Genetic and functional diversity of *Pseudomonas aeruginosa* lipopolysaccharide

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Lipopolysaccharide (LPS) is an integral component of the *Pseudomonas aeruginosa* cell envelope, occupying the outer leaflet of the outer membrane in this Gram-negative opportunistic pathogen. It is important for bacterium–host interactions and has been shown to be a major virulence factor for this organism. Structurally, *P. aeruginosa* LPS is composed of three domains, namely, lipid A, core oligosaccharide, and the distal O antigen (O-Ag). Most *P. aeruginosa* strains produce two distinct forms of O-Ag, one a homopolymer of d-rhamnose that is a common polysaccharide antigen (CPA, formerly termed A band), and the other a heteropolymers of three to five distinct (and often unique dideoxy) sugars in its repeat units, known as O-specific antigen (OSA, formerly termed B band). Compositional differences in the O units among the OSA from different strains form the basis of the International Antigenic Typing Scheme for classification via serotyping of different strains of *P. aeruginosa*. The focus of this review is to provide state-of-the-art knowledge on the genetic and resultant functional diversity of LPS produced by *P. aeruginosa*. The underlying factors contributing to this diversity will be thoroughly discussed and presented in the context of its contributions to host–pathogen interactions and the control/prevention of infection.

**Keywords:** lipopolysaccharide, serotyping, biosynthesis, motility, virulence, seroconversion, bacteriophage, nucleotide sugars

**INTRODUCTION AND HISTORICAL PERSPECTIVES**

To classify the differences among *Pseudomonas aeruginosa* strains isolated from environmental or clinical settings, serological methods have been employed whereby polyclonal antisera were raised against representative strains, then cross-adsorbed with other *P. aeruginosa* strains in order to yield a collection of strain-specific antisera. In order to exploit the specificity of the antibodies within these antisera, a number of immunochemical techniques were employed including slide agglutination tests and gel-diffusion precipitation reactions. The former constitutes a rather simple method of visualizing clumping of bacteria on a microscope slide by a small aliquot of specific antiserum in a matter of seconds, while the latter allows the diffusion of antibodies and antigen, usually bacterial cell lysates, from adjacent sample wells in agarose gels until equivalent amounts of these reagents meet to form visible precipitins (Mutharia and Lam, 2007). Interestingly, the precipitin reactions also provide the resolution to distinguish between cross-reacting groups, thereby giving rise to subgrouping among a specific serotype, for instance, serogroups 2a2b, 2a2c, etc. (Stanislavsky and Lam, 1997). Though tedious, these serotyping methods were the gold standards approximately 30 years ago, and are still being used by certain public health laboratories due to the low cost and rapidity in obtaining results. Since various laboratories from different parts of the world prepare their own antisera, many *P. aeruginosa* serotyping schemes have been established, such as the Homma, Habs, Lanyi and Bergan, and Fisher Immunotyping systems (reviewed in Stanislavsky and Lam, 1997). In order to standardize results stemming from the different serotyping systems, Liu et al. (1983) coordinated a special meeting between the *P. aeruginosa* serotyping experts, with the initiative giving rise to a standardized serotype classification termed the International Antigenic Typing Scheme (IATS), which has 17 serotypes based on Habs serotypes 1–12 plus 5 more serotypes from other serotyping systems (Stanislavsky and Lam, 1997). A subsequent study added 3 more serotypes to bring the total to 20 (Liu and Wang, 1990). Note that from this point onward in this review, the serotypes of *P. aeruginosa* strains mentioned are based on the IATS classification. Although it has been known that variations in the cell-surface lipopolysaccharide (LPS) of *P. aeruginosa* are responsible for serotyping, knowledge of the chemical structures of LPS and the underlying genetics of the biosynthetic process was lacking and prompted new research interests in this area.

The IATS serotyping scheme has been generally effective for classifying *P. aeruginosa* strains that are wildtype organisms producing smooth LPS possessing three distinct domains, namely, lipid A, core oligosaccharide (OS), and O polysaccharides or O antigens (O-Ag). This is not often the case in clinical settings as many of these isolates are found to be either partially lacking or completely devoid of O-Ag. Serotyping of chronic bacterial isolates from cystic fibrosis (CF) patients for epidemiological studies was particularly problematic because a very high proportion of these bacteria were found to be either polytypeable by more than one serotyping antisera or non-typable (NT). This has prompted the generation of new typing reagents based on monoclonal antibodies (mAbs) specific against each of the original 17 IATS serotypes (Lam et al., 1987a,b). In ensuing studies, comparison of the *P. aeruginosa* typing efficiencies with the complete set of anti-O1 to anti-O17 mAbs versus other methods, such as phage-susceptibility typing, pyocin typing, and restriction fragment length polymorphism (RFLP) DNA fingerprinting was performed (Ojenviyi et al., 1989, 1990; Speert et al., 1994); these methods differed substantially in their capacity to indentify unique typing patterns. Although RFLP was deemed to have the greatest discrimination power, LPS-based serotyping appears
QuiNAc is linked to the pseudaminic acid via the 3-hydroxybutanoyl group. In the polymer, QuiNAc is linked to the pseudaminic acid via the 3-hydroxybutanoyl group.

GENETICS OF O-SPECIFIC ANTIGEN BIOSYNTHESIS

Most P. aeruginosa strains produce two different forms of O-Ag, one of which contains a homopolymer of D-rhamnose, arranged in α1→3, α1→2, α1→3 trisaccharide repeats. This homopolymeric D-rhamnan is the common polysaccharide antigen (CPA; formerly termed A band) among P. aeruginosa strains. More details of the properties of CPA will be discussed in later sections. The other form of O-Ag produced by P. aeruginosa is a heteropolymer with three to five distinct sugars in its repeat units, and is named O-specific antigen (OSA; formerly termed B band). The differences of the composition of the O units form the basis of the IATS serotyping scheme. Knirel et al. (2006) have made a significant contribution to our understanding of all the serotypes by systematically elucidating the chemical structures of the OSA of all 20 IATS serotypes (Table 1). To appreciate the complexity and diversity of LPS produced by P. aeruginosa, it is noteworthy that heterogeneity exists in the chain length of either form of O-Ag present on the cell surface used to “cap” the core OS of P. aeruginosa (Figure 1). This accounts for the “ladder-banding” pattern when LPS from P. aeruginosa is analyzed by silver-stained SDS-PAGE gels and Western immunoblotting (Rocchetta et al., 1998a).

The importance of the OSA toward virulence and its use in serotyping has made it an attractive target for genetic studies to help understand both the plasticity of this region of the LPS and the complex steps of OSA biosynthesis. OSA biosynthesis follows the Wzy-dependent pathway model, originally proposed by Whitfield (1995); it involves the sequential activities of a series of integral inner membrane (IM) proteins, for which we have recently obtained comprehensive topological data helping to explain their respective functions (Islam et al., 2010). In this model, OSA sugar repeats are sequentially built on the lipid carrier undecaprenyl pyrophosphate (UndPP) on the cytoplasmic face of the IM. The UndPP-linked OSA repeats are then translocated to the periplasmic face of the IM by the flipase Wzx (Burrows and Lam, 1999), where they are polymerized by Wzy (De Kievet et al., 1995) through a putative catch-and-release mechanism (Islam et al., 2011), to modal lengths regulated by Wzz1 (Burrows et al., 1997) and

Table 1 | O-Antigen repeating units which compose the 20 IATS reference strains.

| IATS serotype | O-specific antigen repeat* |
|---------------|----------------------------|
| O1            | →4)-GalNAc-(1→4)-GlcNAc3NAcA-o-(β1→3)-o-FucNAc-(1→3)-o-QuiNAc-(1→ |
| O2            | →4)-o-ManNAc3NAmAnα-(β1→4)-GuNAc3NAcA-(1→3)-o-FucNAc-(β1→ |
| O5            | →4)-o-ManNAc3NAmAnα-(β1→4)-o-ManNAc3NAcA-(β1→3)-o-FucNAc-(1→ |
| O16           | →4)-o-ManNAc3NAmAnα-(β1→4)-o-ManNAc3NAcA-(β1→3)-o-FucNAc-(β1→ |
| O18           | →4)-o-GuNAc3NAmAnα-(1→4)-o-ManNAc3NAcA-(β1→3)-o-FucNAc-(1→ |
| O20           | →4)-o-GuNAc3NAmAnα-(1→4)-o-ManNAc3NAcA-(β1→3)-o-FucNAc-(1→ |
| O3            | →2)-o-Rha3OAc-(1→6)-o-GlNAc-(1→4)-o-GalNAc4OAcA-(1→3)-o-QuiNAcN4SHb-(β1→ |
| O4            | →2)-o-Rha-(1→3)-o-CucNAc-(1→3)-o-FucNAc-(1→3)-o-QuiNAc-(1→ |
| O6            | →3)-o-Rha-(1→4)-o-GalNAc3OAcAn-(1→4)-o-GalNAcOAc-(1→3)-o-QuiNAc-(1→ |
| O7            | →4)-Pse4OAc5NHRhb7NFox(2→4)-o-Xy1-(β1→3)-o-FucNAc-(1→ |
| O8            | →4)-Pse4OAc5NHRhb7NFox(2→4)-o-β1-(β1→3)-o-FucNAc-(1→ |
| O9**          | →371-Pse4OAc5NHRhb7NBox-(β2→4)-o-FucNAc-(1→3)-o-QuiNAc-(β1→ |
| O10           | →3)-o-Rha2OAc-(1→4)-o-GalNAcA-(1→3)-o-QuiNAc-(1→ |
| O19           | →3)-o-Rha-(1→4)-o-GalNAcA-(1→3)-o-QuiNAc-(1→ |
| O11           | →2)-o-Glc-(β1→3)-o-FucNAc-(1→3)-o-FucNAc-(β1→ |
| O12           | →8)-o-β1-(β1→3)-Lc-CucNAc-(1→3)-o-QuiNAc-(1→ |
| O13           | →1)-o-Rha-(1→3)-o-Rha-(1→4)-o-GalNAc3OAcA-(1→3)-o-QuiNAc-(β1→ |
| O14           | →1)-o-GlNAc-(1→3)-o-Rha-(1→4)-o-GalNAc3OAcA-(1→3)-o-QuiNAc-(β1→ |
| O15           | →2)-o-Rib-(β1→3)-o-GalNAcA-(1→ |
| O17           | →4)-o-Rha-(1→3)-o-ManNAcA-(1→ |

Adapted from Knirel et al. (2008), O-Ag repeats are clustered according to structural similarity. Conserved sugars, colored by backbone structure, are adorned with various side groups depicting the diverse phenotypes generated by the OSA gene cluster throughout all 20 serotypes. Anomeric conformations are α unless marked. Sugars have the pyranose form except ribose in O15. Ac, acetyl; 8β-Leg, 5,7-diamino-3,5,7-tetradec-xylyl-glycero-c-galacto-non-2-ulosonic (8-epi-glycero-namnic) acid; Fo, fomyl; FucN, 2-amino-2,6-dideoxy-galactose; GlcN, 2-amino-2-deoxy-galactose; GlcNA, 2-amino-2-deoxy-glucuronic acid; GuNA, 2-amino-2-deoxy-galuronic acid; ManNA, 2-amino-2-deoxy-mannose; ManN3NAc, 2-amino-2-deoxy-mannuronic acid; N, amino; NAc, acetamido; HAc, acetamido; OAc, O-acetyl; Pse, 5,7-diamino-3,5,7-tetradec-xylyl-glycero-c-galacto-non-2-ulosonic (pseudaminic) acid; QuiN, 2-amino-2,6-dideoxyglucose; Rha, rhamnose; RHB (R)3-hydroxybutanoy; Rib, ribose; SHb, (3S)-3-hydroxybutanoyl. In the polymer, QuiNAc is linked to the pseudaminic acid via the 3-hydroxybutanoyl group. **QuiNAc is linked to the pseudaminic acid via the 3-hydroxybutanoyl group.
Diversity of Surface LPS Glycoforms in a Single Cell

Monosaccharides
- D-ManNAc3ManNAc
- D-ManNAc3NacAc
- D-FucNAc
- D-Rha
- L-Rha
- D-Glc
- D-GalN-L-Ala
- L-D-Hep
- Kdo
- GlcNAc

Wild Type
- ∆wcm/wzt
- ∆wxx
- ∆wzy
- ∆wzz1
- ∆wzz2
- ∆waa1
- ∆waa2

Capped LPS (A band)
- CPA-Capped LPS (B band)
- OSA-Capped LPS (B band) [Core + 1]
- OSA-Capped LPS (B band) [Low MW]
- OSA-Capped LPS (B band) [Long]
- OSA-Capped LPS (B band) [Very Long]

*Terminal GlcNAc is not present in all serotypes of P. aeruginosa.
1Cova lent linkage of the CDA polymer to core OS has yet to be conclusively determined. CPA polymer length averages ~70 D-Rha monosaccharides.
2 modal chain length of ~12–16 and ~22–30 repeat units is conferred by Wzz1.
3 modal chain length of ~40–50 repeat units is conferred by Wzz2.
4 terminal GlcNAc is not present in all serotypes of P. aeruginosa.

