc observed in the minimal amount of

Figure 6. nLC-MS alignment of AH-3 polar flagellin FlaB glycopeptide. The 1501 Da FlaB peptide MTSAPTSGIASSTK (eluting between 17.9 and 18.9 minutes) was modified by the same heptasaccharide glycan with HexNAc residues additionally variably modified by phosphate and methyl groups [9]. These additional variable modifications resulted in four major glycopeptide ion cluster regions, as indicated below each spectrum. The cluster number corresponds to the number of additional phosphate groups modifying the heptasaccharide, and peaks within that cluster possess a variable number of additional methyl groups. Only the most prominent peak within each cluster is annotated. a) Wild type polar flagellen. The heptasaccharide modification was the most abundant form of modification, with only minor amounts of peptide harbouring the 376 Da sugar were observed. The glycopeptide peak intensity increases with the addition of each phosphate group to the glycans. b) AH-2767 gne mutant polar flagellin. In this mutant, the FlaB peptide showed only minor modification with intermediate glycan chains. The unmodified peptide and peptide modified only with the 376 Da sugar were also not observed at high levels. c) AH-3ΔWecP mutant polar flagellin. In contrast to wild type, the predominant modification in AH-3ΔWecP was a single 376 Da sugar. Clustering resulting from the heptasaccharide and additional variable modification similar to that observed in wild type polar flagellin was observed, but with decreased intensity as compared to the monosaccharide modification. Figure legend: solid pentagon = 376 Da sugar; ● = hexose; ■ = HexNAc; ▲ = 102 Da unknown moiety; ● = phosphorylation; ■ = methylation; * = truncated peptide.
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heptasaccharide formed was GalNAc or a derivative of it. We could detect presence of GlcNAc in isolated AH-2767 gne mutant polar flagellins, which is not detected in the corresponding from the wild type, suggesting that this monosaccharide can be substituted into the glycan chain. This fact explains why the major glycosylation in the fagellin of this mutant is Pse, Pse-Hex or Pse-Hex-Hex. Although substitution of GlcNAc did not result in formation of wild type flagellin heptasaccharide glycosylation, a minimum amount could be detected in the mutant perhaps for the permissively of the corresponding glycosyl transferases. For instance C. jejuni PglA showed a relaxed specificity accepting several substrates in formation of the heptasaccharide for N-glycosylation [28,29].

The AH-3ΔWecP mutant showed a slight reduction in motility or in the molecular weight of polar flagellins when compared to the wild type strain. In agreement, the major modification for AH-3ΔWecP polar flagellin heptasaccharide glycosylation, a minimum amount could be detected in the mutant perhaps for the permissively of the corresponding glycosyl transferases. For instance C. jejuni PglA showed a relaxed specificity accepting several substrates in formation of the heptasaccharide for N-glycosylation [28,29].

GalNAc in addition we have begun to shed light on the role of flagellum glycosylation in flagellum formation and motility.
Materials and Methods

Bacterial Strains, Plasmids and Growth Conditions

The bacterial strains and plasmids used in this study are listed in Table 2. E. coli strains were grown on Luria-Bertani (LB) Miller broth and LB Miller agar at 37°C, while Aeromonas strains were grown either in tryptic soy broth (TSB) or agar (TSA) at 30°C. When indicated kanamycin (50 μg/ml), rifampicin (100 μg/ml), spectinomycin (50 μg/ml), tetracycline (20 μg/ml) and chloramphenicol (25 μg/ml) were added to the media.

DNA Techniques

DNA manipulations were carried out essentially according to standard procedures [33]. DNA restriction endonucleases and E. coli DNA polymerase Klenow fragment were obtained from Promega. T4 DNA ligase and alkaline phosphatase were obtained from Invitrogen and GE Healthcare, respectively. PCR was performed using BioTaq DNA polymerase (Ecogen) in a Gene Amplifier PCR System 2400 Perkin Elmer Thermal Cycler. Plasmid DNA for sequencing was isolated by Qiagen plasmid purification kit (Qiagen, Inc. Ltd.) as recommended by the suppliers. Double-strand DNA sequencing was performed by using the Sanger dideoxy-chain termination method [34] with the BigDye Terminator v3.1 cycle sequencing kit (Applied Biosystem). Custom-designed primers used for DNA sequencing were purchased from Sigma-Aldrich. The DNA sequences were compared with those available in the GenBank and EMBL databases at the National Center for Biotechnology Information (NCBI) [35]. The Terminator search program in the GCG Wisconsin package was used to search for factor independent transcriptional terminators. The Neural Network Promoter Prediction, PromScan [36] and PRODORIC [37] were used to search promoter sequences.

