Comment on: CG258 Klebsiella pneumoniae isolates without β-lactam resistance at the onset of the carbapenem-resistant Enterobacteriaceae epidemic in New York City

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Sir,

We read with interest the recent article by Eilertson et al.,1 ‘CG258 Klebsiella pneumoniae isolates without β-lactam resistance at the onset of the carbapenem-resistant Enterobacteriaceae epidemic in New York City’, wherein the authors conduct an investigation of K. pneumoniae causing bloodstream infection in New York City in 1999, 2003–04, 2006, 2009 and 2013. However, we felt that readers would benefit from further discussion of the results in the context of what is known about clonal group (CG) 258, in particular from recent genomic analyses.

In the article by Eilertson et al.,1 CG258 was defined in the discussion section as ‘ST258 and its single allele variants’, referring to alleles of the seven-locus MLST scheme.2 Notably, this definition includes ST11, which: (i) has been shown to be the ancestor of ST258; (ii) differs from ST258 by a single allele (wzi154) that has been shown to be the ancestor of ST11; (iii) includes ST442, which has been shown to be the ancestor of ST258.3 Notably, in the article by Eilertson et al.1, the PCR used to screen for CG258 isolates targets the tonB-79 allele of ST258, which is not present in ST11. Hence, the screen detects only the tonB-79 subgroup of CG258 and is specifically unable to detect ST11 isolates, which may be the most informative members of CG258 in terms of their potential to reveal details of the early emergence of ST258.3–8

Eilertson et al.1 report that the earliest ST258 isolates they identified (2004) carried wzi154, while some later isolates carried wzi29 and occasionally other wzi alleles. This is as expected given the previously reported genomic data, which show that following the initial formation of ST258-wzi154 (KL107), a subsequent ~50 kbp recombination event occurred with an ST42 K. pneumoniae3,4,6–8 importing a new capsule locus harbouring wzi29 to form the ST258-wzi29 subclade (referred to as cps-1 in Eilertson et al.,1 designated KL106 under the standardized nomenclature9). The genomic comparisons therefore support a line of descent from ST11 to ST258-wzi154 to ST258-wzi29 (summarized in Figure 1). Molecular dating analyses estimate that ST258 emerged from ST11 in the mid-1990s3,4 and the ST258-wzi29 subclade emerged ~7–8 years later in the early 2000s.5

The detection of ST258-wzi154 (KL107) in 2003, with ST258-wzi29 (KL106) detected later, is therefore consistent with the prior date on the stepwise evolution of ST258. However, contrary to the statements in the manuscript of Eilertson et al.,1 these data do not suggest that KL107 (wzi154) was the initial capsule type of CG258, either in New York City or globally, as the study captured only ST258 and its direct derivatives that form just one subgroup of CG258 (as discussed above). The ancestral capsule type of the entire CG, or of the ST11 progenitor strain from which ST258 emerged through recombination, remains unknown because CG258 harbours extensive capsule locus diversity (e.g. see Figure 1).3,4,9,10

Transparency declarations
None to declare.

References
Figure 1. Evolutionary history of CG258. Phylogeny and K (capsule) locus data are reproduced from Lam et al.\textsuperscript{10} (outgroup-rooted and recombination-filtered maximum likelihood phylogeny inferred from core genome nucleotide variants). Clades corresponding to each chromosomal ST plus the ST258-wzi29 subclade are highlighted and labelled, followed by the seven-gene MLST designation. K locus and wzi allele numbers are shown by coloured blocks according to the key. Red circles mark the hypothetical most recent common ancestors for which dates have been estimated in published Bayesian analyses: (i) the whole CG;\textsuperscript{5} (ii) all ST258;\textsuperscript{4,5} and (iii) ST258-wzi29 subclade.\textsuperscript{6}
Sir,
We thank Wyres et al.1 for their thoughtful comments on our recent study.2 Subsequent MLST data from a broader set of 195 Klebsiella pneumoniae isolates from the same collection (including all isolates in our recent study) showed no carbapenem-resistant or blakpc-harbouring isolates of the ST11 lineage. Only one ST11 isolate was identified. This isolate dated from 2013 and was fully susceptible to all β-lactam antibiotics tested. Wyres et al.1 correctly state that the PCR assay used to define CG258 only targets the tonB-79 subgroup; however, this oversight is somewhat mitigated by the lack of ST11 isolates from our hospitals over the period studied. Thus, the findings of our recent study2 are representative of regions such as Brooklyn, New York, where ST11 isolates are rare. Based on the noted lack of ST11 isolates we cannot comment on the ancestral capsule type in those isolates. Further study on historical ST11 K. pneumoniae isolates is needed to fill this knowledge gap.

Transparency declarations
None to declare.

References