FIGURE 1 | Representations of the heterogeneity/diversity of the LPS glycoforms present on the surface of a single P. aeruginosa cell. Genetic defects in members of various assembly pathway genes and their resultant changes to the variety of LPS glycoforms present have been indicated, with “✓” or “χ” representing their presence or absence, respectively. Substitutions of various lipid A and core OS sugars with phosphate groups and L-Ala have been indicated with yellow circles and red diamonds, respectively. The OSA polymer from serotype O5 has been displayed as a representative polymer.

Wzz2 (Daniels et al., 2002). The modal length of OSA imparted by Wzz2 (40–50 repeat length) is longer than that imparted by Wzz1 (12–16 repeats and 22–30 repeats; Daniels et al., 2002). However, Wzz1 is apparently more important for virulence than Wzz2 (Kintz et al., 2008). Finally, the complete OSA chain is ligated to lipid A-core by the O-Ag ligase WaaL (Abeyrathne et al., 2005; Abeyrathne and Lam, 2007). Preliminary investigations mapped the OSA cluster to 37 min of the P. aeruginosa PAO1 (serotype O5) genome (Lightfoot et al., 2008).
and Lam, 1993), corresponding to pa3141 to pa3160 in the annotated genome of strain PAO1 (Stover et al., 2000). This first reported LPS OSA cluster was isolated from a cosmids-based genomic library. Clone pFV100 from the library was able to complement mutant get, a Tn5–761 insertional mutant of PAO1, defective in B band (OSA) biosynthesis (Lightfoot and Lam, 1993). Subsequently, Burrows et al. (1996) obtained the sequence of the entire OSA cluster. To characterize the function of the genes encoded in this cluster, knockout mutant constructs were generated for each of the genes and the mutants were examined for their effect on LPS production in P. aeruginosa. The list of OSA biosynthesis genes in this serotype O5 cluster and their functions were determined based on genetic studies as well as biochemical and chemical evidence (Figure 2A; Table 2, Burrows et al., 1996). Following the success in characterizing the O5 OSA biosynthesis locus, the sequences of the O6 (Belanger et al., 1996) obtained the sequence of the entire OSA cluster. To characterize flanked by himD/ihfB (O11 (Dean et al., 1995), the information in this table has been updated with references which indicate that either genetic or biochemical evidence have been collected to demonstrate the function of each of these genes within the OSA biosynthesis cluster of PAO1 serotype O5. Note that in contrast to the genome of P. aeruginosa, which has a relatively high mol% G+C content (67%), the genes within this cluster showed significantly lower mol% G+C, ranging from 45.2 to 61.9%, evidence of horizontal gene transfer.

### Table 2 | P. aeruginosa PAO1 serotype O5 OSA biosynthesis cluster*

| Gene       | Proposed/demonstrated function                                      | Mol% G + C | References                      |
|------------|---------------------------------------------------------------------|------------|---------------------------------|
| wzb/pa3160 | OSA chain length regulator                                         | 49.5       | Burrows et al. (1997), Daniels et al. (2002) |
| wbpA/pa3159 | UDP-N-acetyl-α-glucosamine 6-dehydrogenase                          | 54.5       | Miller et al. (2004)            |
| wbpB/pa3158 | UDP-2-acetamido-2-deoxy-α-glucuronic acid 3-dehydrogenase           | 52.8       | Westman et al. (2009)           |
| wbpC/pa3157 | Possible O-acetyltransferase                                        | 53.1       |                                  |
| wbpD/pa3156 | UDP-2-acetamido-3-amino-2,3-dideoxy-α-glucuronic acid N-acetyltransferase | 53.9       | Wenzel et al. (2005), Westman et al. (2009) |
| wbpE/pa3155 | UDP-2-acetamido-2-dideoxy-α-ribo-hex-3-uluronic acid transaminase   | 52.8       | Westman et al. (2009)           |
| wzy/pa3154 | OSA α-polymerase                                                    | 44.6       | De Kieft et al. (1995), Islam et al. (2010), Islam et al. (2011) |
| wzz/pa3153 | OSA unit flippase                                                   | 49.3       | Burrows and Lam (1999), Islam et al. (2010) |
| hisH2/pa3152 | Imidazole glycerol phosphate synthase subunit                       | 49.3       | King et al. (2009)              |
| hisF2/pa3151 | Imidazole glycerol phosphate synthase subunit                       | 50.0       | King et al. (2009)              |
| wbpG/pa3150 | Amidotransferase                                                    | 44.5       |                                  |
| wbpH/pa3149 | Glycosyltransferase                                                 | 45.6       |                                  |
| wbpI/pa3148 | UDP-N-acetylglucosamine 2-epimerase                                 | 50.2       | Westman et al. (2009)           |
| wbpJ/pa3147 | Glycosyltransferase (GT-4)                                          | 54.5       |                                  |
| wbpK/pa3146 | NAD-dependent epimerase/dehydratase                                 | 56.8       |                                  |
| wbpL/pa3145 | Glycosyltransferase                                                 | 55.5       | Rocchetta et al. (1998a)         |
| wbpM/pa3141 | Nucleotide sugar epimerase/dehydratase                              | 61.9       | Creuzenet and Lam (2001)         |

*Modified from Burrows et al. (1996), the information in this table has been updated with references which indicate that either genetic or biochemical evidence have been collected to demonstrate the function of each of these genes within the OSA biosynthesis cluster of PAO1 serotype O5. Note that in contrast to the genome of P. aeruginosa, which has a relatively high mol% G+C content (67%), the genes within this cluster showed significantly lower mol% G+C, ranging from 45.2 to 61.9%, evidence of horizontal gene transfer.

**EVIDENCE OF HORIZONTAL GENE TRANSFER REVEALS THE PRESENCE OF CONSERVED AND DUPLICATED CLUSTERS IN CERTAIN SEROTYPES**

The high conservation of the wbpM gene and its localization at the 3′ end of the O5 cluster upstream of wbpM (Burrows et al., 1996). This variation helps to explain the diversity of the LPS in P. aeruginosa as a property of the genetic differences among the IATS serotypes. Other factors that influence the OSA diversity are outlined in the following sections.
FIGURE 2 | Organization of the genes within OSA biosynthesis clusters.

(A) OSA biosynthesis cluster of serotype O6 adapted from Burrows et al. (1996). The gene cluster is located on the complementary strand; genes which match the PFAM designation are colored accordingly. Genes not involved in OSA biosynthesis are depicted in gray including a large insertion sequence (IS). (B) Adapted from Raymond et al. (2002), the OSA biosynthesis gene clusters were organized into 11 groups based on sequence conservation. Genes were designated using the PFAM database; specific protein families which occur a minimum of three times throughout all 20 OSA biosynthesis clusters are represented by a specific color. A red outline depicts an ORF with potential transmembrane-spanning domains. Previously identified genes are labeled above the respected cluster if present within the serotype. Insertion sequences (IS) present within genes are depicted by a secondary gray box.
is horizontal gene transfer (HGT), i.e., the direct transmission of genetic information across the species barrier, which often results in similar/identical gene products. Interestingly, Dean and Goldberg (2000) probed all 20 serotypes for the presence of genes that are conserved in the O11 OSA biosynthesis cluster, and discovered that the IATS O17 strain possesses a cryptic copy of the O11 locus within its genome. The O11 cluster was apparently rendered non-functional due to the presence of IS:SPA11B disrupting the 5′ end of the *wbpD* gene. However, when a complete O11 OSA cluster was supplied in *P. aeruginosa*, LPS produced by the complemented strain reacted with anti-O11 antiserum in Western immunoblotting. This suggested that the cryptic O11 serotype genes found within the O17 strain were at one time functional. The presence of a secondary locus within one genome lends support to HGT. Apparently, the OSA LPS genes for the O17 serotype that were acquired or have evolved may have caused a gradual loss of function or suppression to the O11 OSA biosynthesis gene cluster through the incorporation of an IS element. Interestingly, the O17 OSA serotype is very similar to the *Burkholderia cepacia* serogroup O5 antigen, confirming that this may be an ancestral locus which was acquired by HGT (Dean and Goldberg, 2000). Similarly, Raymond et al. (2002) discovered the presence of an additional cryptic locus corresponding to the O3 cluster within the genome of serotype O15, with further investigation revealing that the O3 cluster in the O15 strain was rendered non-functional due to IS elements at two different sites within the cluster.

Evidence that HGT contributes to the diversity of *P. aeruginosa* LPS is not limited to transfer of the entire OSA biosynthesis locus. Varying degrees of genetic information exchange between bacteria are apparent since bacterial pathogens of different genera share conserved metabolic pathways for the biosynthesis of nucleotide sugars. A number of rare sugars are found in the OSA of *P. aeruginosa*, e.g., L-FucNAc (a constituent of the O11 OSA), di-N-acetylated mannuronic acid (d-ManNAc3NAcA, a constituent of the O5 OSA), and a uronamide sugar GalNAcAN (a constituent of O6 OSA). The metabolic pathways for the biosynthesis of the nucleotide-activated precursors for these rare sugars are only found in a few bacterial pathogens but not in humans (Kneidinger et al., 2003; Westman et al., 2008, 2009; King et al., 2010). The biosynthesis of UDP-L-FucNAc from the substrate UDP-α-GlcNAc requires the products of three genes from the serotype O11 OSA cluster, namely, *wbpB*, *wbpC*, and *wbpD*. Two sets of functional homologs of these three genes are found in *Staphylococcus aureus*, namely *cap5E-cap5F-cap5G* and *cap8E-cap8F-cap8G* for the biosynthesis of serogroup-5 and serogroup-8 capsular polysaccharides, respectively (Kneidinger et al., 2003). The complexity of the biosynthesis of the UDP-activated precursor of d-ManNAc3NAcA requires five steps. Indeed, the activity of the encoded products of *wbpA*, *wbpB*, *wbpE*, *wbpD*, and *wbpI*, acting in this particular order, and also starting with UDP-GlcNAc as the starting substrate, have been fully characterized (Westman et al., 2008, 2009). The same pathway is shared by *Bordetella pertussis* as it also possesses d-ManNAc3NAcA as a sugar in its LPS. However, only four of the five genes, *wlbA*, *wlbB*, *wlbC*, *wlbD*, which are functional homologs of *wbpB*, *wbpE*, *wbpD*, and *wbpI*, respectively, are localized within the O-Ag biosynthesis gene clusters. Two copies of the homolog of *wbpA* were found elsewhere in the *B. pertussis* genome (Westman et al., 2008). In a separate investigation, our group has determined that *wbpO*, *wbpP*, and *wbpS* encode the enzymes for the three metabolic steps that convert UDP-GlcNAc by oxidation (WbpO), epimerization (WbpP), and amidotransfer (WbpS) to form the uronamide sugar precursor UDP-GalNAcAN for O6 OSA. Interestingly, functional homologs of these three genes were found in *Escherichia coli* O121. As well, homologs of these three genes were also found in *Francisella tularensis* (King et al., 2010); however, they have not been tested to determine whether they could complement knockout mutants of the *P. aeruginosa* genes to restore production of O6 LPS that contains GalNAcAN in its O-Ag unit. Therefore, *P. aeruginosa* and other bacterial pathogens likely acquired genes for the biosynthesis of the abovementioned UDP-sugars from common ancestors. However, investigation of the evolutionary derivations and lineages of these pathways is beyond the scope of this review.

