Pharyngeal carriage of *Neisseria* species in the African meningitis belt

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**KEYWORDS**
Non-meningococcal *Neisseria*;

**Summary**
Objectives: *Neisseria meningitidis*, together with the non-pathogenic *Neisseria* species (NPNs), are members of the complex microbiota of the human pharynx. This paper investigates the influence of NPNs on the epidemiology of meningococcal infection.

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Pharyngeal carriage; African meningitis belt

Methods: Neisseria isolates were collected during 18 surveys conducted in six countries in the African meningitis belt between 2010 and 2012 and characterized at the rplF locus to determine species and at the variable region of the fetA antigen gene. Prevalence and risk factors for carriage were analyzed.

Results: A total of 4694 isolates of Neisseria were obtained from 46,034 pharyngeal swabs, a carriage prevalence of 10.2% (95% CI, 9.8–10.5). Five Neisseria species were identified, the most prevalent NPN being Neisseria lactamica. Six hundred and thirty-six combinations of rplF/fetA, VR alleles were identified, each defined as a Neisseria strain type. There was an inverse relationship between carriage of N. meningitidis and of NPNs by age group, gender and season, whereas carriage of both N. meningitidis and NPNs was negatively associated with a recent history of meningococcal vaccination.

Conclusion: Variations in the prevalence of NPNs by time, place and genetic type may contribute to the particular epidemiology of meningococcal disease in the African meningitis belt.

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Introduction

The human pharynx hosts a complex microbiota, including bacteria belonging to the genus Neisseria. Most members of this genus are non-pathogenic commensals (non-pathogenic Neisseria, NPNs), which very rarely cause invasive disease, but Neisseria meningitidis (Nm), the meningococcus, is an exception.1 Despite advances in vaccine development, invasive meningococcal disease remains a public health challenge globally and especially in the African meningitis belt, where very large epidemics continue to occur.2,3 despite the recent widespread deployment of a serogroup A meningococcal conjugate vaccine (PsA-TT, MenAfriVac®).4–8

Most pharyngeal carriage studies in the African meningitis belt have focused on the meningococcus,9–12 with little attention paid to other Neisseria species apart from Neisseria lactamica (Nl). In northern Nigeria, pharyngeal carriage of Nl was common, especially in young children13 and genetic exchange among Nm, Nl and unspecified Neisseria species has been demonstrated in The Gambia.14 Molecular epidemiological studies of Nm and Nl conducted in Burkina Faso from 2009 to 2012 reported a high overall prevalence of Nl carriage (18.2%), a higher prevalence of Nl in males than in females, except in those aged 18–29 years, no change between dry and rainy season and no significant changes following vaccination with PsA-TT.10,15–17 It has long been considered likely that carriage of NPNs influences host susceptibility to infection and invasion by Nm, a view supported by the fact that healthy subjects inoculated with Nl exhibit some protection against colonization with Nm.18 Until recently, Nl was considered to be the NPN genetically most similar to Nm. However, recent whole genome sequence (WGS) studies have shown that Neisseria polysaccharea (Np) and Neisseria bergeri (Nb) are more closely related to Nm than Nl.19 Consequently, these bacteria may also influence the epidemiology of Nm colonization and invasion. For this reason we have studied the prevalence of carriage with various Neisseria species in six countries of the African meningitis belt and investigated risk factors for their carriage.

Materials and methods

The methods employed in the MenAfriCar surveys, during which the isolates described in this paper were collected, have been described in detail previously11 and are summarized briefly here. Ethical approval for the surveys was obtained from the ethics committee of the London School of Hygiene & Tropical Medicine and from ethical committees in each partner country. The study was registered with ClinicalTrials.gov (NCT01119482).

Carriage surveys

Bacteria were isolated during 18 cross-sectional surveys conducted in Chad, Ethiopia, Ghana, Mali, Niger and Senegal during 2010–2012 (Fig. 1). Pre-vaccination surveys in Mali, Niger and Chad had a target of 5000 participants and post-vaccination surveys a target of 2000 participants in Mali and Niger and 6000 in Chad. The remaining surveys aimed to recruit 2000 participants. A representative sample of households was obtained either from an updated demographic surveillance system (DSS) or from a census conducted for the study. Within selected households, individuals in four different age groups (0–4 years, 5–14 years, 15–29 years and 30 years or more) were chosen randomly until the required sample size was reached, with a maximum of 5 individuals recruited per household. Once written consent had been obtained, standardized household and individual questionnaires inquiring about risk factors for meningococcal infection were administered. Pharyngeal swabs were obtained using a standardized technique that involved swabbing both the posterior-pharyngeal wall and the tonsils.20
Bacteriology