Construction of Defined Mutants

The chromosomal in-frame manC deletion mutant A. hydrophila AH-3ΔmanC was constructed by allelic exchange as described by Milton et al. [38]. Briefly, upstream (fragment AB) and downstream (fragment CD) of manC gene was independently amplified using two sets of asymmetric PCRs. Primer pairs A-ManC (5’-CGGCCTTATTGTCAAGGGGACCTG-3’) and B-

Figure 10. A/Electron microscopy of whole cells from A. hydrophila AH-3::WecX mutant (1) and the mutant complemented with pBAD-WecX (2) stained according to Experimental Procedures. Bar represents 1μ. B/Western blots using specific antiserum against purified polar flagellins. Molecular weights are indicated. Cytoplasmic fractions (C), Whole membrane fractions (M), and partially purified polar flagellins (F), the fractions were separated as indicated in Materials and Methods section. Strains: 1, A. hydrophila AH-3 (wild type); 2, AH-3::WecX mutant, and 3 the mutant complemented with pBAD33-WecX.

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Figure 11. Molecular model proposed for the polar flagellum glycosylation. We suggest that WecX is able to link CMP-Pse-like to Und-P, and several different glycosyltransferases are able to form sequentially the heptasaccharide linked to Und-P, being first the glycosyltransferases that add Hex (two of them), then the glycosyltransferases that add GalNAc or a derive of it (three of them), and finally a putative transferase for the unknown glycan of 102 Da. Then an enzyme, that we named OTase like, transfers in O-glycosylation basis these heptasaccharide to the threonine/serine amino acids of the polar flagellin. Once the flagellin molecules are glycosylated could be transported to produce the polar flagellum. Gne is the enzyme that converts UDP-GlcNAc in UDP-GalNAc, and their lack jeopardizes the UDP-GalNAc formation.

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broth and LB Miller agar at 37°C, while Aeromonas strains were grown either in tryptic soy broth (TSB) or agar (TSA) at 30°C. When indicated kanamycin (50 μg/ml), rifampicin (100 μg/ml), spectinomycin (50 μg/ml), tetracycline (20 μg/ml) and chloramphenicol (25 μg/ml) were added to the media.

DNA Techniques

DNA manipulations were carried out essentially according to standard procedures [33]. DNA restriction endonucleases and E. coli DNA polymerase Klenow fragment were obtained from Promega. T4 DNA ligase and alkaline phosphatase were obtained from Invitrogen and GE Healthcare, respectively. PCR was performed using BioTaq DNA polymerase (Ecogen) in a Gene Amplifier PCR System 2400 Perkin Elmer Thermal Cycler. Plasmid DNA for sequencing was isolated by Qiagen plasmid purification kit (Qiagen, Inc. Ltd.) as recommended by the suppliers. Double-strand DNA sequencing was performed by using the Sanger dideoxy-chain termination method [34] with the BigDye Terminator v3.1 cycle sequencing kit (Applied Biosystem). Custom-designed primers used for DNA sequencing were purchased from Sigma-Aldrich. The DNA sequences were compared with those available in the GenBank and EMBL databases at the National Center for Biotechnology Information (NCBI) [35]. The Terminator search program in the GCG Wisconsin package was used to search for factor independent transcriptional terminators. The Neural Network Promoter Prediction, PromScan [36] and PRODORIC [37] were used to search promoter sequences.