**EVIDENCE OF DIVERSITY IN LPS EXPRESSION IN *P. AERUGINOSA* DUE TO CHROMOSOMAL INSERTIONS**

Large chromosomal inversions (LCIs), which can be induced by ISs, persist throughout Gram-negative bacteria and have altered the balance between chromosomal stability and variability. Clonal isolates taken from CF patients were found to contain additional, non-conserved LCIs (Schmidt et al., 1996). *P. aeruginosa* “clone C,” found in various geographic locations and in non-CF patients (Kidd et al., 2011), possesses a particular additional sequence, IS6100, which was first discovered in *Mycobacterium tuberculosis* and known to cause genetic rearrangements in heterologous hosts. The LCIs of the “clone C” strains have been associated with *wbPM* (pa3141) disruption causing loss of the OSA (Kresse et al., 2003). The genomes of two CF isolates were compared to that of the wildtype PAO1, with regions of high variability localized to the OSA biosynthesis locus. Targeted sequencing of the OSA cluster provided a high-resolution view of the genetic changes that arise during clonal infection of CF patients. Genomes of all late-stage clinical isolates contained ISs, as judged by gel electrophoresis (Spencer et al., 2003). The OSA region of a clonal isolate (strain 1–60) was nearly identical to serotype O1 except for the presence of a large 1.5-kb IS, which disrupted a reading frame responsible for the biosynthesis of an unspecified nucleotide sugar. Additionally, a second strain (2–164) was found to contain a 2-bp deletion in *wbPQ*, which has not yet been characterized. Both of these observed mutations caused defects in LPS production and conferred a NT phenotype. Interestingly, some of the earlier NT strains did not contain the IS elements (Spencer et al., 2003). Strain-specific islands were discovered with high similarity to previously identified sequences (e.g., PAGI-1 in O-Ag biosynthesis) as well as some phage-related sequences; however, these contained a low mol% G + C content possibly indicating acquisition from other bacterial species (Spencer et al., 2003), since the *P. aeruginosa* genome has a high mol% G + C content. Whole-genome sequencing has begun to provide snapshots of these genetic changes, which occur during clonal infection; future work using this approach may provide a roadmap for navigating the infection stages of *P. aeruginosa*.

**LYSOCENIC CONVERSION BY BACTERIOPHAGE ALTERS THE OSA SEROTYPE TO PROMOTE RESISTANCE**

A natural part of bacteriophage biology is engagement of a lyso-"recall" cycle wherein a temperate phage invades a bacterial cell and incorporates its genetic material into the host genome, leading to the...
propagation of phage DNA during host cell replication. This part of the phage life cycle could also lead to the incorporation of exogenous bacterial genes into the host genome (Miller et al., 1974). The D3 phage is a temperate phage that readily infects P. aeruginosa (Cavenagh and Miller, 1986). LPS-specific phage-dependent conversion of the OSA occurs throughout Gram-negative bacteria (Robbins et al., 1965; Bagdian et al., 1966). Holloway and Cooper (1962) observed that after strain PAO1 (serotype O5) was infected with D3 phage, at least 20% of the new colonies from subcultures tested could not be agglutinated by O5-specific typing antiserum. The mechanism of the changes occurring at the bacterial cell surface was unknown at the time. In a later study, a similar observation was made by the Hancock group who showed that lysogenized PAO1 cells had an altered LPS phenotype (Hancock et al., 1983), and this change was later determined to be due to the addition of an acetyl group at position 4 of the fucosamine residue and an alteration in the linkage of the trisaccharide repeating unit bond from \( \alpha \) to \( \beta \) to \( \beta \) (Kuzio and Kropinski, 1983).

In a more recent study by our group, a region of the D3 genome was found to contain a gene that hybridized in Southern blots to an O-Acetylation gene found in the PAO1 genome (Newton et al., 2001). This region spans 3.6 kb of the D3 genome, and contains three genes: (i) oac (encodes an O-acetylase), which adds an O-acetyl group to the OSA FuCNAc residue; (ii) \( \text{wzy}_3 \) (encodes a \( \beta \)-polymerase) which polymerizes the OSA repeats to form \( \beta \) to \( \beta \) to \( \alpha \) linkages; and (iii) iap (encodes an \( \alpha \)-polymerase inhibitor, a small 31 amino acid peptide with a single transmembrane domain). Iap possesses the activity to render Wzy non-functional thereby preventing the formation of the usual \( \alpha \) to \( \beta \) to \( \beta \) linkage among O-Ag subunits of strain PAO1 (Newton et al., 2001). Expression of this operon in trans produced an LPS banding pattern similar to that of LPS from O16, and these bands reacted strongly with the anti-O16 mAb MF47-4 in Western immunoblotting but not with the anti-O5 mAb MF15-4 (Newton et al., 2001). O-Ag subunits of serotypes O2 and O16 are linked by \( \beta \) to \( \beta \) to \( \beta \) bonds, and both serotype strains contain a functional \( \text{wzy}_3 \), which is actively being repressed by a previously undetected \( \text{iap} \). Interestingly, the \( \text{wzy}_3 \) gene in O2 and O16 was not localized to the OSA biosynthesis cluster, further supporting the idea that this could be a D3 phage-associated gene. The authors initially suggested that \( \text{wzy}_3 \) and \( \text{iap} \) identified in P. aeruginosa O2 and O16 strains are xenologs of the D3 phage. However, the mol% G + C of \( \text{wzy}_3 \) and \( \text{iap} \) respectively are inconsistent when compared to the mol% G + C values of both the PAO1 and the D3 phage genomes; therefore, the authors attributed the origin of these genes to an exogenous source (Kaluzny et al., 2007).

It has become dogma that once P. aeruginosa has colonized the lungs of CF patients, it adapts to a biofilm lifestyle to protect the bacterial community from hostile host defense and promote survival. The emergence of mucoid colony morphology is a hallmark of such adaptation and provides selective pressures that favor the occurrence of mutator phenotypes. These mutators in the bacterial community accumulate genetic changes resulting in altered phenotypes and gene expression profiles. The cause of the mutator phenotype is the inactivation/suppression of the methyl-directed mismatch repair (MMR) mechanism used to repair DNA replication errors and decrease the likelihood of homologous recombination events (Kunkel and Erie, 2005). Mutations in this region are known to cause loss of DNA proof reading thereby increasing the likelihood of mutant phenotypes which may convey survival advantages (Lecerc et al., 1996). This “hypermutable” phenotype was observed in 20% of P. aeruginosa isolates collected from CF airways and absent in isolates collected from blood and from the airway of non-CF patients (Oliver et al., 2000). One of the major genes responsible for the MMR mechanism is mutS; inactivation of this gene effectively shuts off the MMR cascade. Genetic analysis of the mutS gene from the previously characterized hypermutable P. aeruginosa strain JMSMA7 revealed a 3.3-kb IS and an 8-nucleotide repeat which flanked a 54-nucleotide deletion (Oliver et al., 2000). Interestingly, contained within the 3.3-kb IS lies another 1.2-kb IS element that is 90% homologous to IS222 from the P. aeruginosa D3 phage (Kropinski et al., 1994; Oliver et al., 2000). This region was interpreted as a veritable “hot-spot” of recombination within the mutS gene promoting its loss of function and increasing the number of mutations (Oliver et al., 2002). An in-depth longitudinal study was undertaken by Smith et al. (2006) to examine the presence of the mutator phenotype in CF patients. The information gained from this study was used to further characterize the hypermutable isolates and determine the locations of these genetic errors. Sequencing of genes in two CF isolates taken at 6 months and a comparable isolate taken at 96 months during time course experiments revealed the accumulation of 68 mutations, wherein 13 were associated with virulence factors including genes associated with O-Ag biosynthesis (Smith et al., 2006). A continuing in-depth study using the genomes of these two isolates revealed an apparent increase in mutations within genes responsible for antibiotic resistance, as well as in lasR, a gene responsible for the regulation of quorum sensing in P. aeruginosa (Mena et al., 2008). The inactivation of lasR and subsequently rhlI in the hypermutable phenotype may result in a decrease of migA expression as indicated earlier by Yang et al. (2000). MigA is a glycosyltransferase responsible for the uncapped core OS phenotype, under the control of the RhlR–RhlI system (described below); when inactivated, the levels of MigA decrease causing an accumulation of “core-plus-one,” a LPS glycoform with a single OSA repeat (Yang et al., 2000; Figure 1), a less virulent form of LPS. This degree of genotype plasticity is proposed to aid in the survival of bacteria within stressful environments during the transition to a chronic infection (Smith et al., 2006).

QUORUM-SENSING ALTERS CORE OS STRUCTURE THAT AFFECTS O-AG LIGATION

Cell density-dependent gene regulation plays an integral role in cell-surface virulence factor expression, altering gene expression based on population size (Whiteley et al., 1999). As previously mentioned, most P. aeruginosa CF isolates from chronically infected patients are devoid of OSA and begin to favor the typical “rough” phenotype, i.e., producing LPS that lacks the full-length O antigen side chain, being non-motile and non-piliated (Hancock et al., 1983). Although previous studies have shown the genetic relationship of the alginate-based mucoid phenotype (Ma et al., 1998), the regulation of changes governing the transition from smooth to rough LPS is not well defined. The “mucus inducible gene” migA (described above) is a putative glycosyltransferase and was discovered when P. aeruginosa was grown in the presence of CF mucus (Wang et al., 1996). Expression of the migA gene was found to be a result of
quorum sensing (Whiteley et al., 1999), dependent on the release of autoinducer molecules by P. aeruginosa into the local environment. When the bacterial density increases to an appropriate level, the local concentration of autoinducer molecules reaches a critical threshold past which they can consistently bind to transcription regulators across the entire cell population, resulting in alternative gene expression across the entire community (Fuqua et al., 1994). The migA gene was found to be under the control of RhlI–RhlR regulatory system, one of the two principal quorum-sensing systems, due to its severe decrease in expression in a rhlI–rhlR double knockout as compared to wildtype. Furthermore, it possesses an upstream las-box-like sequence (CT-N_{11})-AG), known to be a quorum-sensing recognition site for quorum-sensing regulators (Yang et al., 2000). Interestingly, overexpression of migA resulted in a loss of “core-plus-one” but not the higher molecular weight LPS, leading the authors to speculate the existence of a secondary glycosyltransferase (Yang et al., 2000). Recently, that second glycosyltransferase was identified by our group as wapR, located 158-bp downstream of the core biosynthesis locus (Poon et al., 2008). Genetic and biophysical analysis of these two genes and their products revealed that both are responsible for core modifications in LPS biosynthesis (Figure 3). The expression of these genes results in an altered core structure: migA is responsible for the addition of the l-rhamnose^4 (l-Rha^4) in the α1 → 6 linkage, preventing the addition of O-Ag to lipid A-core by the O-Ag ligase WaaL (Yang et al., 2000; Poon et al., 2008). The wapR gene is responsible for the addition of the core sugar l-Rha^4, creating the favored linkage site for the addition of the O-Ag by WaaL (Poon et al., 2008). A genomic knockout of wapR resulted in the loss of higher molecular weight LPS bands, corresponding with the addition of O-Ag to lipid A-core. Work is currently underway in our laboratory to establish the relationship between migA and wapR gene expression. The cell density-dependent effects on gene regulation are of particular importance as CF isolates are often found in mature biofilms, a direct byproduct of quorum sensing.

THE AGE OF TECHNOLOGY BROADENS GENE EXPRESSION INVESTIGATIONS

In silico-based algorithms and bioinformatic methods have moved to the forefront of genomic research, performing a variety of predictions to provide a global view of genome expression profiles. When combined with microarray data, in silico modeling can be applied to examine an entire genome, particularly for investigating up and down regulations of genes throughout a longitudinal experiment. Using this approach, Oberhardt et al. (2010) examined the gene expression in a number of CF isolates throughout a progressive CF infection by a clonal strain of P. aeruginosa. They observed that the genes relating to LPS and capsule biosynthesis were downregulated during chronic CF infection. The development of new computer-based programs and ever-improving novel DNA sequencing techniques will help to broaden our understanding of the polymorphic nature of OSA of P. aeruginosa during CF infection.