NPNs were isolated using the same conventional microbiology techniques that were employed for the detection of Nm described previously. Briefly, pharyngeal swabs were plated onto modified Thayer Martin agar plates and incubated 24–48 h at 37 °C in 5% CO₂; oxidase and gram stain testing identified oxidase positive, Gram-negative diplococci. Further biochemical tests (ortho-nitrophenyl-b-galactoside, -glutamyl transpeptidase and tributyrin) were used in each site to differentiate between putative Nl and Nm isolates and members of the genus Moraxella.

Molecular methods

A boiled cells suspension of each oxidase positive, Gram negative diplococci (OPGND) isolate was sent to the Department of Zoology at the University of Oxford for molecular typing, where Sanger sequencing was used to characterize gene targets as described in the Supplement. Sequences were assembled using SeqSphere (http://www.ridom.de/seqsphere/) and imported into the isolate’s record previously created in a BIGSdb database.

Neisseria speciation

Amplification and sequencing of a 413 bp fragment of the rplF gene was used to differentiate among Neisseria species as described previously. A phylogeny based on the f_rplF alleles from Bennett et al. (2014) and the unique alleles found in this study was reconstructed using the Neighbor-Joining algorithm and the Kimura 2-parameter substitution model in MEGA version 6.0. For isolates that did not yield results for the rplF assay, sequencing of the rrna gene, encoding 16S rRNA, was used to confirm the presence or absence of a bacterium and to determine its genus. Only rplF confirmed NPNs were included in this study with the exception of four Nm speciated on the basis of the 16S rRNA and the porA sequences. New alleles were investigated by a BLAST of obtained sequences against the 16S rRNA sequence data of the EzTaxon server (http://www.ezbiocloud.net/eztaxon).

Genetic diversity

Sequencing of the variable region of the fetA gene (feta_VR) was used to assess the genetic diversity of the Neisseria identified as described in the Supplementary methods. The sequences were assembled as for the other targets using SeqSphere (http://www.ridom.de/seqsphere/) and imported into the isolate’s record previously created in a BIGSdb database.

For isolates for which feta_VR could not be amplified, an assay identifying the absence of the fetA gene (fetA null, fnl) was employed, using primers placed on genes on each side of the fetA gene: thdF and fetB as described in the Supplementary methods.
Statistical methods

Analyzes were performed using Stata v12.0 (StataCorp, Texas). Survey design and potential household clustering were taken into account using the survey commands in Stata. Carriage prevalence of each of the different Neisseria species, together with 95% confidence intervals, was calculated for each country and each survey. Risk factors for carriage of Nm and NPNs were assessed simultaneously using multinomial logistic regression. Each risk factor was considered in turn using univariable, multinomial logistic regression. A multivariable model was then constructed including country, age group and sex a priori and any variable with a p-value < 0.1 in the univariable analyses; only the variables with a p value < 0.05 in the multivariable analysis were kept in the final model. As a final check, dropped variables were re-entered into the model one at a time and the p-values re-examined; the variable was retained as significant if the p value was < 0.05.

Results

Prevalence of carriage with Neisseria species

A total of 4694 of the 46034 pharyngeal swabs collected yielded a Neisseria species, giving a carriage prevalence of 10.2% [95% CI, 9.8–10.5%]: 696 Neisseria were identified out of the 946 OPGND samples received from Chad; 838 out of the 994 received from Ethiopia; 446 out of the 544 received from Ghana; 298 out of the 504 from Mali; 1644 out of 2321 from Niger and 773 out of the 971 received from Senegal. The most frequently isolated species was Nl with a 5.6% point prevalence [95% CI, 5.3–5.8%], followed by Nm at 3.6% [95% CI, 3.4–3.8%], Np at 0.6% [95% CI, 0.5–0.7%], Nb at 0.2% [95% CI, 0.2–0.3%] and Neisseria subflava (Ns) at 0.05% [95% CI, 0.03–0.1%] (Supplementary Table 1). Twenty isolates from Chad were identified as belonging to the Neisseria genus but did not cluster with any known species on the Neighbor-Joining Tree (NJT; data

Table 1  Factors associated with carriage of Neisseria meningitidis and non-pathogenic Neisseria; results from a multinomial multivariable logistic regression.

