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Research Paper

The diversity of *Klebsiella pneumoniae* surface polysaccharides

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*Klebsiella pneumoniae* is considered an urgent health concern due to the emergence of multi-drug-resistant strains for which vaccination offers a potential remedy. Vaccines based on surface polysaccharides are highly promising but need to address the high diversity of surface-exposed polysaccharides, synthesized as O-antigens (lipopolysaccharide, LPS) and K-antigens (capsule polysaccharide, CPS), present in *K. pneumoniae*. We present a comprehensive and clinically relevant study of the diversity of O- and K-antigen biosynthesis gene clusters across a global collection of over 500 *K. pneumoniae* whole-genome sequences and the seroepidemiology of human isolates from different infection types. Our study defines the genetic diversity of O- and K-antigen biosynthesis cluster sequences across this collection, identifying sequences for known serotypes as well as identifying novel LPS and CPS gene clusters found in circulating contemporary isolates. Serotypes O1, O2 and O3 were most prevalent in our sample set, accounting for approximately 80 % of all infections. In contrast, K serotypes showed an order of magnitude higher diversity and differ among infection types. In addition we investigated a potential association of O or K serotypes with phylogenetic lineage, infection type and the presence of known virulence genes. K1 and K2 serotypes, which are associated with hypervirulent *K. pneumoniae*, were associated with a higher abundance of virulence genes and more diverse O serotypes compared to other common K serotypes.

Keywords: *Klebsiella pneumoniae*; seroepidemiology; surface polysaccharide; K antigen and O antigen; vaccine target.

Abbreviations: CPS, capsule polysaccharide; ESBL, extended spectrum beta lactamases; MLST, multilocus sequence typing; TPP, thiamine pyrophosphate; UndPP, undecaprenyl-pyrophosphate; LPS, Lipopolysaccharide.

Data statement: All supporting data, code and protocols have been provided within the article or through supplementary data files.

Data summary

Supplementary dataset S1 lists the ENA accession numbers for the 573 publicly available *K. pneumoniae* whole-genome Illumina read sets analysed in this study (http://www.ebi.ac.uk/ena).

Representative *rfb* and *cps* locus sequences have been deposited in GenBank, see Table S2 for the list of GenBank accession numbers (http://www.ncbi.nlm.nih.gov/genbank/).
Introduction

*Klebsiella pneumoniae* is a leading cause of hospital- and community-acquired infections (including urinary tract infections, pneumonia, bacteraemia and soft tissue infections), primarily afflicting the young and immunocompromised, despite being part of the normal human intestinal microbiota and able to colonize the skin and nasopharynx of healthy individuals (Podschun & Ullmann 1998; Broberg et al. 2014). *K. pneumoniae* are naturally resistant to antibiotics such as amino-penicillins and carboxy-penicillins. Increasingly, treatment options are diminishing, leaving third-generation cephalosporins and carbapenems as the remaining alternative. However, the emergence of isolates carrying genes encoding extended spectrum beta lactamases (ESBL) and carbapenemases has raised alarm because it removes these last line treatment options and effectively negates the use of a whole class of antibiotics, with few alternatives. Therefore multiple agencies including the WHO, US Centers for Disease Control and Prevention and the UK Department of Health singled out *K. pneumoniae* as a global health concern (Boucher et al. 2009; Ahmad et al. 2012).

Historically *Klebsiella* isolates have been classified into serotypes and tracked using typing antisera. Serotyping is based on the recognition of distinct variations of surface-exposed polysaccharides, namely O-antigens and K-antigens, by specific antibodies, resulting in different O and K serotypes. O-antigens are the outermost part of the lipopolysaccharide (LPS), whereas K-antigens belong to the bacterial capsule (CPS). The number of serotypes has been estimated to be eight for O-antigens and 77 for K-antigens (Orskov, Fife-Asbury 1977; Trautmann et al. 1997; Edwards, Fife 1952; Edmunds 1954).

Multivalent protein-conjugate polysaccharide vaccines have been demonstrated to be highly successful and effective against bacterial pathogens, such as *Streptococcus pneumoniae* (Center, 2007). In *K. pneumoniae*, polyvalent vaccines based on the K-antigen have been developed and reached Phase I trials in humans (Edelman et al. 1994; Campbell et al. 1996). However, the high diversity of K-antigens and the confusing seroepidemiology render a vaccine with a broad coverage complex to develop and thus very costly. Compared with other *Enterobacteriaceae*, such as *Escherichia coli* [161 defined O serotypes (Iguchi et al. 2015a)] and *Shigella flexneri* [at least 47 O serotypes (Talukder et al. 2003)], *Klebsiella* has a surprisingly low number of reported O serotypes which promises a more viable alternative for vaccine development compared with K-antigen-based vaccines (Ahmad et al. 2012). Whole-genome sequence data allows us to verify and assess the diversity of these clusters and also estimate their frequency in contemporary isolates to assess the contribution of isolates bearing these markers.

The O-antigen biosynthesis enzymes are encoded on the *rfb* locus. To date, seven O-antigen clusters have been defined for *K. pneumoniae*, associated to serotypes O1, O2, O3, O4, O5, O8 and O12 (Table S1) and the *rfb* O-antigen biosynthetic pathway is well described (Raetz & Whitfield, 2002; Kalnych et al. 2014; Whitfield & Trent 2014). In essence it is an ABC-transporter-dependent pathway functionally composed of three types of enzymes: those responsible for (i) biosynthesis of nucleotide-activated sugars, (ii) polysaccharide repeat-unit synthesis and (iii) assembly of the repeat units and transport across the membrane (flippases).