Construction of Defined Mutants

The chromosomal in-frame manC deletion mutant A. hydrophila AH-3ΔmanC was constructed by allelic exchange as described by Milton et al. [38]. Briefly, upstream (fragment AB) and downstream (fragment CD) of manC gene was independently amplified using two sets of asymmetric PCRs. Primer pairs A-ManC (5’-CGGCCTTATTGTCAAGGGGACCTG-3’) and B-

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ManC (5'-CCCATCCAC TAACTTAAACACCTTTG-GAAAATGCTCACG-3') and C-ManC (5'-TGTTTTAAAGT- TAGTTGATGGGATTGAAATACGCTCCGGTA-3') and D- ManC (5'-CCGGGATCTCTAATGAGGACGTCTCGCTG-3') amplify DNA fragments of 830 (Man-AB) and 798 (Man-CD) bp for manC in-frame deletion. DNA fragments AB and CD were annealed at the overlapping regions (underlined letters in primers A and D). The fusion products were purified, annealed at the overlapping regions (underlined letters in primers A and D), ligated into pBAD33 digested and phosphatase-treated pBAD33 vector independently site is double-underlined in primer A and D), ligated into Bgl II digested and phosphatase-treated pBAD33 vector independently [38], electroporated into E. coli MC1061 (Δpir), and plated on chloramphenicol plates at 30°C to obtain pDM4-ManC plasmid. Plasmid pDM4-ManC with mutated gene was transferred into A. hydrophila AH-405 by triparental mating using the pBAD33 vector, or digested with Sma I and ligated into pBAD33 vector. This plasmid was used to construct the pBAD-ManC plasmid. Plasmid pBAD-ManC and pBAD-WecX containing the complete manC and wexX of A. hydrophila AH-3 under the arabinose promoter (pBAD) on pBAD33 [40] were obtained, respectively. Oligonucleotides 5'-TCCCCCGGAAGCGATGCTGCTGAATC-3' and 5'-CCCAGCTTTAACGCAACCACATAGCTG-3' generated a band of 1687 bp containing the manC gene. Oligonucleotides 5'-GTCTCTCCCTGCTTACC-3' resulting in an ampiclon of 586 bp that was subcloned in pGEM. After EcoRI digestion the plasmid pGEMT-wexX was treated with E. coli DNA polymerase I (Klenow fragment) to create blunt ends in order to ligate the band to pCM100 (suicide plasmid, Δpir dependent, [40]) to obtain plasmid pCM-WecX. This plasmid was used to isolate wexX deficient mutants from AH-3 and AH-3ΔWecP (AH-3:WecX and AH-3ΔWecP:WecX, respectively) by a single recombination event according to previously described procedures [39].

**Plasmid Constructions**

Plasmid pBAD-ManC and pBAD-WecX containing the complete manC and wexX of A. hydrophila AH-3 under the arabinose promoter (pBAD) on pBAD33 [40] were obtained, respectively. Oligonucleotides 5'-TCCCCCGGAAGCGATGCTGCTGAATC-3' and 5'-CCCAGCTTTAACGCAACCACATAGCTG-3' generated a band of 1687 bp containing the manC gene. Oligonucleotides 5'-GTCTCTCCCTGCTTACC-3' resulting in an ampiclon of 586 bp that was subcloned in pGEM. After EcoRI digestion the plasmid pGEMT-wexX was treated with E. coli DNA polymerase I (Klenow fragment) to create blunt ends in order to ligate the band to pCM100 (suicide plasmid, Δpir dependent, [40]) to obtain plasmid pCM-WecX. This plasmid was used to isolate wexX deficient mutants from AH-3 and AH-3ΔWecP (AH-3:WecX and AH-3ΔWecP:WecX, respectively) by a single recombination event according to previously described procedures [39].

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Flagella Purification

A. hydrophila AH-3 was grown in TSB for the polar flagellum purification. For the isolation of lateral flagella the strains were grown in TSA and plates washed with 100 mM Tris (pH = 7.8). Cells were in both cases collected by centrifugation at 5000 x g, and suspended in the same Tris buffer. Flagella were removed from the cells by shearing in a vortex with a glass bar for 3–4 minutes, and then passing repetitively (minimum six times) through a syringe. Cells were removed by centrifugation at 8000 x g for 30 minutes, and the supernatant centrifuged at 18000 x g for 20 minutes. From the remaining supernatant the flagella were pelleted by ultracentrifugation at 100 000 x g for 48 hours. The band containing the flagella was collected, the cesium chloride removed by extensive dialysis against the same buffer (100mM Tris 2mM EDTA). Purified flagella were analyzed by SDS-PAGE or by glycosylation chemical studies.