| Factor                  | Number (carriers) | Adjusted RRR Neisseria meningitidis (95% CI) | Adjusted RRR Non-pathogenic Neisseria (95% CI) |
|-------------------------|-------------------|---------------------------------------------|-----------------------------------------------|
| Age                     |                   |                                             |                                               |
| <1 year                 | 2074 (207)        | 0.40 (0.29, 0.55)                           | 3.11 (2.60, 3.73)                             |
| 1–4 years               | 8291 (1355)       | 0.70 (0.59, 0.82)                           | 5.90 (5.23, 6.65)                             |
| 5–14 years              | 12,563 (895)      | 1.49 (1.31, 1.69)                           | 2.59 (2.29, 2.94)                             |
| 15–29 years             | 11,863 (4372)     | 1.0                                         | 1.0                                           |
| 30+ years               | 11,243 (206)      | 0.59 (0.50, 0.68)                           | 0.51 (0.43, 0.61)                             |
| Sexa                    |                   |                                             |                                               |
| Female                  | 26,619 (1701)     | 1.0                                         | 1.0                                           |
| Male                    | 19,296 (1313)     | 1.34 (1.21, 1.48)                           | 0.87 (0.80, 0.94)                             |
| Country                 |                   |                                             |                                               |
| Chad                    | 13,396 (584)      | 1.0                                         | 1.0                                           |
| Ethiopia                | 5970 (450)        | 7.11 (5.43, 9.29)                           | 1.70 (1.44, 2.01)                             |
| Ghana                   | 5209 (253)        | 4.81 (3.64, 6.35)                           | 1.25 (1.03, 1.51)                             |
| Mali                    | 8837 (219)        | 1.45 (1.05, 1.99)                           | 0.48 (0.40, 0.58)                             |
| Niger                   | 8213 (1112)       | 11.39 (9.00, 14.43)                         | 3.58 (3.13, 4.01)                             |
| Senegal                 | 4409 (427)        | 10.85 (8.28, 14.23)                         | 2.39 (1.97, 2.89)                             |
| Area                    |                   |                                             |                                               |
| Urban                   | 19,462 (1398)     | 1.0                                         | 1.0                                           |
| Rural                   | 26,572 (1637)     | 1.44 (1.09, 1.60)                           | 0.97 (0.88, 1.06)                             |
| Crowdingb               |                   |                                             |                                               |
| <2 people per room      | 16,299 (889)      | 1.0                                         | 1.0                                           |
| ≥2 people per room      | 29,679 (2127)     | 1.27 (1.12, 1.45)                           | 1.08 (0.98, 1.19)                             |
| Kitchen location        |                   |                                             |                                               |
| Open air                | 17,805 (1274)     | 1.0                                         | 1.0                                           |
| Inside house            | 12,741 (1027)     | 1.32 (1.09, 1.60)                           | 0.90 (0.79, 1.02)                             |
| Separate hut           | 14927 (677)       | 0.94 (0.76, 1.17)                           | 0.96 (0.84, 1.09)                             |
| Missing information     | 504 (38)          | 1.06 (0.57, 1.98)                           | 0.77 (0.51, 1.17)                             |
| Vaccinated recently     |                   |                                             |                                               |
| with meningitis vaccine |                   |                                             |                                               |
| No                      | 31,338 (2060)     | 1.0                                         | 1.0                                           |
| Yes, <1 year ago        | 9048 (545)        | 0.71 (0.59, 0.84)                           | 0.82 (0.73, 0.92)                             |
| Yes, 1–3 years ago      | 4543 (356)        | 0.51 (0.42, 0.63)                           | 0.94 (0.81, 1.10)                             |
| Don’t know/missing      | 1020 (55)         | 0.68 (0.47, 1.00)                           | 0.62 (0.46, 0.84)                             |

The final multivariable logistic regression model included age group, sex, country, season area, crowding, kitchen location and recent vaccination.

Other risk factors used in the univariable model but not significant in the adjusted model: smoking, living in a house with smokers, cooking fuel, respiratory symptoms and attendance of social gatherings.

a Sex not reported for 119 individuals.
b Data not reported for 57 individuals.

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Factors influencing the prevalence of carried Neisseria species

Country, season, age, gender, and a history of recent vaccination against Nm were associated with carriage of NPNs. Additionally, area of residence, household crowding and kitchen location were significant risk factors for carriage of Nm (Table 1).