Both the O1 and O2 antigen polysaccharide chains are based on a repeat-unit designated as D-galactan I. The difference being that the O1 antigen is capped by a distal D-galactan II unit, whereas O2 is not (Vinogradov et al. 2002; Kol et al. 1992). D-galactan II is the only known O-antigen polysaccharide in *Klebsiella* for which biosynthesis is enabled by genes (*wbb*Y and *wbb*Z) unlinked to the *rfb* cluster (Hsieh et al. 2014). The O1/O2 biosynthetic cluster occurs in two variants (Fig. 1a). Variant 1 consists of the transporter genes (*wzm* and *wzt*), three glycosyltransferases responsible for repeat-unit synthesis and an UDP-galactopyranose mutase, responsible for the synthesis of the polysaccharide subunit D-galactan I (Clarke & Whitfield 1992). Variant 2 is extended and carries three additional putative glycosyltransferases encoded by *gmlABC*. These additional genes are thought to explain the existence of a recently defined O2 subtype whereby the *gml* gene products modify D-galactan I, to give the chemically and antigenically distinct D-galactan III explaining the different subtype (Kelly et al. 1995; Szijártó et al. 2015). The serology of O1 isolates is proposed to be unchanged by GmlABC, because in this instance D-galactan II (encoded by WbbY and WbbZ) is the dominant antigenic epitope (Szijártó et al. 2015).

O-antigen polysaccharide structure and *rfb* sequence of *K. pneumoniae* serotypes O3 and O5 are identical to those of *E. coli* serotypes O9a and O8, respectively (Vinogradov et al. 2002; Iguchi et al. 2015b). Biosynthesis enzymes of O3 and O5 differ in the sequence of their mannosyltransferase (*WbdA*) and methyltransferase (*WbdD*). WbdD, in complex with WbdA, regulates the mannose chain length by capping the growing chain with a phosphate and methyl

Impact statement

Vaccines offer a potential remedy against the increasing threat of pan-drug-resistant *Klebsiella pneumoniae* strains. However the high diversity of surface antigens poses a challenge for vaccine design. This work is the first, to our knowledge, to catalogue naturally occurring polysaccharide antigen biosynthesis gene clusters in a globally representative collection of *K. pneumoniae* isolates, enabling us to identify novel serotypes and perform an epidemiological analysis. We show that only three O-antigen serotypes account for the majority of infections, offering a promising target for vaccine design.
group in O3 and a methyl group only in O5 (Hagelueken et al. 2012, 2015).

For completeness the O8 polymer is identical to the O1 antigen, with the exception of the partial O-acetylation of the d-galactan I repeat unit in O8 (Kelly et al. 1993) and serotype O9 may be considered as a subtype of serogroup O2 (Hansen et al. 1999; Trautmann et al. 1997; Kelly & Whitfield 1996). Serotype O9 Wzm and Wzt sequences share >95% identity to the O8 Wzm and Wzt orthologues (Fang et al. 2015) whilst most members of the Enterobacteiraeae have one conserved LPS core, two types of outer cores have been reported for K. pneumoniae; they differ in a set of three genes within the waa operon (Regué et al. 2005), which is independently located in the genome.

The seroepidemiology of K. pneumoniae has previously shown that for human-host-associated isolates the most prevalent O serotypes are O1, O2 and O3, with O1 being dominant in human disease (Trautmann et al. 1997;
Hansen et al. 1999; Yu et al. 2007; Fang et al. 2015). The observed prevalence of O4, O5 and O12 differs among the studies, whereas the significance of serotypes O8 and O11 remain equivocal (Trautmann et al. 1997). For O11 this is due to the lack of available rfb sequences and publicly available O11 isolates, so they could not be considered in this study. In addition, little is known about the association between O serotype and disease presentation, the only exception being the observation that serotype O3 isolates are more commonly isolated from blood samples compared with urine samples (Trautmann et al. 1997; Hansen et al. 1999).

The K-antigen biosynthesis enzymes are encoded on the cps (capsule polysaccharide synthesis) locus. To date the cps gene clusters of the 77 serologically defined K-types and nine additional distinct cps operons have been identified and published (Pan et al. 2015; Chung The et al. 2015; Wyres et al. 2015). The biosynthesis pathway is a Wzy-dependent polymerization pathway, identical to E. coli Group 1 capsule synthesis (Whitfield 2006; Larue et al. 2009; Shu et al. 2009; Bushell et al. 2013).

There is some confusion in the literature regarding the prevalence of K serotypes and their association with disease outcome (Cryz et al. 1986; Fung et al. 2000, 2002; Yu et al. 2007). This may be explained by the higher diversity of the K serotypes and possible cross-reactivity, making them much harder to define. In some studies K1 and K2 showed the highest prevalence and were associated with poorer disease outcome (Fung et al. 2000, 2002; Yu et al. 2007). In addition, these K serotypes have been associated with the emergence of hypervirulent K. pneumoniae causing community-acquired invasive infections such as pyogenic liver abscess, which has become of particular concern in parts of Asia (Shon et al. 2013). Hypervirulent K1 strains generally belong to the ST23 lineage, while K2 is found in more diverse backgrounds (Struve et al. 2015; Bialek-Davenet et al. 2014). However, in many studies outside of Asia, including an extensive seroepidemiological study of European and North American cases, K1 isolates are rare (Cryz et al. 1986).