Cell Fractions

A. hydrophila cells grown in tryptic soy broth at 25°C were harvested by centrifugation (5000 x g), washed with 20 mM MgCl2 in 100 mM Tris buffer (pH = 8.0), and resuspended in the same cold buffer, and French press cell lysis was performed. The lysates were centrifuged at 5000 x g to remove unbroken cells. After centrifugation at 4°C for 1 h at 115,000 x g, the cytoplasmic fraction remained in the supernatant, whereas the whole membrane fraction was retained in the pellet.

Immunological Methods

Western blot of purified flagella was performed as briefly described. After SDS-PAGE, immunoblotting was carried out by transfer to polyvinylidene fluoride membranes (Millipore Corp., Bedford, Mass.) at 1.3 A for 1 h in the buffer of Towbin et al. [42]. The membranes were then incubated sequentially with 1% bovine serum albumin, specific anti-polar flagellins polyclonal serum [27] (1:2000), alkaline phosphatase-labeled goat anti-rabbit immunoglobulin G, and 5-bromo-4-chloroindolylphosphate disodium-nitroblue tetrazolium. Incubations were carried out for 1 h, and washing steps with 0.05% Tween 20 in phosphate-buffered saline were included after each incubation step.

Motility Assays (Swarming and Swimming)

Freshly grown bacterial colonies were transferred with a sterile toothpick into the centre of swarm agar (1% tryptone, 0.5% NaCl, 0.5% agar) or swim agar (1% tryptone, 0.5% NaCl, 0.25% agar). The plates were incubated face up for 16–24 h at 25°C and motility was assessed by examining the migration of bacteria through the agar from the centre towards the periphery of the plate. Moreover, swimming motility was assessed by light microscopy observations in liquid media.

Transmission Electron Microscopy (TEM)

Bacterial suspensions were placed on Formvar-coated grids and negative stained with a 2% solution of uranyl acetate pH 4.1. Preparations were observed on a Hitachi 600 transmission electron microscope.

Electrospray Liquid Chromatography Mass Spectrometry

Mass spectrometry studies of intact flagellin proteins were carried out using 1–5 μg of protein, exactly as described previously [9]. Glycopeptides were mapped using tandem mass spectrometry methods, exactly as reported in our previous work [9]. Briefly, purified flagellins (50 to 200 ng) were digested overnight with trypsin (Promega, Madison, WI) at a ratio of 30:1 (protein/enzyme, vol/vol) in 50 mM ammonium bicarbonate at 37°C. Protein digests were analyzed by nano-liquid chromatography MS/MS (nLC-MS/MS) using a Q-TOF Ultima Hybrid quadrupole-tandem time of flight MS (Waters, Milford, MA) [44]. Front end peptide separation was carried out using online nanoAcquity ultrahigh-pressure liquid chromatography system (Waters, Milford, MA) using previously detailed setup [9]. MS/MS spectra were acquired using data dependent mode or in targeted analyses on doubly, triply, and quadruply charged ions in collision induced dissociation (CID) mode. Unmodified peptides were identified using Mascot (Matrix Science, London, UK) as described in our earlier work [43]. Glycopeptide MS/MS spectra were de novo sequenced, with peptide type y and b ions highlighted for confirmation of the peptide sequence.

Statistical Analysis

Results are expressed as the mean ± standard deviation (SD) of three to four experiments. Student’s t test was used to compare mean values. Differences were considered significant when P values were <0.05.

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Author Contributions

Conceived and designed the experiments: SM KMF SMT MW RM JMT. Performed the experiments: SM KMF SMT MW RM JMT. Analyzed the data: SM KMF SMT MW RM JMT. Contributed reagents/materials/analysis tools: SM KMF SMT MW RM JMT. Wrote the paper: SM KMF SMT MW RM JMT.

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