STRUCTURE, GENETICS, AND FUNCTION OF CPA (A BAND) DISCOVERY OF CPA

In the 1980s, CPA was detected to be a P. aeruginosa antigen that reacted with a human monoclonal antibody produced by Sawada’s group (Sawada et al., 1985). Further analysis found the antigen to be a neutral polysaccharide composed of α-rhamnose (α-Rha) that is separable as well as structurally and immunologically distinct from the predominant OSA (Yokota et al., 1987; Kocharova et al., 1988; Rivera et al., 1988). In a study by McGroarty’s group, the authors used gel filtration chromatography and a buffer containing deoxycholate, which helped to disperse LPS from micelles and aggregates, to obtain two separate fractions of surface polysaccharide from P. aeruginosa (Rivera et al., 1988). The fast-migrating fraction contained OSA (named B band) that was reactive to serotype-specific antibodies, while the slower-migrating fraction (CPA, i.e., A band) did not react with any of the mAbs that the Lam laboratory had raised against the IATS serotypes. Upon receiving the CPA from McGroarty, our group raised a number of hybridoma cell lines (e.g., N1C9, N1F10) that secreted mAbs that are specific against...
CPA. Using mAb N1F10 in dot-blot and Western-immunoblotting analyses, the presence of CPA was detected in 14 of the 20 P. aeruginosa IATS serotypes. The strains that did not react to the mAb belong to serotypes O7, O12, O13, O14, O15, and O16 (Lam et al., 1989; Currie et al., 1995). However, a majority of clinical P. aeruginosa isolates examined since then were found to be devoid of OSA but produce CPA as the predominant surface polysaccharide antigen (Lam et al., 1989; Weisner et al., 2007). In a separate study, CPA was found to be a receptor for bacteriophage A7, indicating its significance as a cell-surface polysaccharide component of P. aeruginosa and its role in the co-evolution with temperate bacteriophages (Rivera et al., 1992).

**STRUCTURAL STUDY OF CPA**

Structural elucidation of CPA from different P. aeruginosa strains has been reported. Yokota's group was the first to discover that the predominant structure of the common polysaccharide from P. aeruginosa strain IID 1008 (ATCC 27584; Yokota et al., 1987) reactive to a human mAb is composed of a trisaccharide repeating unit: \( \alpha_1 \rightarrow \delta\text{-Rha}(\alpha_1 \rightarrow 3)\delta\text{-Rha}(\alpha_1 \rightarrow 2)\delta\text{-Rha}(\alpha_1 \rightarrow \cdot) \). Our group prepared CPA from PAO1 (serotype O5;Arsenault et al., 1991) and by using high-field nuclear magnetic resonance (NMR) and mass spectrometry (MS) methods, we collected data to show that this neutral polysaccharide contains a \( \delta\text{-rhamnan} \) with the same structural repeat as demonstrated by Yokota et al. (1987). Two other groups who examined CPA from a serotype O7 strain (Lanyi–Bergan classification; Kocharova et al., 1988) and strain PA103 (serotype O11; Choudhury et al., 2005) also observed the same trisaccharide repeat structure. However, aside from the predominant \( \delta\text{-Rha} \) trisaccharide repeat units, small amounts of other components have also been reported with less consistency among the different studies. For example, the presence of 3-\( \text{O} \)-methylrhamnose, ribose, mannose, glucose, and a 3-\( \text{O} \)-methylhexose have been detected from CPA prepared from strain PAO1 (Arsenault et al., 1991); 3-\( \text{O} \)-methyl-6-deoxyhexose, glucose, xylose, alanine, galactosamine, and phosphorus have all been reported as components of CPA fractions for strain IID 1008 (ATCC 27584; Yokota et al., 1987). Glucose, mannose, and phosphate have been found in the study of CPA from the O7 strain (Kocharova et al., 1988), while mannose and GlcNAc, as well as small amounts of \( \text{O} \)- and \( \text{N} \)-acyetyl substitutions, were found in the CPA from strain PA103 (Choudhury et al., 2005).

These differences reported in the minor components of CPA could be due to several factors including genuine strain differences, the details in the preparation of CPA, and the sensitivity and accuracy of the detection methods used. In conclusion, the precise structure of CPA has not been defined to date. Further study is currently underway in our lab to determine the identity of the initiating sugar, the terminal sugar, the position and amount of other sugar components, as well as the degree and position of the methylation and acetylation modifications of CPA from strain PAO1.

**GENETICS OF CPA BIOSYNTHESIS**

A cosmid clone pFV3 that is able to restore CPA biosynthesis in a CPA-deficient mutant (strain AK1401) was first obtained from a cosmid library of P. aeruginosa prepared from strain PAO1. Mutant strain AK1401 was a phage-resistant mutant derived from the PAO1 background (Lightfoot and Lam, 1991). When expressed in trans, a recombinant strain of P. aeruginosa expressing CPA also contained the pFV3 plasmid. In previous genetic studies in this laboratory, a cosmid library of P. aeruginosa strain PA103 was analyzed to find cosmid clones that would restore CPA biosynthesis in CPA-deficient mutant (strain AK1401) to a wild-type phenotype. Out of 20 of the 20 IATS reference strains examined, 14 of the 20 isolates (O7, O13, O14, O15, and O16) contained a three-gene cluster, \( \text{wbpY} \), \( \text{wbpZ} \), \( \text{wzm} \), encoding putative rhamnosyltransferases responsible for the specific linkages of the \( \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 2 \) bonds that form the \( \delta\text{-Rha} \) trisaccharide repeating unit (Rocchetta et al., 1998a). The last two genes, \( \text{wzm} \) and \( \text{wzt} \) (pa5449, pa5448, and pa5447), encode putative rhamnosyltransferases responsible for the specific linkages of the \( \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 2 \) bonds that form the \( \delta\text{-Rha} \) trisaccharide repeating unit (Rocchetta et al., 1998a). The last two genes, \( \text{wzm} \) and \( \text{wzt} \) (pa5449, pa5448, and pa5447), encode putative rhamnosyltransferases responsible for the specific linkages of the \( \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 2 \) bonds that form the \( \delta\text{-Rha} \) trisaccharide repeating unit (Rocchetta et al., 1998a). The last two genes, \( \text{wzm} \) and \( \text{wzt} \) (pa5449, pa5448, and pa5447), encode putative rhamnosyltransferases responsible for the specific linkages of the \( \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 3, \alpha_1 \rightarrow 2 \) bonds that form the \( \delta\text{-Rha} \) trisaccharide repeating unit (Rocchetta et al., 1998a).
and might have disrupted the promoter region of this gene cluster, which would explain the lack of CPA expression in this strain. We also found that this genomic region from strain PA7 showed a much lower sequence identity to that of the other three strains. The strains PA01, PA14, and LESB58 share more than 95% sequence identity in most of the proteins encoded by genes of the CPA biosynthesis cluster, while PA7 showed less than 50% identity to these three strains. This may explain why the cosmids pFV3 did not complement CPA biosynthesis in this serotype while still being able to do so for the other five serotypes that lack CPA LPS. Interestingly, we found CPA biosynthesis in this serotype while still being able to do so for strains. This may explain why the cosmid pFV3 did not complement the sis cluster, while PA7 showed less than 50% identity to these three strains. The strains also found that this genomic region from strain PA7 showed a much lower sequence identity to that of the other three strains. The strains might have disrupted the promoter region of this gene cluster, which would explain the lack of CPA expression in this strain. We also found that this genomic region from strain PA7 showed a much lower sequence identity to that of the other three strains. The strains PA01, PA14, and LESB58 share more than 95% sequence identity in most of the proteins encoded by genes of the CPA biosynthesis cluster, while PA7 showed less than 50% identity to these three strains. This may explain why the cosmids pFV3 did not complement CPA biosynthesis in this serotype while still being able to do so for the other five serotypes that lack CPA LPS.

**STRUCTURE OF INNER CORE OS**

Composition of the sugar constituents in inner core OS is identical among *P. aeruginosa* strains and it consists of two residues of 3-deoxy-d-manno-ocululosonic acid (KdoI and KdoII) and two residues of L-glycero-d-manno-heptose (HepI and HepII, Figure 4). The inner core is highly conserved when compared to other Gram-negative bacteria. A distinguishing feature of the *P. aeruginosa* inner core is a high degree of phosphorylation that is essential for viability of *P. aeruginosa*, since mutation of either of the two genes (wapP and wadP) encoding kinases that are responsible for adding phosphate groups to the inner core heptoses is lethal (Walsh et al., 2000). In most *P. aeruginosa* strains, three phosphorylation sites have been identified; positions 2 and 4 on HepI and position 6 on HepII. In a CF isolate, HepII has been shown to possess an additional phosphorylation site on position 4 (Knirel et al., 2006). In theory, monophosphate, diphosphate, or triphosphate groups might occupy each of the phosphorylation sites (Kooistra et al., 2003; Choudhury et al., 2005; Bystrova et al., 2006). Some of the analyzed *P. aeruginosa* strains have shown the non-stoichiometric substitution of a phosphate group on position 2 of HepI by ethanolamine-phosphate or ethanolamine-diphosphate (Bystrova et al., 2006). However, Knirel et al. (2006) suggested that the actual content of phosphates and ethanolamine-phosphates in the inner core might be higher, since these phosphate groups can be released from the core during preparation of LPS prior to structural analyses. In addition to phosphorylation, HepII is modified with another non-carbohydrate substituent, namely an O-carbamoyl group on position 7 (Beckmann et al., 1995).

**STRUCTURE OF OUTER CORE OS**

The outer core of *P. aeruginosa* LPS is composed of one d-galactosamine (GalNAc), one L-Rha, and three or four d-glucose (GlcI–GlcIV) residues. GaNAc is further substituted on position 2 with an alanoyl (Ala) group or in some truncated core structures with an acetyl group (Sanchez Carballo et al., 1999; Choudhury et al., 2008). The outer core OS of *P. aeruginosa* has a unique feature; it exists in two structurally distinct glycoforms, called “uncapped” and “capped” (King et al., 2009). These two glycoforms basically differ in position and linkage of an L-Rha residue in each structure. The capped glycoform is covalently attached to O-Ag on L-Rha8 that is 1,3-linked to GlcI, whereas the uncapped glycoform cannot be substituted with O-Ag and it contains an L-Rha8 that is 1,6-linked to GlcII (Figure 4). The presence of this L-Rha8 likely causes steric hindrance and prevents attachment of O-Ag to the uncapped core OS; it is also possible that this 1,6-linked L-Rha cannot be recognized by WaaL (O-Ag ligase) as part of the core OS receptor during the O-Ag ligation process.

Another variability in sugar composition of core OS is the presence or absence of a fourth Glc residue (GlcVI) in the uncapped core OS; among the core OSs of the 20 IATS serotypes, GlcVI is present in only nine (O2, O5, O7, O8, O10, O16, O18, O19, and O20; De Kievit and
for a number of bacteriophages and pyocins (Jarrell and Kropinski, 1981a,b; Temple et al., 1986; Yokota et al., 1994; Michel-Briand and Baysse, 2002). For more details, see the section in this review on the role of LPS in host–pathogen interactions. In earlier studies of \textit{P. aeruginosa} LPS biosynthesis, bacteriophages E79 and 2 Lindberg were used to select for spontaneous rough mutants that also had truncation within the core OS (Jarrell and Kropinski, 1981a,b; Dasgupta et al., 1994). Two varieties of outer core truncations providing resistance to bacteriophages have been described, more specifically, the outer core of these structures in the mutant bacterial strains were composed of either (i) the first residue of outer core, GalN (representing the deepest truncation of core OS structure ever found in \textit{P. aeruginosa}) or (ii) a disaccharide GalN–GlcII (Masoud et al., 1994; Sadovskaya et al., 1998; Sanchez Carballo et al., 1999). Similarly, pyocins (i.e., “defective phages”) produced by \textit{P. aeruginosa}, that are able to kill susceptible cells of the same species but are unable to replicate within cells (Michel-Briand and Baysse, 2002), have been used to select for spontaneous rough mutants aiming to generate ones with truncations within the core region (Koval and Meadow, 1977). Another strategy to search for strains with core OS truncations has been to select for spontaneous mutants that would be resistant to a cationic antibiotic; our group used increasing concentrations of gentamicin to select for drug-resistant mutants (Dasgupta et al., 1994). However, the core OS structures from these gentamicin and pyocin-resistant strains have yet to be elucidated, and as such their chemical compositions remain unknown.