Country: The prevalence of carriage of Neisseria species overall varied significantly among countries with Niger having the highest point prevalence (19.9% [95% CI, 18.9–20.9]), followed by Senegal (17.5% [95% CI, 16.1–18.8]), Ethiopia (13.8% [95% CI, 12.8–14.8]), Ghana (8.5% [95% CI, 7.6–9.4]), Chad (5.3% [95% CI, 4.9–5.7]) and Mali (3.4% [95% CI, 2.9–3.8]) (Supplementary Table 1). The distribution of the different species of Neisseria by country is shown in Fig. 1. The prevalence of Nm carriage varied significantly among countries, being highest in Senegal (8.0% [95% CI, 7.1–9.0]) and lowest in Mali (0.9% [95% CI, 0.7–1.1]). There were also major differences in the prevalence of carriage of NPNs by country and some NPNs were not identified in some countries, for example no Nbf were isolated in Ethiopia and Senegal and no Ns in Ghana. Ethiopia was the only country where the prevalence of Nm carriage was higher than that of Ni.

Year and season: The prevalence of carriage of Neisseria species overall varied little over the three years of the study: 10.0% [95% CI, 9.4–10.5%] in the first survey, 11.1% [95% CI, 10.6–11.7%] in the second survey and 9.4% [95% CI, 8.9–9.9%] in the third survey (Table 2). The prevalence of carriage with both Nm and NPNs was also similar over time, with Ni being the most carried species, regardless of survey or season. There was, however, variation in the prevalence of carriage of NPNs over time at the country level: for example, there was an increase in the prevalence of Ni between surveys 1 and 2 in both Chad (from 1.1% to 5.9%) and Ghana (from 0.8% to 6.3%) and an increase in Nm prevalence between survey 2 and 3 in Senegal (from 3.1% to 19.8%) (Supplementary Fig. 1). The relative risk of carriage of NPNs was significantly higher during the dry compared to the rainy season with an adjusted Relative Risk Ratio (aRRR) of 0.78, [95% CI, 0.70–0.86], whereas the opposite was true for Nm with an aRRR of 1.53, [95% CI, 1.35–1.74] in the dry season.

Age and gender: The relative risk of carrying NPNs overall was higher in children aged less than 15 years compared to the young adult age group (age 15–29 years) with an aRRR of 3.11 [95% CI, 2.60–3.73] for those <1 year, 5.90 [95% CI, 5.23–6.65] for the 1–4 year olds and 2.59 [95% CI, 2.29–2.94] for the 5–14 year olds. The oldest age group >30 years had a lower aRRR of 0.51 (95% CI, 0.43–0.61). The prevalence of carried Ni was highest in the 1–4 year old age group, reaching a peak of 14.1% [95% CI, 13.3–14.8%] in contrast to carriage of Nm, which reached a peak of 5.2% [95% CI, 4.8–5.6%] in the 5–14 year age group. Similarly to Ni, Np carriage also reached a peak prevalence of 1.7% [95% CI, 1.4–1.9%] in the 1–4 year old age group (Fig. 2).
Carriage prevalence of \( \text{Nb} \) and \( \text{Ns} \) was too low to identify an overall trend in age group distribution. Males had a lower risk of carrying NPNs than females (aRRR of 0.87 [95% CI, 0.80–0.94]) but a higher risk of carrying \( \text{Nm} \) (aRRR of 1.34 [95% CI, 1.21–1.48]).

**Vaccination history:** A history of vaccination within the past 12 months with any meningococcal vaccine was associated with a decrease in \( \text{Nm} \) carriage, as reported previously,\(^8\) but additionally with a decreased risk of carrying NPNs (aRRR of 0.82 [95% CI, 0.73–0.92]). In the three countries where MenAfriVac\(^k\) was introduced during the course of the study (Chad, Mali, and Niger) a significant decrease in the prevalence of carriage of \( \text{Ni} \) from 6.4% to 4.9% was observed. An overall decrease in carriage of \( \text{Nm} \) was also observed in the three countries but this reduction was not consistent in all countries with Mali experiencing an increase from 0.6% to 1.2% (Supplementary Fig. 2).

**Other risk factors:** Area of residence (rural vs urban), crowding, cooking with cow dung or straw, kitchen location, and attendance at social gatherings in the past week all had a significant impact on the odds of carrying NPNs in the univariable regression model, but their effect was not significant in the multivariable model. Some of these risk factors were retained in the final model as, although they had no effect on NPN carriage, they were significant for \( \text{Nm} \) carriage.

