

## Comparison of childhood pneumococcal carriage and serotype distribution between inpatients with pneumonia and healthy controls in Nepal prior to pneumococcal conjugate vaccine introduction

Rama Kandasamy<sup>1,2</sup>, Michael Carter<sup>1,2</sup>, Meeru Gurung<sup>3</sup>, Shrijana Shrestha<sup>3</sup>, Imran Ansari<sup>3</sup>, Madhav C. Gautam<sup>3</sup>, Susan Ndimah<sup>1,2</sup>, Stephen Thorson<sup>3</sup>, Katherine L. O'Brien<sup>4</sup>, Sarah Kelly<sup>1,2</sup>, David R. Murdoch<sup>4</sup>, Dominic F. Kelly<sup>1,2</sup>, and Andrew J. Pollard<sup>1,2</sup>

<sup>1</sup>Oxford Vaccine Group, Department of Paediatrics, University of Oxford, Oxford, United Kingdom. <sup>2</sup>NIHR Oxford Biomedical Research Centre, Oxford, United Kingdom. <sup>3</sup>Paediatric Research Unit, Patan Academy of Health Sciences, Kathmandu, Nepal. <sup>4</sup>Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, USA. <sup>5</sup>Department of Pathology, University of Otago, Christchurch, New Zealand.

## BACKGROUND

- Some pneumococcal serotypes are infrequently carried by healthy children but represent a large proportion of disease isolates.
- Detection of serotype-specific pneumococcal colonization in children with pneumonia, although not diagnostic of etiology at an individual level, likely enriches the group of cases for true pneumococcal pneumonia.
- We aimed to assess serotype-specific carriage among Nepalese children admitted to hospital with physician diagnosed pneumonia and compare it with that of healthy community controls prior to PCV10 introduction.

## METHODS

- Community children aged 6-24 months without pneumonia were recruited from the outpatient department of Patan Hospital, Kathmandu, Nepal.
- All children aged 2 months to 14 years admitted with clinician diagnosed pneumonia to Patan Hospital were approached for enrolment into the study and those aged 6-24 months included in this analysis.
- Nasopharyngeal swabs were collected from recruited children, processed according to WHO guidelines, and serotype determined by Quellung reaction (Serum Staten Institute, Denmark).