Lam, 1994; Bystrova et al., 2006). Additionally, a variable non-sugar substitution found in outer core OS is O-acetylation. \textit{P. aeruginosa} strains display diversity in the degree of O-acetylation ranging from none to five acetyl groups (Bystrova et al., 2006). Since O-acetylation is non-stoichiometric, attempts to precisely identify and quantify all acetylated residues have been difficult. Thus far, only two specific residues have been found to be O-acetylated: Rha, particularly when this residue is not further linked to GlcII, and GlcII, as observed in a number of \textit{P. aeruginosa} mutants with defects in core OS biosynthesis producing truncated core OS (Bystrova et al., 2006; Poon et al., 2008).

![Diagram of LPS structures](https://example.com/lps_structures)

**FIGURE 4 | Structures of uncapped and capped glycoforms.** The structures of the uncapped core oligosaccharide (A), and capped core oligosaccharide (B) are depicted. All the sugars shown have α configuration unless otherwise indicated. Asterisks depict variable substitutions, acetylation sites are shown since they have not been precisely identified. Ala, alanine; Cm, carbamoyl; Etn, ethanolamine; GalN, 2-amino-2-deoxy-galactose (galactosamine); Glc, glucose; Hep, L-glycero-D-manno-heptose; Kdo, 3-deoxy-D-manno-oct-2ulosonic acid; Rha, rhamnose. Adapted from King et al. (2009).

**SPONTANEOUS CORE TRUNCATIONS**

As mentioned above, phosphorylation of inner core heptoses is essential for \textit{P. aeruginosa} cells. Each constituent of the entire inner core seems to be required for viability of \textit{P. aeruginosa}; thus far all of the truncated OS structures that have been examined exhibited a complete inner core, with defects only found in the outer core residues (Sadovskaya et al., 1998; Sanchez Carballo et al., 1999; Kooistra et al., 2003; Choudhury et al., 2005). Preliminary observations also suggested that GaN, the first residue of the outer core, might be essential for viability. This assumption was based on the inability to construct a mutant of \textit{wapG}, which encodes a glycosyltransferase presumably responsible for transferring a GaN residue to the Hep residue in the core (Matewish, 2004). A spontaneous mutation causing truncation of core OS under a selective pressure could be a significant source of core diversity. Core OS has been reported to be a receptor for a number of bacteriophages and pyocins (Jarrell and Kropinski, 1981a,b; Temple et al., 1986; Yokota et al., 1994; Michel-Briand and Baysse, 2002). For more details, see the section in this review on the role of LPS in host–pathogen interactions. In earlier studies of \textit{P. aeruginosa} LPS biosynthesis, bacteriophages E79 and 2 Lindberg were used to select for spontaneous rough mutants that also had truncation within the core OS (Jarrell and Kropinski, 1981a,b; Dasgupta et al., 1994). Two varieties of outer core truncations providing resistance to bacteriophages have been described, more specifically, the outer core of these structures in the mutant bacterial strains were composed of either (i) the first residue of outer core, GaN (representing the deepest truncation of core OS structure ever found in \textit{P. aeruginosa}) or (ii) a disaccharide GaN–GlcII (Masoud et al., 1994; Sadovskaya et al., 1998; Sanchez Carballo et al., 1999). Similarly, pyocins (i.e., “defective phages”) produced by \textit{P. aeruginosa}, that are able to kill susceptible cells of the same species but are unable to replicate within cells (Michel-Briand and Baysse, 2002), have been used to select for spontaneous rough mutants aiming to generate ones with truncations within the core region (Koval and Meadow, 1977). Another strategy to search for strains with core OS truncations has been to select for spontaneous mutants that would be resistant to a cationic antibiotic; our group used increasing concentrations of gentamicin to select for drug-resistant mutants (Dasgupta et al., 1994). However, the core OS structures from these gentamicin and pyocin-resistant strains have yet to be elucidated, and as such their chemical compositions remain unknown.
GENETICS OF CORE DIVERSITY DUE TO GLYCOSYLTRANSFERASES

*Pseudomonas aeruginosa* genomes possess a gene cluster (pa4996–pa5012) encoding proteins that are known to be involved in core OS biosynthesis: HldE, MsbA, PA4998, WaaL, WapR, PA5001, PA5002, PA5003, WapH, WapO, PA5006, WapQ, WapR, WaaP, WapG, WaaC, and WaaE. The respective function of each of these proteins (either demonstrated or proposed based on homology comparisons) has been summarized (Table 3; for more detailed description, see a recent review by our group (King et al., 2009). Apart from the presence or absence of the Glc\(^{IV}\) residue as a terminal sugar in uncapped core OS, the structures of core OS among *P. aeruginosa* strains are relatively conserved with no strain-to-strain variability in sugar composition. This observation is consistent with the discovery that the genes in the locus for core biosynthesis are well conserved among *P. aeruginosa* strains whose genomes have been sequenced. The amino acid identity of the gene products in this locus from one strain to the next is relatively high ranging from 77 to 100%. In addition to genes that encode kinases and heptosyltransferases, this cluster contains genes that are involved in the transportation of lipid A and ligation of O-Ag to the core (msbA and waaL, respectively). Two other genes, pa5002 and pa5003 have not been characterized; therefore, their functions are at present unknown.

Our group has reported the characterization of two transferase genes, migA (described above) and waaA, that are important for core OS biosynthesis, but are localized outside the usual core LPS gene locus. The waaA gene encodes a putative Kdo transferase (King et al., 2009). More recently, we have characterized a homolog of migA that is designated as *wapR*. The *wapR* gene turned out to be localized within the core OS gene cluster. Characterization of *WapR* showed that it is responsible for the transfer of L-Rha\(^{a}\), the outer core rhamnose residue that is 1,6-linked to the Glc\(^{IV}\) of the capped core OS (Poon et al., 2008). We hypothesize that the ratio between expression levels of *migA* and *wapR* would influence the amount of capped and uncapped core OS being produced. Studies to understand the regulation of *wapR* are currently underway in our laboratory. Both putative rhamnosyltransferase enzymes, MigA and WapR, presumably utilize the same substrates, TDP-L-Rha as a sugar donor and core OS lacking L-Rha residues as an acceptor molecule. Therefore, they compete for the enzyme substrates and should impact on amounts of capped and uncapped glycoforms being produced. For instance, overproduction of MigA favors synthesis of uncapped core, whereas an elevated level of WapR activity leads to higher amounts of capped core substituted with O-Ag. It is plausible that under certain conditions migA is upregulated while *wapR* is downregulated (or vice versa), which may consequently influence the amount of O-Ag on the cell surface of *P. aeruginosa*.

We recently discovered yet another transferase gene called *wapB*, which shares sequence homologies with both migA and *wapR*. The *wapB* gene encodes a putative 1,2-glucosyltransferase that is required for transfer of terminal Glc\(^{IV}\) to uncapped core OS (Kocíncová et al., 2011). As not all of the *P. aeruginosa* strains possess Glc\(^{IV}\) as a terminal sugar residue in their uncapped core OS, not all of the *P. aeruginosa* strains have the *wapB* gene. For instance, this gene is absent from the sequenced genome of *P. aeruginosa* PA7.

Other variability in the structure of the *P. aeruginosa* core OS arises from the different degrees of phosphorylation (including ethanolamine-phosphate) and acetylation patterns. However, the variability of these phosphatidyl or acetyl substitutions is nonstoichiometric and the genetic elements that account for these minor substitutions are unknown at present. Additionally, it has yet to be determined what role (if any) genetic events such as mutations or mobile element insertions in core OS genes have played in the truncation of core OS, when rough LPS mutants were selected for resistance to bacteriophages/pyocins or antibiotics. Therefore, there are obvious areas concerning core biosynthesis and regulation that require further investigation.

**Table 3 | Genes involved or potentially involved in the core OS biosynthesis of *P. aeruginosa***

| Core Biosynthesis Gene Cluster (PA4996–PA5012) | Proposed/demonstrated function | References |
|------------------------------------------------|---------------------------------|------------|
| *hidE* (pa4996) | Heptose biosynthesis | King et al. (2009) |
| *msbA* (pa4997) | Transport lipid A-core | Ghanie et al. (2007) |
| *pa4998* | Kinase | King et al. (2009) |
| *waaL* (pa4999) | O-antigen ligase | Abeyrathne et al. (2005), Abeyrathne and Lam (2007) |
| *wapR* (pa5000) | Glycosyltransferase (Rha\(^{a}\)) | Poon et al. (2008) |
| *pa5002* | Glycosyltransferase | King et al. (2009) |
| *pa5003* | Unknown | |
| *wapH* (pa5004) | Glycosyltransferase (Glc\(^{IV}\)) | Matewish (2004) |
| *wapQ* (pa5005) | Carbamoyltransferase | King et al. (2009) |
| *pa5006* | Kinase | King et al. (2009) |
| *wapQ* (pa5007) | Heptose kinase | Walsh et al. (2000) |
| *wapP* (pa5008) | Heptose kinase | Walsh et al. (2000), To (2006) |
| *waaP* (pa5009) | Heptose kinase: position 4 of Hepl | Walsh et al. (2000), Zhao and Lam (2002), Zhao et al. (2002) |
| *wapG* (pa5010) | Glycosyltransferase (GalN) | |
| *waaA* (pa5011) | Glycosyltransferase (Hep\(^{a}\)) | De Kievit and Lam (1997) |
| *waaA* (pa5012) | Glycosyltransferase (Hep\(^{a}\)) | De Kievit and Lam (1997) |

**GENES LOCATED OUTSIDE OF THE CORE BIOSYNTHESIS CLUSTER**

| PA4996 | Glycosyltransferase (Glc\(^{IV}\)) | Kocíncová et al. (2011) |
| PA4997 | Glycosyltransferase (Rha\(^{a}\)) | Poon et al. (2008) |
| PA4998 | Glycosyltransferase: (Kdo\(^{a}\), Kdo\(^{b}\)) | King et al. (2009) |

* Indicates genes involved in cellular processes other than core biosynthesis. Bold font indicates that biochemical or genetic evidence of involvement in *P. aeruginosa* core biosynthesis exists.

**STRUCTURE AND FUNCTION DIVERSITY OF LIPID A**

The lipid A domain of LPS is a glucosamine-based lipid that anchors the LPS in the bacterial outer membrane (OM). The structure of *P. aeruginosa* lipid A has been studied in great detail using both MS and NMR techniques, and it has been reviewed by several groups (Knirel et al., 2006; King et al., 2009; Moskowitz and Ernst, 2010). Similar to lipid A of other Gram-negative bacteria, *P. aeruginosa*
lipid A is composed of a diglucosamine biphosphate backbone [4-P-β-d-GlcNβ(1 → 6)-α-d-GlcNβ(1 → P)], and O- and N-linked primary and secondary fatty acids. The major structural differences of lipid A are observed in the number, the position, the nature of the linked acyl groups, and the modification of the phosphate groups (Figure 5). For example, the chain lengths of the fatty acids attached to lipid A of P. aeruginosa (C12/C14) are shorter than those of E. coli and Salmonella enterica sv. Typhimurium (C13/C16; Trent, 2004). Heterogeneity of lipid A structure has also been observed within different P. aeruginosa strains, which arises from different growth conditions and isolation sources.