Until now our knowledge of the nature and diversity of O- and K-operative and, by proxy, the antigens they encode is based on sequences of a limited number of reference isolates. Recently, the whole-genome sequences of more than 500 isolates collected from environmental samples, plants, mammals and non-human primates, as well as those from asymptomatic human carriage, from cases of invasive disease and from both the clinical and community setting were sequenced and published (Holt et al. 2015; Ellington, 2016; Chung The et al. 2015; Wand et al. 2015) (Table 1). These studies highlighted that the ability to cause invasive infections is not determined by lineage but is associated with the presence of virulence factors such as siderophore systems and the mmpA mucosity factor, although these virulence determinants are overrepresented in lineages associated with the hypervirulence phenotype (Holt et al. 2015; Bialek-Davenet et al. 2014; Struve et al. 2015).

Using this comprehensive whole genome dataset we set out to describe the genetic diversity of O-antigen and K-antigen and LPS core biosynthesis gene clusters and classify them by molecular serotyping. Using these data we then set out to determine if there was an association between O-, K- or LPS core types and disease outcome, phylogenetic lineage or the presence of other known virulence-related genes.

Methods

Bacterial isolates. Publicly available genome data derived from four different K. pneumoniae collections were analyzed in this study, totaling 573 sequenced isolates (Table 1). The global dataset consists of a globally representative collection from six different countries including isolates from different hosts and different infection types (Holt et al. 2015). The UK hospital dataset consists of a collection from the Cambridge University Hospitals NHS Foundation Trust and contains invasive isolates biased towards those with resistance to third-generation cephalosporins collected over a period of seven years (Ellington 2016). The Nepal hospital dataset contains human isolates from a single Nepalese hospital outbreak in 2012 (Chung The et al. 2015). The preanti-biotic dataset contains strains isolated before the widespread use of antibiotics (pre 1949) (Wand et al. 2015).

Genome data, generated in the above studies by paired end Illumina sequencing, were sourced from the European Nucleotide Archive (accession numbers are listed in Supplementary dataset S1, available in the online supplementary material). Reads were de novo assembled using Velvet (Zerbino & Birney 2008) and Velvet Optimiser, and the resulting assemblies annotated using Prokka (Seemann 2014), as described previously (Holt et al. 2015).

In silico serotyping. O serotyping was performed on the basis of the polysaccharide ABC transporters (flipases; Wzm and Wzt). The specificity of transporters to polysaccharide types has been noted previously (Cuthbertson et al. 2007). The protein sequences of Wzm were located and extracted from the assembled and annotated contigs using tblastn (Camacho et al. 2009) and grouped based on sequence clustering using CD-HIT (Fu et al. 2012) with an identity threshold of 95 %. This threshold was empirically derived and shown to be able to differentiate between the previously described rfb clusters (Table S1. The resulting groups were assigned the O serotype of known Wzm sequences (Table S1), which were included in the clustering step. In cases where Wzm was missing, Wzt homology was used for assignment. Any rfb loci which differed from previously described loci in their sequence and gene content were assigned to putative novel O serotypes. Novel rfb loci sequences were given the designation OL,[n], etc, (OL=LPS locus, to differentiate those defined on the basis of rfb locus sequences from those defined serologically; n is a numeric identifier beginning from n=100) (Fig. S1, Table S2).
differentiation of O1 from O2 serotypes was based on the presence of WbbY and WbbZ in O1 (NCBI accession number KJ451390, strain NTUH-K2044 (Hsieh et al. 2014)). LPS core types were assigned as described above, using WaaL clustering with an identity threshold of 80 %. The waa gene cluster was extracted by locating WaaC and WabG using tblastn. Several operons contained a contig break or stretches of low sequencing coverage; when no waaL could be identified or did not fit into any of the two main groups, these were marked as unassigned.

Due to the much higher diversity of the K antigen, grouping was performed based on the full-length cps sequence, which was extracted by locating wzi and wzc using tblastn (Camacho et al. 2009) and extending the presumed locus until a gene on the opposite strand appeared. The extracted cps locus DNA sequences were grouped using UCLUST (Edgar 2010) based on a 95 % identity. The groups were assigned to a serotype if they matched to any of the 86 previously described cps locus sequences (Pan et al. 2015; Chung The et al. 2015; Wyres et al. 2015).

Additionally, a database of wzi (Brisse et al. 2013) and wzc (Pan et al. 2013) sequences with known serotypes was used to verify the classifications. Any cps clusters which differed from previously described loci in their sequence and gene content were assigned to putative novel K serotypes. Putative full-length novel clusters were confirmed by generation of alternative assemblies using SPAdes 3.6.1 (Bankevich et al. 2012) and inspection of the resulting assembly graph using Bandage (Wick et al. 2015). Sequences were confirmed as full length if they spanned from the 5’ galF to the 3’ ugd gene. Full-length clusters were given the designation KL[n], etc: KL= capsule locus, to differentiate those defined on the basis of cps locus sequences from those defined serologically; n is a numeric identifier beginning from n=101 (although the first two loci (KL101 and KL102) have already been named KN1 and KN2 in the literature, and thus we will continue to use these designations here). Representative nucleotide sequences of full length cps and rfb clusters were annotated using Prokka (Seemann 2014) followed by manual curation, and deposited in GenBank (Fig. S1, Table S2). Novel wzi alleles were added to the K. pneumoniae whole-genome sequence typing database BIGSdb at the Institut Pasteur (http://bigdadb.web.pasteur.fr/) (Bialek-Davenet et al. 2014).