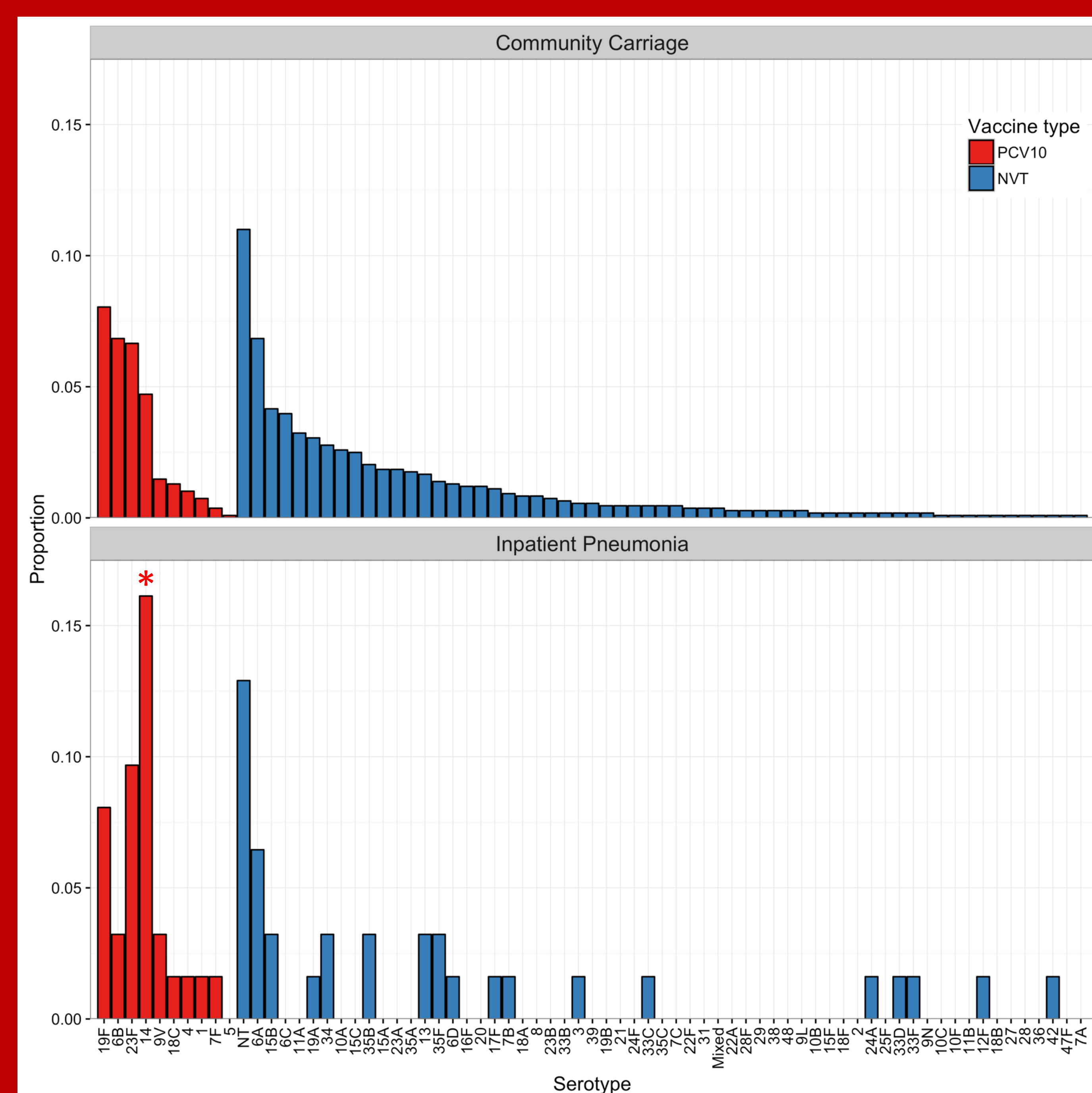
## RESULTS

- Between April 2014 and August 2015 we enrolled 1751 healthy children aged 6-24 months and 164 children with clinician diagnosed pneumonia aged 6-24 months.
- Carriage prevalence differed between the groups ( $p < 0.0001$ ):
  - Community children: 65% (95% CI 62.8-67.2; 1141/1751)
  - Hospitalized pneumonia children: 37.8% (95% CI 30.9-46.3; 62/164)
- In the pneumonia cohort 27% (95% CI 16.6-39.7; 17/63) of carriers had had antibiotics in the week before admission compared with 46.5% (95% CI 36.6-56.7; 47/101) of non-carriers ( $p = 0.0140$ )
- Pneumococcus colonized pneumonia cases were more likely to have a PCV10 strain than were colonized community children ( $p = 0.0166$ ).
  - Community children: 31.2% (338/1082; 95% CI 28.5-34.1%)
  - Hospitalized children with pneumonia: 46.8% (95% CI 34-59.9, 29/62)

## FUNDING

This study is funded by Gavi, The Vaccine Alliance.

## Serotype-specific pneumococcal carriage in children admitted with pneumonia compared with healthy community controls



**Figure 1. Serotype-specific pneumococcal carriage distribution in paediatric inpatients with pneumonia at Patan hospital, Kathmandu and healthy community controls prior to PCV10 introduction.** Children were aged 6-24 months from Kathmandu, Nepal and had not received a pneumococcal conjugate vaccine had a single nasopharyngeal swab collected and any isolated pneumococci serotyped by Quellung. NT = non-typeable. \*Significantly higher odds of being identified in inpatients compared with community controls ( $p = 0.0011$ )

## CONCLUSIONS

- Carriage prevalence in inpatients with pneumonia is significantly lower than community controls.
- Antibiotic usage prior to presentation is a significant modifier of carriage prevalence in inpatients with pneumonia.
- Assessment of pneumococcal carriage of paediatric inpatients with pneumonia may provide improved insight into PCV effect on serotypes infrequently isolated from healthy children.