The lipid A of laboratory-adapted P. aeruginosa strains (including PAO1, PAK, and PA14) grown in rich medium was found to exhibit a hexa-acylated form (~25%; Figure 5 lipid A form A) or a penta-acylated form (~75%; Figure 5 lipid A form B; Bhat et al., 1990; Kulshin et al., 1991; Ernst et al., 1999; Moskowitz et al., 2004). The hexa-acylated form is a symmetrically acylated structure, with each sugar containing an N-linked 12:0(3-OH) [1-3-hydroxylauroyl] group at the 2 position, an O-linked 10:0(3-OH) [1-3-hydroxycosanoyl] group at the 3 position, and a secondary 12:0 (lauroyl) group linked to the 3-OH group of the primary N-linked 12:0(3-OH) at position 2 (lipid A form A). Either or both of the secondary acyl groups could be non-stoichiometrically 2-hydroxylated (Figure 5 red labeled), converting 12:0 to 12:0(2-OH) (Kulshin et al., 1991). The penta-acylated lipid A form, which is the predominant (~75%) lipid A form of laboratory P. aeruginosa strains, lacks the primary O-linked acyl substitution at position 3 of the GlcNβ (Figure 5, lipid A form B; Kulshin et al., 1991).

As has been demonstrated in enteric bacteria such as S. enterica (Guo et al., 1997, 1998), P. aeruginosa lipid A adopts different modifications in response to different environmental conditions encountered such as different growth medium and during human infection. For example, clinically isolated P. aeruginosa produce lipid A with modified structures. Lipid A from P. aeruginosa isolated from patients with acute clinical infections (such as blood, ear, eye, and urinary tract) or chronic non-CF infections such as bronchiectasis has a penta-acylated form that lacks the secondary acyl group at position 2 of the sugar GlcNβ (Figure 5 lipid A form C; Ernst et al., 1999, 2003). Three different types of modifications have been observed for the lipid A structure of P. aeruginosa isolates from CF patient airways. The first type of modification is observed in the majority of CF P. aeruginosa isolates: the addition of an O-linked secondary palmitate (16:0) to the 3-OH group of 10:0(3-OH) at the 3′ position in sugar GlcNβ (Figure 5 lipid A form D, blue; Guo et al., 1997, 1998). Less than half of CF isolates examined by the Miller group showed a second type of modification to lipid A, the non-stoichiometric addition of aminoarabinose (4-amino-4-deoxy-L-arabinose, Ara4N) to either or both of the

![Figure 5](https://www.frontiersin.org)
terminal phosphates (Figure 5 lipid A form D, green; Ernst et al., 2007). A hepta-acylated lipid A (Figure 1 lipid A form E) found only in P. aeruginosa isolates from CF patients with very severe lung disease resulted from a third type of modification: the retention of the primary O-linked 10:0(3-0H) group at the 3 position of GlcNAc (Ernst et al., 2003, 2007). Interestingly, the lipid A modifications observed in CF airway isolates (i.e., addition of palmitate and aminoarabinose) could be induced in laboratory strains, strains isolated from the environment, or non-CF clinical isolates under certain growth conditions including low Mg\(^{2+}\) concentration or the presence of cationic antimicrobial peptides (CAMPs) such as polymyxin B, indicating the enzymatic pathways for these modifications were intact and inducible in these strains (Bhat et al., 1990; Ernst et al., 1999, 2003; Bedoux et al., 2004).

PROPOSED BIOSYNTHESIS AND MODIFICATION PATHWAY OF LIPID A

Over the past 20 years, the biosynthesis pathway of lipid A has first been experimentally determined in E. coli as well as other enteric bacteria and reviewed by Raetz and Whitfield (2002) and Trent (2004). Bioinformatics analysis indicates that the general scheme of lipid A biosynthesis pathways are highly conserved among Gram-negative bacteria, although various modifications are present in different bacteria (Trent, 2004). In P. aeruginosa, in contrast to the extensive and detailed structural elucidation of lipid A, most aspects of the biosynthesis pathway have not been investigated experimentally, perhaps due to the fact that the genetics of lipid A biosynthesis were so thoroughly studied in enteric bacteria and that the pathway is relatively conserved. The current assumption about the biosynthesis pathways is mainly based on the identification of homologs to E. coli genes. Since all of the corresponding homologous genes encoding enzymes required for the biosynthesis of the important lipid A structure precursor lipid IV\(_a\) (Figure 5); including the primary acyltransferases LpxA and LpxD, the nucleotide LpxH, the disaccharide synthase LpxB, and the kinase LpxK) were identified in the genome of P. aeruginosa, the biosynthesis of the lipid IV\(_a\) analog of P. aeruginosa is thought to be executed in a conserved manner as in E. coli and S. enterica. A detailed description of the proposed biosynthesis pathway of lipid IV\(_a\) was recently reviewed by our group (King et al., 2009). Not surprisingly, the lipid A primary acyl chain length of different bacteria is determined by the chain length preference of the acyltransferase LpxA and LpxD (Dotson et al., 1998; Wyckoff et al., 1998).

Different forms of lipid A could then be synthesized and modified from lipid IV\(_a\) by different enzymes depending on the environmental conditions (a proposed pathway is depicted in Figure 5). Two secondary acyltransferases (LpxL1 and LpxL2, encoded by paa3243 and paa0011, respectively) are proposed to transfer the secondary lauroyl groups to lipid IV\(_a\) to form the hexa-acyl lipid A (Figure 1, lipid A form A; Mohan et al., 1994; King et al., 2009). However, unlike in E. coli and S. enterica sv. Typhimurium, the addition of the first core sugar 3-deoxy-d-manno-oct-2-ulosonic acid (Kdo) to the lipid IV\(_a\) is not strictly necessary for the attachment of the secondary acyl groups in P. aeruginosa (Goldman et al., 1988). The non-stoichiometric 2-hydroxylation of the secondary lauroyl group is proposed to be catalyzed by corresponding homologs of S. enterica sv. Typhimurium LpxO (Gibbons et al., 2000, 2008). In P. aeruginosa, two putative LpxO homologs (LpxO1 and LpxO2, encoded by paa4512 and paa1936, respectively) were present; presumably each would hydroxylate one of the two secondary acyl chains (King et al., 2009). A newly identified deacylase enzyme PagL is believed to be involved in the removal of the primary 10:0(3-0H) group at the 3 position of GlcNAc of the hexa-acylated lipid A (Figure 5, lipid A form A) to form the penta-acylated lipid (Figure 5, lipid A form B), and to convert the hepta-acylated lipid A (Figure 5, form E) to hexa-acylated lipid A (Figure 5, form D) in CF isolates (Ernst et al., 2006). As stated earlier, the lipid A from isolates of patients with acute infection or non-CF chronic infection lacks the secondary C\(_{12}\) acyl chain at GlcNAc (Figure 5, lipid A form C; Ernst et al., 1999, 2003). This could be due to the lack of expression of one of the secondary acyltransferases lpxL, or the later removal of the secondary C\(_{12}\) acyl group from lipid A (Figure 5, form A) by a novel deacylase enzyme. The OM palmitoyl transferase protein PagP was originally identified in Salmonella and is responsible for the addition of palmitate to lipid A (Guo et al., 1998). A PagP homolog in P. aeruginosa is proposed to catalyze the addition of palmitate to lipid A forms A and B to synthesize lipid A forms D and E that are found in CF patient airway isolates, respectively (Trent, 2004). The hexa-acylated lipid A (form D) could also result from the removal of the primary 10:0(3-0H) group from the hepta-acyl lipid A (form E) by the deacylase enzyme PagL. It is not known which of the two steps, i.e., the addition of palmitate or the removal of the primary 10:0(3-0H), occurs first. However, it has been confirmed that the occurrence of the hepta-acylated lipid A (form E) found only in isolates from CF patients with severe infection was due to the loss of PagL function (Ernst et al., 2006). Although an in vitro assay indicated that a PagP-like enzyme was present in the membranes (Trent, 2004) of P. aeruginosa, to date, no pagP homolog has been identified in the genome (Stover et al., 2000). This could be due to low sequence similarity between these homologs. Similar to Salmonella, the addition of palmitate is regulated by the PhoP–PhoQ two-component regulatory system, as a phoP null mutant of P. aeruginosa is not able to synthesize palmitate-modified lipid A under inducing conditions for wildtype strains (such as limited Mg\(^{2+}\); Ernst et al., 1999). The biosynthesis pathway of aminoarabinose and its addition to lipid A has been well characterized in E. coli and S. enterica (reviewed by Trent, 2004 and Bishop, 2005). The gene agiL (also called pmrF) and the pmrHFIJKLM (also referred to as the arnBCADETF) gene cluster have been identified and characterized to encode enzymes required for the biosynthesis of aminoarabinose as well as its transport across the IM and its ligation to lipid A (Trent, 2004; Bishop, 2005). These homologous genes were identified in a single cluster in the genome of P. aeruginosa PAO1 (pa3552–pa3559; Stover et al., 2000). As in E. coli and S. enterica, it was found that both PhoP–PhoQ and another two-component regulatory system PmrA–PmrB are involved in the regulation of the expression of these genes and the addition of aminoarabinose to lipid A (Moskowitz et al., 2004; Moskowitz and Ernst, 2010). Both systems can respond to Mg\(^{2+}\)-limiting conditions (Mcphee et al., 2003, 2006); moreover, the PmrA–PmrB system can also respond to the presence of CAMPs including polymyxin B causing activation of the transcription of the pmr gene cluster (Mcphee et al., 2003).


**BIOLOGICAL IMPLICATIONS OF LPS DIVERSITY IN P. AERUGINOSA**

**COLONIZATION AND PERSISTENCE**

Physiological adaptability and metabolic plasticity have long been hallmarks of *P. aeruginosa* survival, traits that were consistent with the annotated genome sequence for strain PAO1, which revealed a wealth of predicted genes involved in various synthesis, assembly, metabolic, regulatory, and pathogenesis paradigms (Stover et al., 2000). This genomic abundance reflects the remarkable versatility of this bacterial species to colonize and persist in diverse niches ranging from different environmental settings to host animal tissues (Wolfgang et al., 2003). While several characteristics of *P. aeruginosa* facilitate initial substratum colonization and later-stage persistence, each is either directly or indirectly affected by variations in the phenotype of the LPS present on the cell surface.

In the aqueous environment, cells of *P. aeruginosa* swim with the aid of a polar flagellum (Feldman et al., 1998), but this mode of motility was impaired in mutants that lacked wildtype smooth LPS due to defects in either core OS biosynthesis, yielding truncated core OS (Lindhout et al., 2009), or ligation of OSA and CPA to the core OS, in the case of a *wapA* mutant (Abeyrathne et al., 2005). Swarming motility of *P. aeruginosa* cells, which is the process of translocation on semi-solid surfaces dependent on flagella, type IV pili, and rhamnolipid (a wetting agent produced by *P. aeruginosa*; Kohler et al., 2000), was similarly compromised in the aforementioned mutants with truncated core OS. The motility defects in the mutants were found to be a result of changes in cell-surface properties such as cell-to-cell and cell-to-substratum adhesion forces. Both types of adhesive forces became significantly increased in the mutants, as measured by AFM. This is in contrast to the initial thoughts that LPS defects might have somehow affected flagella-driven motility. In fact, when the mutant cells were examined by standard light microscopy, and the swimming speed of individual cells was quantified, no difference could be discerned between the mutant and the wildtype parent strain. The increased adhesive properties retarded all outward motility from the population of *P. aeruginosa* cells (Lindhout et al., 2009).