**Seroepidemiology of human K. pneumoniae infections.**

To analyze the prevalence of different O and K serotypes in human disease and their putative association with disease outcome, the 216 human-associated isolates of the global collection were used to investigate their seroepidemiology.

Three different infection properties were examined for association between O and K serotypes and LPS core variant: (i) infection site: we focused on isolates sampled from human blood (representing bacteraemia), sputum (pneumonia) and urine (urinal tract infections, UTI), isolates from other sites were excluded from this category, (ii) acquisition type: hospital-acquired isolates had been sampled in patients from 48 h after admission to hospital, community-acquired isolates had been sampled within 48 h of admission to hospital, isolates where no acquisition type was recorded were excluded from this category, (iii) infection status: carriage isolates are intestinal samples without any infection, non-invasive isolates are infections without bacteraemia (such as wound infection, pneumonia, UTI) and invasive isolates are from infections of normally sterile sites (such as blood).

<table>
<thead>
<tr>
<th>Dataset</th>
<th>Note</th>
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<td>Global</td>
<td>289 isolates; Human and environmental isolates, from six countries (Australia, Indonesia, Laos, Singapore, Vietnam, USA), sampled to maximize diversity and exclude members of a clonal outbreak, metadata includes invasiveness, acquisition type and sample site. Invasiveness status of human isolates consists of three types: carriage (isolates not considered to be the cause of an infection), non-invasive (pneumonia, urinary tract infection, wound infection; with no recorded bacteraemia) and invasive (isolated from normally sterile sites: blood, CSF, intra-ocular, pleural, pericardial, joint fluids, deep-seated tissue abscesses). Acquisition type is either community acquisition (isolated within 48 hours of admission to hospital) or hospital acquisition (isolated after 48 hours after admission)</td>
<td>(Holt et al. 2015)</td>
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<tr>
<td>UK Hospital</td>
<td>162 isolates; Collection from Cambridge University Hospitals NHS Foundation Trust in the UK over a period of seven years, invasive isolates isolated from normally sterile sites. Biased selection for antimicrobial resistance to three or more of six antimicrobial classes (penicillins, amoxicillin-clavulanate, aminoglycosides, fluoroquinolones, trimethoprime and third-generation cephalosporins), metadata includes sample site</td>
<td>(Ellington 2016)</td>
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<tr>
<td>Nepal Hospital</td>
<td>88 isolates; Nepalese hospital outbreak from May to December 2012, consisting mainly of two clonal lineages; randomly selected blood cultures.</td>
<td>(Chung The et al. 2015)</td>
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<td>Preantibiotic</td>
<td>34 isolates; Collection of isolates isolated before the widespread use of antibiotics (pre 1949). No additional metadata available.</td>
<td>(Wand et al. 2015)</td>
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Gene presence. A database of putative virulence genes was compiled (Table S3) and the presence of these genes was characterized as described previously (Chung The et al. 2015).

Multi locus sequence typing (MLST). MLST sequence types for all isolates were determined directly from sequence reads using SRST2 (Inouye et al. 2014) to type against the seven-locus MLST scheme (Diancourt et al. 2005). Sequence types (ST) are listed in Supplementary dataset S1.

Comparative gene analysis and phylogenetic trees. Comparative gene analysis and core genome definition was performed using Roary (Page et al. 2015), using a blastp percentage identity of 95 % and a core definition of 99 %. Phylogenies were inferred from either a SNP alignment generated by mapping reads to the K. pneumoniae MGH78578 reference sequence (Fig. S2) or from a concatenated alignment of core genes extracted from Roary output (Page et al. 2015) (Fig. 2). Single-gene alignments (Fig. 1b and Fig. S3) were performed using Clustal Omega (Sievers et al. 2011). Phylogenies were inferred from these alignments by running RAxML using a gamma distribution to model site-specific rate variation and 100 bootstrap replicates (Stamatakis 2006).

Results

Genetic structures of the rfb gene cluster

The rfb gene cluster was located from the genome sequence data and classified on the basis of Wzm and Wzt protein sequence. The rfb cluster was universally located between the cps and his gene clusters in all of our samples (Fig. S1). It was evident from these data that there were rfb operons that did not match any of the known rfb gene clusters (i.e. they did not belong to O1, O2, O3, O4, O5, O8 or O12), either as a result of different gene content or differing nucleotide sequence. These clusters were defined as novel geno-/serotypes. Out of the 573 isolates analyzed in this study, 533 isolates could be assigned to six known O serotypes, 36 isolates to five novel serotypes, and four isolates remain unassigned due to contig breaks or missing sequencing coverage (Supplementary dataset S1).

The clusters themselves range in size from seven to ten genes (8–10 kb). Both O1/O2 variants, O3, O4, O5 and O12, and four novel rfb clusters (OL101, OL102, OL103 and OL104) were observed in our dataset (Fig. 1a). The novel cluster, OL104, is identical in gene repertoire to O3 and O5, both synthesizing a mannose polymer repeated subunit. OL104 possesses the same WbdA as O3, that contains two mannosyltransferase domains and is distinct from the longer O5 WbdA, that contains three mannosyltransferase domains, which have been shown to have functional consequences in the mannosyl linkages they are able to synthesize (Greenfield et al. 2012). Within the isolates designated as serotype O3, two types of WbdD, the chain-length regulator, were observed. One (present in 34 of a total of 86 O3 isolates screened; 40 %) is identical to the E. coli O9a WbdD (Uniprot accession number Q47592), whereas the other type (present in the remaining 52 isolates; 60 %) has a deletion of 25 aa at the C-terminal region of its protein kinase domain. The WbdD of the isolates designated as serotype OL104 shows the same deletion (Fig. 1b). We speculate that OL104 and the two O3 WbdD variants might actually represent serologically distinct subtypes of the O3 serogroup.

The K. pneumoniae O12 antigen is composed of an N-acetylmuramic acid and rhamnose polymer (Vinogradov et al. 2002). Based on their gene content and order, O12, OL101 and OL102 are also closely related rfb clusters differing by the presence and absence of single genes: Compared with the O12 cluster, OL101 includes an additional glycosylhydrolase located on the opposite strand, while OL102 lacks the terminal glycosyltransferase gene but possesses an additional rhamnan synthesis gene (Fig. 1a).

It is of note that the direction of transcription in all of the rfb operons run antiparallel to those genes belonging to the adjoining his operon, with exception noted above, the O4 cluster and the extended variant of the O1/O2 O-antigen cluster (O1/O2 Variant 2) (Fig. 1a). The O4 rfb cluster is the only rfb cluster in K. pneumoniae containing a transposase. The O4 antigen is based on a galactose and ribofuranose polymer (Vinogradov et al. 2002).

When assessing the genetic diversity of the waa LPS core biosynthesis operon we detected both of the known core types, but found no conclusive evidence for novel core types.

Genetic structures of the cps gene cluster

The cps gene cluster showed a strikingly higher level of diversity both in sequence and gene content, compared to the rfb clusters (Fig. S1). Clusters ranged in size from 20.5 to 21.6 kb.

Of the 573 isolates analyzed, 387 (68 %) were assigned to 52 known K serotypes by comparison to previously described cps clusters representing the serologically typed reference strains (Pan et al. 2015) (Supplementary dataset S1). Three isolates (0.5 %) where only a partial sequence was available were assigned to a serotype with lower confidence due to only a partial match to the reference sequence. A total of 165 isolates (29 %) for which the cps sequence was > 5 % different to those of any of the known cps clusters were putatively assigned to 68 novel serotypes. Eighteen isolates (3 %) remained unassigned because no wzi and wzc sequences could be identified in the genomes. Among the putative novel serotypes, 21 distinct full-length cps sequences were identified (Fig. S1). Five of these sequences matched or were transposase variants of those previously described in (Wyres et al. 2015) and (Chung The et al. 2015), including one which was a transposase-negative
Seroepidemiology of *Klebsiella pneumoniae*

Fig. 2. Phylogeny of a randomly selected subset of isolates. Lineages are labeled according to the scheme of (Holt et al. 2015), subdividing the *K. pneumoniae* species into subspecies Kpl (K. *pneumoniae*), Kpll (K. *quasipneumoniae*) and Kplll (K. *variicola*). O and K serotypes and LPS core types, as well as the geographical origin, are denoted by colored squares as indicated.
variant of the cps cluster from *K. pneumoniae* HS11286 (genome accession NC_016845.1).

Consistent with results from previous studies, the first gene in all cps clusters was *galF* (Ebrecht et al. 2015) followed by a putative glucose phosphatase (*cpsACP*), the JUMPstart sequence (‘just upstream of many polysaccharide starts’) followed by the translocation and surface assembly genes (*wzi, wza, wzb* and *wzc*) (Rahn et al., 1999). Although the genes responsible for transport into the periplasm and polymerization (*wzx* and *wzy*), UndPP-Linkage (*wbaP* or *wcaJ*) and a 6-phosphogluconate dehydrogenase (*gnd*, a housekeeping gene not required for CPS synthesis (Chen et al. 2010)) were universally present, they were not found at fixed locations in these clusters. The cps clusters are usually terminated by UDP-glucose 6-dehydrogenase gene (*ugd*). The gene encoding glucose dehydratase (*rfG*) separates the cps and rfb clusters.

### The distribution of the rfb and cps gene clusters across *K. pneumoniae* phylogeny

To understand the distribution of the *rfb*, *cps* and *waa* gene clusters across the *K. pneumoniae* species we reconstructed a whole-genome-based phylogeny for all isolates included in this study and show their distribution (Fig. 2). This highlights that, with some exceptions, *rfb* clusters are not restricted to any particular clade. In our dataset the O1 and O2 rfb loci are only found in the KpI subspecies (Rahn et al. 2015)), for the other *rfb* clusters there is extensive evidence of horizontal gene transfer of the different *rfb* clusters between closely and more distant related lineages. This is further confirmed by the non-concordance of the phylogeny of the *rfb* gene clusters and the whole-genome phylogeny [O1 and O2: Fig. S2(a); O3, O5 and Novel 4: Fig. S2(b)].