Biofilm formation is the preferred mode of growth for *P. aeruginosa* cells clinging to rocks in fluvial streams or surviving in the lungs of CF patients in a chronic infection situation (Hall-Stoodley et al., 2004). Following planktonic growth of *P. aeruginosa*, attachment to a substratum is a prerequisite for establishing long-term colonization at a particular site to eventually adapt to a biofilm mode of growth. Biofilms are intricate surface-associated bacterial communities that confer survival advantages to the cells residing within and for which flagellar-mediated motility is important for their maturation in *P. aeruginosa* (O’toole and Kolter, 1998; Klausen et al., 2003). Consistent with the motility defects described above, the same mutants of *P. aeruginosa* lacking complete core OS and the distal CPA and OSA moieties were found to form biofilms with significant differences in mechanical and structural properties when compared to those of wildtype bacteria. This evidence was collected from a variety of quantitative measurement studies to determine changes in ultrastructures, biophysical properties, cell–cell adhesion forces, and viscoelasticity of mutant and wildtype strains using a technique called microbead force spectroscopy (Lau et al., 2009a). Several bacterial strains including knockout mutants disrupted in *migA*, *wapR*, and *rmlC*, respectively, and with defined core OS truncation characteristics were compared to their wildtype PAO1 parent strain in these studies. Significant changes were observed in cell mechanical properties among the mutant strains compared to the wildtype PAO1. The functions of *migA* and *wapR* have been described earlier; the *rmlC* gene is responsible for TDP-4-Rha biosynthesis and hence a *rmlC* mutant produces a defective core OS truncated at the Rhaα and Rhaβ residues in the two glycoforms of the core OS (Figure 4). The data from these studies revealed that truncation of core OS enhanced both adhesive and cohesive forces by up to 10-fold, whereas changes in instantaneous elasticity were correlated with the presence of O-Ag. Using AFM to raster-scan bacterial cells in air in contact mode for each of the aforementioned four strains showed differences in the texture of the surface “smoothness.” Interestingly, LPS-“smooth” strain wildtype PAO1 with O-Ag exhibits rougher surface topography than rough strains, i.e., the mutant bacterial strains of *migA*, *wapR*, and *rmlC* (Figure 6). Using confocal laser scanning microscopy to quantify biofilm structural changes in these mutants, we observed that textural parameters varied with adhesion or the inverse of cohesion, while areal and volumetric parameters were linked to adhesion, cohesion, or the balance between them. Microcolonies formed by cells of the wildtype PAO1 had round perimeters, while the microcolonies formed by the mutant strains had more irregular edges (Figure 7; Lau et al., 2009b). These studies support the importance of O-Ag in the formation of cellular structures and the physiology of *P. aeruginosa* in a biofilm mode of growth. In a study by Ivanov et al. (2011), they showed that changes in the relative proportion of OSA modalities as well as the outright loss of OSA result in reduced virulence of *P. aeruginosa* consistent with diminished surface adhesive forces, further supporting the role of LPS-mediated adhesion in *P. aeruginosa* persistence. The observations made in these recent studies substantiated an earlier report in which rough mutants of *P. aeruginosa* lacking OSA had an LD₅₀ that was 1000X higher than that of a wildtype strain in a mouse infection model (Cryz et al., 1984).

**PROPERTIES OF P. AERUGINOSA LPS THAT INFLUENCE ITS INTERACTION WITH THE DEFENSE SYSTEMS OF THE HOST**

The genomic stability of *P. aeruginosa* is such that environmental isolates have been found to maintain the potential for pathogenicity through the conservation of numerous virulence-associated genes required for host infection (Wolfgang et al., 2003). Undoubtedly, these genetic traits contribute to the success of the bacterium as an opportunistic pathogen. In humans, opportunistic infection by *P. aeruginosa* is often seen in patients with varying degrees of compromised host defenses such as those suffering from damaged corneal lenses, cancer, AIDS, severe burn wounds, and CF (Lyczak et al., 2000). As with the more environmental physiological aspects described above, LPS of *P. aeruginosa* plays multiple key roles in the interaction between the pathogen and the infected host through direct engagement or evasion of innate and adaptive immune system responses.

During the course of infection, it is a general understanding that the humoral adaptive immune response has a lag phase of between 5 days to a week from initial infection detection; in that time period, the innate immune system plays an essential role in engaging and...
managing the infection through a variety of mechanisms. One such mode of action involves the recognition of pathogen-associated molecular patterns (PAMPs) by various pattern recognition receptors in the host (Janeway and Medzhitov, 2002). Arguably, the most widely described reaction of the innate immune system to LPS is that of endotoxicity resulting from recognition of the distal lipid A moiety by the host Toll-like receptor 4 (TLR4) complex, which consists of the TLR4 receptor, the co-receptor component MD-2, and the GPI-linked LPS binding protein CD14 (Da Silva Correia et al., 2001; Hajjar et al., 2002; Palsson-Mcdermott and O’neill,
the TLR4 complex (Hajjar et al., 2002). In contrast, LPS with penta-acylated lipid A is less potent. Compared to enterobacterial lipid A which has the hexa-acyl form with longer chain fatty acids (C14 versus C10–C12), *P. aeruginosa* lipid A showed a significantly lower toxicity in an experimental animal model infection (Takada and 2004; Miller et al., 2005). As discussed earlier, during adaptation of *P. aeruginosa* to the CF airway, the normally penta-acylated lipid A of the bacterium (Figure 5, lipid A form B) was modified to a hexa-acylated form (Figure 5, lipid A form D; Ernst et al., 2003), becoming a more robust proinflammatory stimulus mediated by the TLR4 complex (Hajjar et al., 2002). In contrast, LPS with penta-acylated lipid A is less potent. Compared to enterobacterial lipid A which has the hexa-acyl form with longer chain fatty acids (C14 versus C10–C12), *P. aeruginosa* lipid A showed a significantly lower toxicity in an experimental animal model infection (Takada and

![Image](image_url)

**FIGURE 7** Images from confocal laser scanning microscopy analyses of *P. aeruginosa* cells that illustrate the changes in biofilm structure resulting from truncation in the LPS core in mutant strains as comparing to the wildtype bacteria. Average projections (top panels) and midpoint cross sections (bottom panels) of representative microcolonies of (A) wildtype strain PAO1, (B) migA mutant, (C) wapR mutant, and (D) mliC mutant are shown. Reproduced from Lau et al. (2009b), with permission from Copyright Clearance Centre.
Kotani, 1992; Backhed et al., 2003). On the surface, direct stimulation of the innate immune system by *P. aeruginosa* would appear to be a curious survival strategy for a pathogen that establishes chronic infections in CF patients; however, this strongly points to an as-yet-to-be-determined survival advantage for *P. aeruginosa* conferred by constant inflammation of the airway epithelium. Conversely, this sequence of proinflammatory responses has been recognized as a direct contributing factor to the gradual deterioration of lung capacity in CF patients with *P. aeruginosa* infection (Hoiby et al., 2010).

Modifications to the lipid A domain of LPS may also confer other survival advantages. The host innate immune system produces a wide range of CAMPs, and some of them (such as cecropin and human defensin) directly interact with bacterial membranes (Hancock and Diamond, 2000). The addition of positively charged aminoarabinose to the phosphate groups of lipid A reduces the permeability of the OM or the affinity of lipid A for those CAMPs, and has clearly been shown to promote *P. aeruginosa* resistance to CAMPs; thus, such modifications may increase bacterial survival rates during host colonization (Ernst et al., 1999, 2000; Moskowitz et al., 2004; Trent, 2004).

The principal genetic defect resulting in altered physiology of patients with CF is a loss of function of the CF transmembrane conductance regulator (CFTR), a chloride ion channel important for maintaining the optimal viscosity of mucus membranes. Impaired CFTR function results in the production of mucus that is highly viscous, creating an ecological niche in the respiratory tract similar to a biofilm matrix, ideal for colonization by *P. aeruginosa* (Gibson et al., 2003). However, in addition to the increased viscosity of the lung environment, a deficiency in binding of *P. aeruginosa* by CFTR also contributes to pathogenesis. Uptake of bacteria by airway epithelial cells has been implicated as a key mechanism for clearance of *P. aeruginosa* and CFTR was shown to play a role as a receptor and contribute to epithelial internalization (Bajmoczi et al., 2009). Murine cells lacking recombinant human CFTR, as well as those expressing the predominant ΔF508 CFTR mutant, displayed significant impairment in the binding and endocytosis of *P. aeruginosa*, compared to expression of the wildtype human protein (Pier et al., 1997), with the core OS portion of the LPS molecule of the bacterium implicated as the ligand responsible for uptake (Pier et al., 1996). As such, in individuals with wildtype CFTR, clearance of *P. aeruginosa* is mediated by uptake of the infecting bacteria via binding to CFTR and subsequent internalization. However, studies of experimental eye infections demonstrated that CFTR-mediated internalization of *P. aeruginosa* by corneal epithelial cells is essential to the development of keratitis (Zaidi et al., 1999), a potentially severe infection of the ocular lens that is often developed by individuals that frequently wear contact lenses (Robertson et al., 2007). In this scenario, the core OS-mediated interaction of CFTR with *P. aeruginosa* is detrimental to the human host rather than beneficial.

Activation of the complement cascade, which is comprised of more than 30 serum and cell-surface proteins, is another response of the innate immune system essential to combating infection. Three known mechanisms lead to activation of the complement system, namely, the classical (CCP), alternative (ACP), and mannann-binding lectin (MBL) pathways, each of which results in activation of C3 to yield C3b; in turn, this can lead to opsonization of pathogens and engulfment by phagocytes or surface deposition of complement components C5b, C6, C7, C8, and C9 to form the lytic membrane-attack complex (Dunkelberger and Song, 2010). In a murine model of *P. aeruginosa*-induced pneumonia, C3-deficient mice displayed higher mortality rates than C3-expressing mice. The C3-deficient mortality rates were similar to those observed for mice deficient in ACP-specific factor B, but not for mice deficient in CCP- and MBL-specific C4, indicating a critical role for the ACP in controlling initial *P. aeruginosa* pulmonary infection. Phagocytic cells from both the C3-expressing and C3-deficient mice displayed a reduced capacity in vitro to bind and take up *P. aeruginosa* in the presence of C3-deficient serum compared to C3-containing serum, suggesting that phagocytic clearance of the bacterium via C3-mediated opsonization constitutes a part of the protection afforded by the ACP against *P. aeruginosa* infection (Mueller-Ortiz et al., 2004).

Clinical isolates of *P. aeruginosa* from CF patients, deficient in long-chain O-Ag production, were found to be sensitive to pooled normal human serum (PNHS) when compared to subcultured derivatives of the parental strains with restored O-Ag biosynthesis, which were found to be serum resistant (Schiller et al., 1984). The same derivative strains with restored O-Ag biosynthesis were subsequently shown to activate (i.e., consume) more complement from PNHS per bacterial organism than their O-Ag-deficient parental counterparts. Serum-sensitive strains were observed to accumulate more C3 than serum-resistant strains, with the C3 accumulation also occurring more rapidly in the former; both of these phenotypes were reduced upon prior treatment of the PNHS with ethylene glycol tetraacetic acid (EGTA, which blocks the CCP), suggesting that the CCP is important for maximal deposition of C3. Analysis of the form of C3 deposited on the surface of the various serum-sensitive and serum-resistant strains indicated equivalent amounts of C3b and iC3b (proteolytically inactivated C3b) in the former, while the latter contained mostly iC3b (Schiller et al., 1989). Taken together, these results indicate that complement-mediated responses to *P. aeruginosa* are important parts of the infection resolution mechanism, which is directly affected by the presence of O-Ag on the bacterial surface.

Sugar-binding proteins known as lectins, of which there are several classes, are also known to play a role in the innate immune response to *P. aeruginosa* infection. Surfactant proteins A (SP-A) and D (SP-D) are part of the calcium-dependent, collagenous C-type lectin family, containing carbohydrate-recognition domains. In the lung, they are secreted into the mucus lining the respiratory tract, agglutinating, neutralizing, and opsonizing (Holmskov et al., 2003). SP-A and SP-D have been shown to bind both rough and smooth strains of *P. aeruginosa* (Butler et al., 2003), with SP-D interacting specifically with LPS (Kishore et al., 1996); incidentally, both surfactant proteins have been shown to be important for clearance of *P. aeruginosa* (Levine et al., 1998; Mariencheck et al., 1999; Restrepo et al., 1999; Ni et al., 2005) but are present at reduced levels in the lungs of CF patients (Postle et al., 1999). In a separate investigation, the signal from biotin-labeled LPS from
P. aeruginosa was shown to overlap with that obtained with antibodies against the S-type lectin galectin-3 via Western blotting of human corneal epithelial proteins, suggesting binding of the LPS by galectin-3. Additionally, pretreatment of scarified corneas in whole-eye culture with mAbs against either the outer core OS from P. aeruginosa or galectin-3 was found to inhibit binding of the bacterium to the cornea (Gupta et al., 1997). Coupled with the importance of SP-A and SP-D described above, the capacity of galectin-3 in binding of P. aeruginosa indicates a significant role for LPS-dependent lectin-mediated clearance of P. aeruginosa infection via the innate immune response.