The two *waa* LPS core types are equally not restricted to particular clades in the phylogeny, although we note a biased distribution with respect to the O serotypes. Out of the 573 isolates in the full dataset, 70 isolates (12 %) encoded core type 2; these core 2 strains are almost exclusively (65 isolates, 93 %) associated with serotype O1. The remaining isolates are core 1 (446, 78 %) or unassigned (57, 10 %). One previous study (Reguè et al. 2005) investigated the distribution of LPS core types in 100 *K. pneumoniae* strains based on PCR and dot blots and found a ratio of LPS core 1 to LPS core 2 isolates similar to our data (79 : 19), and also noted the association of LPS core 2 with serotype O1 (10 out of 34 O1 strains); however we did not observe an association with K serotype K2 as has been previously reported. Our analysis shows an elevated number of strains carrying both the LPS core type 2 and K2 or K45, however this is likely to be strongly influenced by two clonal expansions within the sample collection we analysed (Fig. 2). Furthermore, we could only observe LPS core type 2 in subspecies KpI (Fig. 2).

Due to the high diversity, K serotype switching is more difficult to analyze. As an example, K1 isolates belong mainly to two lineages, one of which (ST23) is associated with the hypervirulent phenotype, whereas the K2 serotype is much more distributed across the phylogenetic tree, including lineages associated with the hypervirulent phenotype (e.g. ST25) and other non-hypervirulent lineages (e.g. ST14, Fig. 2). This is consistent with previous reports which showed that many *K. pneumoniae* sequence types include multiple CPS types (Wyres et al. 2015; Holt et al. 2015; Brisse et al. 2009).

### O and K Serotype epidemiology

Looking across the full dataset of the 573 isolates the majority were genotyped as O1 (296 isolates, 52 %), followed by O2 (91, 16 %), O3 (86, 15 %), O5 (33, 6 %), OL101 (26, 5 %), O4 (18, 3 %) and O12 (9, 2 %). Each of the OL102 to OL104 serotypes occur in less than 1 % of isolates. The O1/O2 extended rfb cluster variant 2 was observed in 112 (38 %) of O1 genotyped isolates and 52 (57 %) O2 isolates (Supplementary dataset S1).

The seven most common O serotypes described above were associated with 54, 39, 40, 22, 14, 7 and 9 distinct K serotypes respectively, suggesting there is a relationship between the number of representatives of each O serotype in our dataset and the number of distinct K serotypes with which each is associated. This relationship is best explored using only the global isolate set, which is not biased towards any particular *K. pneumoniae* lineage or isolate source. Within this dataset O1 is the most common O serotype (n=133) and was associated with 43 distinct K serotypes. O3 is the next most common O serotype (n=45), followed by O2 (n=39) and O5 (n=24), these were associated with 29, 30 and 20 distinct K serotypes, respectively. Among the same isolate set the most common K serotype was K2 (n=20), associated with two distinct O serotypes (O1 and O2). K1 (n=15) and K64 (n=14) are the next most common serotypes and are associated with two and four O serotypes respectively. All other K serotypes are represented by eight or fewer isolates in our collection and were each associated with just one or two distinct O serotypes. These data indicate that re-assignment of K and O types occurs frequently in *K. pneumoniae*.

To investigate the seroepidemiology in human disease, we analyzed the association of O and K serotypes with the three different recorded infection properties (infection site, acquisition type or infection status). Significant correlations were found between K2 and invasive infections (odds ratio 4.1, p=0.008), and between OL101 O type and asymptomatic carriage (OR 8.5, p=0.009) (Table 2). Of special interest is also K1, which was virtually restricted to community-acquired isolates (OR 8.4, p=0.016) and completely missing in carriage-associated isolates (p=0.024). No correlation was found for any
Association between serotype and virulence gene content

Multiple virulence factors have been defined for *Klebsiella* (Lawlor et al. 2007; Broberg et al. 2014) including three siderophores (yersiniabactin, salmochelin and aerobactin), the cytotoxin colibactin and *rmpA/rmpA2*, a regulator of CPS overexpression (Cheng et al. 2010). Fimbrial adhesins are also known to play a major role in biofilm formation and are thus also classified as virulence factors. We determined if the distribution of these known virulence functions was correlated with O- or K-serotype using the existing global data set. The most notable association was the significant enrichment of an array of siderophore and colibactin clusters and the mucosity regulator *rmpA* with K serotypes K1 and K2 (Fig. 4a). Since K1 and K2 isolates were almost exclusively also O1 serotypes, the same effect was observed in O1. These results are perhaps unsurprising given that these virulence genes are overrepresented in the hypervirulent *K. pneumoniae* lineages, which are also associated with serotypes K1 and K2 (Holt et al. 2015; Bialek-Davenet et al. 2014).

Two types of fimbriae are known to be present in *K. pneumoniae* type 1 and type 3 fimbriae. Whilst type 3 fimbriae are known to bind to different human cell types, ranging from tracheal epithelia to renal tubular cells, as well as being important for binding to plants and abiotic surfaces (Khater et al. 2015), type 1 fimbriae appear more specific, so far they have been shown to be important for adhesion to uroepithelial cells and also for binding to plants (Klemm, Schembri 2000). Both type 1 and type 3 fimbriae are also associated with biofilm formation in *K. pneumoniae* type 3 fimbriae more strongly than type 1 (Schroll et al. 2010; Stahlhut et al. 2012; Klemm, Schembri 2000). Our data showed that serotype O1 isolates had a significantly higher likelihood of possessing type 1 fimbriae compared with any other O serotype (odds ratio 9.7, \( p=1.1 \times 10^{-16} \)), type 3 fimbriae (20 isolates without type 1 or type 3 fimbriae were almost exclusively also O1 serotypes, the same effect was observed in O1. These results are perhaps unsurprising given that these virulence genes are overrepresented in the hypervirulent *K. pneumoniae* lineages, which are also associated with serotypes K1 and K2 (Holt et al. 2015; Bialek-Davenet et al. 2014).

![Fig. 3. Distribution of serotypes in human *K. pneumoniae* isolates in the global collection. The eight most common O-types (left bars) and ten most common K-types (right bars) are shown. Other serotypes are shown in dark grey.](http://mgen.microbiologyresearch.org)
Table 2. Distribution of serotypes and LPS core types in human K. pneumoniae isolates in the global collection (n=216)

The eight most common O-types and ten most common K-types are shown. In the sample site category, samples from other than blood, urine or sputum were excluded. In the acquisition mode category, samples where no acquisition mode was reported were excluded.

<table>
<thead>
<tr>
<th>Sample site</th>
<th>Acquisition</th>
<th>Infectiousness</th>
</tr>
</thead>
<tbody>
<tr>
<td>O serotype</td>
<td>Blood</td>
<td>Urine</td>
</tr>
<tr>
<td></td>
<td>n=67</td>
<td>n=47</td>
</tr>
<tr>
<td>O1</td>
<td>32</td>
<td>21</td>
</tr>
<tr>
<td>O2</td>
<td>10</td>
<td>9</td>
</tr>
<tr>
<td>O3</td>
<td>12</td>
<td>9</td>
</tr>
<tr>
<td>O4</td>
<td>5</td>
<td>7.5%</td>
</tr>
<tr>
<td>O5</td>
<td>4</td>
<td>6.0%</td>
</tr>
<tr>
<td>O12</td>
<td>2</td>
<td>3.0%</td>
</tr>
<tr>
<td>OL101</td>
<td>1</td>
<td>1.5%</td>
</tr>
<tr>
<td>OL104</td>
<td>0</td>
<td>0.0%</td>
</tr>
<tr>
<td>Others†</td>
<td>1</td>
<td>1.5%</td>
</tr>
<tr>
<td>K serotype</td>
<td></td>
<td></td>
</tr>
<tr>
<td>K1</td>
<td>3</td>
<td>4.5%</td>
</tr>
<tr>
<td>K2</td>
<td>9</td>
<td>13.4%</td>
</tr>
<tr>
<td>K5</td>
<td>2</td>
<td>3.0%</td>
</tr>
<tr>
<td>K16</td>
<td>3</td>
<td>4.5%</td>
</tr>
<tr>
<td>K23</td>
<td>0</td>
<td>0.0%</td>
</tr>
<tr>
<td>K27</td>
<td>2</td>
<td>3.0%</td>
</tr>
<tr>
<td>K28</td>
<td>2</td>
<td>3.0%</td>
</tr>
<tr>
<td>K54</td>
<td>0</td>
<td>0.0%</td>
</tr>
<tr>
<td>K62</td>
<td>3</td>
<td>4.5%</td>
</tr>
<tr>
<td>K64</td>
<td>3</td>
<td>4.5%</td>
</tr>
<tr>
<td>Others†</td>
<td>40</td>
<td>59.7%</td>
</tr>
<tr>
<td>LPS core type</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Type 1</td>
<td>43</td>
<td>64.2%</td>
</tr>
<tr>
<td>Type 2</td>
<td>7</td>
<td>10.4%</td>
</tr>
<tr>
<td>Others†</td>
<td>17</td>
<td>25.4%</td>
</tr>
</tbody>
</table>

* Unidentified or not listed † Significant correlation (see text)

virtually absent in all other K serotypes (p=4.82×10−57, Fisher’s exact test). This K1-associated alginate lyase is located just upstream of the thiamine pyrophosphate (TPP) biosynthesis cluster (Fig. 4c). The only other commonly occurring alginate lyase gene in the dataset was present within the K14 cps cluster and has a sequence identity to the K1 associated alginate lyase below 45 %. Alginate lyases enable the use of alginate as carbon and energy source (Wong et al. 2000).

Discussion

The rfb and cps gene clusters give rise to the dominant serotypical properties of K. pneumoniae and are therefore priority vaccine candidates for the treatment of this increasingly multidrug-resistant pathogen. We have catalogued the naturally occurring diversity of these gene clusters within a large collection of isolates taken from different geographic and clinical settings, hosts and disease manifestations. In doing so we identified six known and five novel O-antigen clusters and 45 known and 18 novel complete K-antigen biosynthesis gene clusters. Of particular interest is a previously undescribed rfb cluster (OL101), occurring in 5 % of isolates in our dataset, originating in Europe, Asia and North America and found in human, bovine and environmental samples. We also identified a further 46 putative cps clusters (for which only partial sequences could be obtained) which appear novel. Although this may be an overestimate as some of these may represent transposase variants or divergent forms of other cps types or result from assembly errors; additional work will be required to investigate and validate these further.