It has been demonstrated that the type III secretion system (T3SS) of the bacterium mediates direct virulence-promoting primary interactions between P. aeruginosa and the host. This pathogenesis mechanism involves the highly regulated contact-dependent insertion of a needle-like injector assembly into a host cell by the surface-associated bacterium, followed by the extrusion of four principal effector proteins (ExoS, T, U, Y) from the bacterium directly into the cytosol of the host cell. These type III effector proteins display a wide range of substrate specificities and in turn perturb a diverse range of host cell functions (Hauser, 2009). Intriguingly, upon comparison of isogenic LPS mutants to wildtype P. aeruginosa, the relative levels of expression of exoS and exoT (encoding effectors) exxA (encoding the positive T3SS regulator), and pcrV (encoding the injection-facilitation component) displayed marked increases in the absence of OSA. Even more pronounced increases in gene expression were observed in the simultaneous absence of OSA and CPA, suggesting a strong regulatory association between the presence/absence of capped core OS and T3SS effector expression (Augustin et al., 2007). Intracellularly, the production of ExoS and PcrV were found to be reflective of upregulated gene expression. P. aeruginosa mutants defective in OSA and CPA were also found to confer increased cytotoxicity in vitro and in vivo in lung epithelial cells and a murine model of infection, respectively (Augustin et al., 2007), indicating that the presence of O-Ag is an important factor contributing to the regulation and expression of T3SS effectors.

Notwithstanding either innate immune system interplay or T3SS activity, the host–pathogen interaction dynamic is also applicable with P. aeruginosa as the host and bacteriophage as the infectious agents. To date, multiple motifs on the LPS molecule of P. aeruginosa have been identified as receptors for various bacteriophages. As discussed earlier, bacteriophage A7 was found to bind and hydrolyze the β-Rha of CPA in a P. aeruginosa mutant devoid of OSA (Rivera et al., 1992). In addition, specificity for OSA has also been identified through examination of bacteriophage D3; lysogenization with the phage resulted in a loss of adsorption with the same phage during subsequent infection cycles (Holloway and Cooper, 1962), a phenotypic alteration later identified to be a result of acetyl-group addition at the β-Fuc residue as well as alteration of the linkage stereochemistry between OSA repeat units from α1 → 4 to β1 → 4, resulting in serotype conversion. Lysogenization by D3 also impeded the binding ability of the unrelated LPS-specific phage E79 (Kuzio and Kropinski, 1983). The core OS of the bacterium is not only an important ligand for interaction with eukaryotic cells, but it is also a receptor for bacteriophages. Phages ϕCTX, ϕPLS27, E79, and H22 have all been shown to recognize the core OS domain of P. aeruginosa LPS (Meadow and Wells, 1978; Jarrell and Kropinski, 1981b; Temple et al., 1986; Yokota et al., 1994).

Analogous to the binding of P. aeruginosa by bacteriophage is the interaction of pyocins with the surface of the bacterium. Pyocins are chromosomally encoded by over 90% of P. aeruginosa strains and are inducible upon treatment with certain mutagenic agents. Depending on the type of pyocins, which might behave like lytic proteins or phage-like particles, they can disrupt the membranes of both related and unrelated bacteria, but which normally have no effect on the host strain (Michel-Briand and Bayssé, 2002). Three principal pyocin types have been described, with the R-type (rod-shaped) pyocins closely resembling the contractile tails of bacteriophage. Specifically, R-type pyocins are related to ϕCTX bacteriophages (Hayashi et al., 1994), with core OS serving as a receptor. Five subtypes of R-type pyocins (R1–R5) have been described, each with different killing spectra. Using a collection of well-defined LPS-deficient P. aeruginosa knockout mutants generated by our group, the RhaΔ, GlcΔ, and GlcΔ/GlcΔIII core sugars (Figure 5) were identified as receptors for R1, R2, and R5 R-type pyocins, respectively. Therefore, spontaneous truncations of the core OS provide resistance to some of the R pyocins. The presence of OSA on the surface of P. aeruginosa was demonstrated to provide a “shield” against pyocin-mediated killing (Köhler et al., 2010).

TREATMENT AND PREVENTION OF INFECTION

Upon infection with P. aeruginosa, leading treatment regimens often involve doses of antibiotics such as oral ciprofloxacin, or aerosolized drugs including TOBI® (inhaled tobramycin therapeutic), or colymycin (inhaled colistin–lysin) that are currently being used to treat CF patients nowadays (Anderson, 2010). Tobramycin belongs to the family of aminoglycosides that inhibits bacterial protein biosynthesis via irreversible binding to the 30S bacterial ribosome (Edson and Terrell, 1999). Aminoglycosides such as gentamicin have also been shown to disrupt the cell envelope of P. aeruginosa (Martin and Beveridge, 1986; Walker and Beveridge, 1988), leading to lysis of the bacterium (Kadurugamuwa et al., 1993a). Irrespective of its mechanism of bactericidal activity, the initial event in gentamicin-mediated treatment of infection involves ionic binding of the drug to the surface of P. aeruginosa. However, comparison of a wildtype strain with isogenic mutants lacking various LPS glycoforms indicated a higher affinity of OMs containing OSA for gentamicin. Furthermore, OSA-expressing strains were more susceptible to killing via gentamicin treatment, with viability reduced almost 50% in strains simultaneously expressing OSA and CPA as a result of higher antibiotic binding than in any other strain (Kadurugamuwa et al., 1993b). The importance of OSA in gentamicin binding has direct implications for the treatment of CF patients, as chronic P. aeruginosa isolates often lose the ability to synthesize OSA, while LPS capped with CPA is maintained (Hancock et al., 1983; Lam et al., 1989).

Given the ability of various phages to specifically and selectively target their respective host bacterium, combined with their overall lack of eukaryotic epitope recognition, the use of phage to control bacterial infection is once again gaining attention in the medical community after a prolonged period of research dormancy in the field (Hanlon, 2007). This is of particular interest in light of the
continued development of widespread resistance to antibiotics, including isolates of *P. aeruginosa*, a bacterium already known to possess high intrinsic antibiotic resistance. As such, with a range of *P. aeruginosa*-specific bacteriophages identified (Ceyssens and Lavigne, 2010), phage therapy could serve as an alternative or complementary method to the treatment and management of infections with this bacterium. One such example is the recently identified LPS-specific phage JG024 which was found to efficiently lyse a wide range of environmental and clinical isolates of *P. aeruginosa* (Garbe et al., 2010).

Rather than intervention through therapeutic measures after an infection has occurred, researchers have pursued prevention of *P. aeruginosa* colonization through the use of immunogenic vaccines to foster sustained adaptive immunity (Stanislavsky and Lam, 1997). Various vaccine formulations have been tested in a range of healthy and compromised individuals, using mixtures of LPS from different serotypes, different LPS components, immunogenic toxins conjugated to LPS components, and inactivated whole-cell preparations, all traditionally administered via injection (Stanislavsky and Lam, 1997; Sedlak-Weinstein et al., 2005). While certain vaccine trials involving whole-molecule LPS preparations have yielded increased anti-LPS antibody titres, most LPS vaccines have been characterized by various toxic side effects, yielding inconsistent results with varying degrees of sustained protection (Sedlak-Weinstein et al., 2005). Furthermore, certain instances of non-*Pseudomonas* microbe replacement have also been identified following testing of LPS vaccines (Pennington et al., 1975; Jones et al., 1979). To circumvent issues of pyrogenticity associated with purified LPS used for injection, lipid A-core OS from *P. aeruginosa* (and three other species) was reconstituted in liposomes and used to immunize rabbits, resulting in a well-tolerated vaccine that induced the production of cross-reactive anti-core OS antibodies against a large panel of pathogenic Gram-negative bacteria expressing both rough and smooth LPS (Bennett-Guerrero et al., 2000); this same liposomal formulation was later demonstrated to reduce the induction of TNF-α production in vitro when compared to the equivalent amounts of the purified LPS constituents (Erridge et al., 2002). As such, liposomal reconstitution may provide a means of reducing the endotoxic effects of LPS vaccine while still maintaining their protective efficacy.

In comparison to immunization with whole-molecule LPS, OSA vaccines are well tolerated. Correlation between the length of the OSA polymer and the production of protective antibodies was demonstrated in mice, with OSA containing over 18 repeat units inducing 50- to 100-fold increases (Macintyre et al., 1986). In CF patients not previously infected by *P. aeruginosa*, immunization with an octavalent OSA vaccine conjugated to exotoxin A was found to elicit high levels of anti-OSA antibodies in the serum (Lang et al., 1995), a trend sustained with yearly immunizations for a decade (Zuercher et al., 2006). These regular immunizations with the octavalent OSA-exotoxin A conjugate vaccine were found to delay the occurrence of colonization and reduce the frequency of chronic infection in young CF patients (Lang et al., 2004), demonstrating promise as a potential commercialized vaccine to prevent *P. aeruginosa* infection. Another method to induce protective immunity against *P. aeruginosa* has been through the generation of recombinant immunogens involving the OSA from certain serotypes. Heterologous expression of *P. aeruginosa* O11 OSA in an attenuated strain of *S. enterica* sv. Typhimurium, followed by intranasal immunization, resulted in the production of OSA-specific antibodies in the serum of immunized mice. Complete protection with the immunization was also provided against respiratory challenges until 6 months post-vaccination as well as against infection from a burn wound, while only partial protection was provided against corneal infection (Digiandomenico et al., 2007). This is a novel method for the stimulation of adaptive immunity against *P. aeruginosa*. However, despite the data from some of these studies demonstrating efficacy of the vaccine formulations in animal models, obvious hurdles must be overcome before the vaccines can be administered to CF patients. One of these would be the development of an effective and accurate diagnostic method to determine whether or not a young CF patient has been colonized by *P. aeruginosa*. Infected patients produce high amounts of antibodies against LPS. Therefore, administering a vaccine to a patient who has previously been colonized by *P. aeruginosa* can have deleterious effects due to the formation of immune complexes between specific antibodies against *P. aeruginosa* surface antigens, such as LPS, and the vaccine antigens. The presence of these immune complexes could lead to serum sickness. Ultimately, the continued pursuit of an effective anti-*P. aeruginosa* vaccine involving LPS (whole or in part) is an important avenue of research that will undoubtedly benefit millions of people suffering from a range of medical conditions.

Considering all of the aforementioned biological implications of LPS diversity in *P. aeruginosa*, it is indisputable that the bacterium follows a very complex infection dynamic, involving a multitude of simultaneous interactions with the host, many of which it can exploit to further pathogenesis and its continued survival upon infection; yet a majority of these virulence mechanisms are affected, either directly or indirectly, by the LPS of the pathogen. As such, continued research into the biosynthesis and assembly of this important cell-surface virulence factor is essential to clearly understanding the colonization process and eventually controlling or inhibiting infection by *P. aeruginosa*.

**CONCLUSION**

In this review, we have provided a comprehensive account of the genetics for the biosynthesis of each of the three regions of *P. aeruginosa* LPS. We have also provided a critical review of the information in the literature pertaining to knowledge of how the variation in producing each region of the LPS can impart diversity to this species. External factors such as temperate bacteriophages that recognize LPS epitopes as receptors, followed by internalization and propagating the viral life cycle, also influence diversity of LPS. All of these different factors account for the heterogeneity of *P. aeruginosa*, not only of the chain length of the O-Ag, but also of the chemical make up that accounts for the different serotypes. We close out the discussion by addressing the importance of LPS in host–pathogen interactions and the strategies that researchers use to target this cell-surface glycolipid to develop effective vaccines or novel antimicrobial interventions against *P. aeruginosa* infections. Suggestions for future research have been integrated throughout the various sections.
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