This genetic catalogue of rfb and cps clusters enabled in silico serotyping of whole genome sequences. We focused on the seroepidemiology by analyzing the human-associated isolates from the global dataset (Holt et al. 2015), where clinical parameters such as infection site, type and acquisition
notable in the K serotype analysis was the distribution of serotype K1 sequences: most K1 isolates in our study belong to one of two lineages, ST23 and ST82 and showed a strong association with community-acquired infections, belonging to one of two lineages, ST23 and ST82 and showed a strong association with community-acquired infections, consistent with previous reports and the known association of ST23 with the hypervirulent phenotype (Tsay et al. 2002; Yu et al. 2007). K1 isolates were found in six out of seven countries covered by our genome collection. However, despite the fact that K1 is regarded as one of the two most common serotypes it was only found once in 162 isolates collected over a period of seven years in a UK hospital (this study) and is absent in another study based on 703 isolates in 13 hospitals located in Western Europe and Northern America (Cryz et al. 1986). This contrasts with reports from Taiwan, China and South Africa where K1 appears to have been dominant for a considerable period of time (Fung et al. 2000, 2002; Yu et al. 2007; Peng et al. 1991; Luo 1990) and has been associated with comparatively higher prevalence of hypervirulent infections in these countries, mostly linked to the ST23 lineage (Bialek-Davenet et al. 2014).

When looking for genes closely associated with K1 we identified the presence of a gene predicted to encode an alginate lyase isozyme. Outside of K1 the only other alginate lyase detected is distantly related and part of the K14 cps cluster. Hence, this gene is almost exclusively found in isolates of the K1 serotype (including those of both major K1 lineages). Pseudomonas aeruginosa is known to produce alginate during biofilm biogenesis in chronic lung infections of cystic fibrosis patients. Alginate lyase enables cell detachment from the biofilm (Boyd & Chakrabarty 1995; Ramsey & Wozniak 2005). However it is not known whether K. pneumoniae produces alginate biofilms. Moreover, since it is unusual to find alginate-lyase-producers, like the K1 K. pneumoniae produces alginate biofilms. Moreover, since it is unusual to find alginate-lyase-producers, like the K1 K. pneumoniae, that do not use alginate as primary carbon source, it has been proposed that alginate lyase production could be related to coinfection of P. aeruginosa and K. pneumoniae in cystic fibrosis patients (Wong et al. 2000). Given the above it is suggested that the association with cystic fibrosis is important, but probably it is opportunistic, and its role in K. pneumoniae is much broader.

We found no association of LPS core type with the isolates’ infection sites, infection types and acquisition modes. This is in contrast to a study showing that LPS core type 2 contributes to the level of virulence in K. pneumoniae, although its mechanism is so far unknown (Regué et al. 2005).

Virulence genes (siderophores, colibactin, rpmA and fimbrae) are expected to be overrepresented in isolates from infections compared with those from carriage or environmental sources and are particularly overrepresented in
lineages associated with hypervirulent invasive disease (Holt et al. 2015), thus the significantly high presence in K. pneumoniae serotype O1 isolates was anticipated. However, their significantly lower abundance in isolates of serotype O2 or O3, which are both also prevalent in human infection, was more surprising.

The striking difference in the distribution of the two major fimbriae types found in K. pneumoniae is also of note. Type 3 fimbriae are essential for biofilm formation (Schroll et al. 2010), while type 1 fimbriae are important for adhesion to uroepithelial cells and are thus considered virulence factors for urinary tract infection (Stahlhut et al. 2012). The differential distribution of fimbriae in K. pneumoniae probably allows isolates to adhere to different receptors and perhaps exploit or specialize in different niches, the presence of type 1 fimbriae in K2 isolates and the notable absence of type 1 fimbriae in K1 isolates is intriguing but will require more targeted sampling to unravel its true biological significance.

It is suspected that certain CPS types are able to influence the accessibility of the LPS O-antigen, possibly masking it; the reports are however inconclusive: Hsieh et al. (2012) described the masking of the O1 antigen by K1 capsule but not by K2, whereas Szijártó et al. (2015) showed that the O1 antigen is accessible by antibodies irrespective of the capsule type. The potential masking of O-antigen by the capsule warrants further research and is the crucial next step towards a polysaccharide-based vaccine against K. pneumoniae infection.

Our data indicate that O and K serotypes are frequently reassorted within the K. pneumoniae population. There are far fewer distinct O serotypes than K serotypes but among the O serotypes, those that are most common are generally associated with a higher number of distinct K serotypes. While there are lineage-specific clustering effects (Fig. 2), these data support the hypothesis that the rfb and/or cps clusters are shuffled within the K. pneumoniae population via horizontal gene transfer. However given the wide diversity of K serotypes, much larger strain collections will need to be examined in order to detect a statistically significant divergence from random re-arrangement of CPS and LPS loci.

K. pneumoniae is considered to be a significant threat to human health with the rates of infection increasing globally and appearing to be driven by increasing levels of antimicrobial resistance to front line antimicrobials. The short-term solution has been to turn to old drugs such as colistin that are associated with significant nephrotoxicity. However, the development of resistance to colistin is rapid and explained by both intrinsic mechanisms, such as point mutations (Cheng et al. 2015), as well as by the acquisition of genes by lateral gene transfer (Liu et al. 2015). With this in mind we found that O-serotype prevalence and distribution were stable with regards to different infection types and sites. In addition although O serotype switching has been a common occurrence in the evolution of K. pneumoniae K serotypes show a much bigger variance (Wyres et al. 2015). Consequently O antigens, defined in detail in this study, offer a promising target for vaccine design that warrant further research.

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References


Seroepidemiology of *Klebsiella pneumoniae*


**Data Bibliography**