### Genetic diversity of identified *Neisseria* species

Forty-two different alleles were identified for the \( rplf \) fragment (\( f_rplf \)). \( \text{Ni} \) was the most diverse species with 17 \( f_rplf \) alleles, the most frequent of which was \( f_rplf \) 6 (1453 isolates, 56.1%). Eleven alleles were found for \( \text{Nm} \) the most frequent being \( f_rplf \) 2 (749 isolates, 44.6%) and \( f_rplf \) 1 (707 isolates, 42.1%); four \( f_rplf \) alleles were identified in \( \text{Np} \), the most frequent of which was \( f_rplf \) 9 (207 isolates, 71.4%). Four alleles were also found among the \( \text{Nb} \) alleles with \( f_rplf \) 62 (57) and \( f_rplf \) 69 (46) representing 91.2% of these isolates. Finally, seven alleles were observed for \( \text{Ns} \), with predominance of \( f_rplf \) 43 (15 isolates, 62.5%) (Supplementary Table 2). A Neighbor Joining tree was used to represent the phylogeny of the \( rplf \) fragment present in the *Neisseria* isolates of this study in relation to the original isolates used to create the \( f_rplf \) assay\(^22\) (Supplementary Fig. 3).

The diversity of \( f\text{eta} \) alleles varied by country (Fig. 3A; Supplementary Table 3); Niger had the highest number of \( f\text{eta}_\text{VR} \) alleles (111) and Ghana had the least (55). Some allele variability was seen between surveys, for example the proportion of the \( \text{F1-1} \) allele increased from 61 (3.63%) in survey 2 to 342 (23.38%) in survey 3 and similar changes were observed for others alleles in survey 3 (Fig. 3B; Supplementary Table 4). A total of 234 different alleles were identified across all species; 75 of these were found in only one isolate; 184 in fewer than 20 isolates. A total of 194 alleles were observed for \( \text{Ni} \), 80 for \( \text{Nm} \); 35 for \( \text{Np} \); 21 for \( \text{Nb} \); and 16 for \( \text{Ns} \). Most alleles were found predominantly in only one species, for example 99.61% of \( \text{fnl} \) alleles were found in \( \text{Nm} \);\(^29,30\) all F5-84 variants were only detected in \( \text{Ni} \), all F5-1 were exclusive to \( \text{Nm} \), all F11-4 were found only in \( \text{Np} \) and all F1-169 and all F1-193 variants were only observed in \( \text{Ns} \). Some variants, however, were shared among different species, e.g. F1-72, F1-21, F2-24, F6-3 (Fig. 3C; Supplementary Table 5). Neither the \( f\text{eta}_\text{VR} \) nor the \( \text{fnl} \) fragment was successfully amplified in 643 isolates, which were designated Not Determined (ND). As defined by \( f_rplf \) and \( f\text{eta}_\text{VR} \) alleles, there were 636 different *Neisseria* strain types identified, with almost half of these (297, 46.70%) observed only once. There was appreciable variation in the frequency of the strain types observed more than 20 times (Table 3).

### Discussion

Although the introduction of PsA-TT into the African meningitis belt has had a major impact on serogroup A epidemics,\(^8,31\) the region will remain at risk of meningococcal disease until comprehensive vaccines targeting all serogroups are available.\(^7\) The reasons for the unique epidemiology of the African meningitis belt remain poorly understood,\(^4,5\) making it difficult to predict when and where epidemics caused by non-serogroup A meningococci might occur. Variations in the prevalence of NPN species, which potentially contribute cross immunity through subcapsular antigens, could play a role in determining susceptibility to a potentially epidemic strain. A number of studies have indicated the movements of genes encoding various protein antigens from NPN to \( \text{Nm} \);\(^4,33\)\(^4,33\) and variants of the FetA antigen have been previously shown to be shared amongst *Neisseria* species.\(^14,34\) This antigen is known to generate protective responses in humans.\(^35\) Colonization with NPNs also affects colonization of humans in experimental studies.\(^18\)

The novel sequence based techniques employed in this study enabled \( \text{Nm} \) and NPN to be identified and characterized rapidly and cost effectively from the very large numbers of samples obtained in the African centers. The \( rplf \) assay\(^22\) achieved reliable speciation, which would not have been possible with conventional methods such as 16S rRNA gene sequencing. An indication of diversity within species was achieved by sequencing the variable region of the gene encoding FetA (\( f\text{eta}_\text{VR} \)), an outer membrane protein (OMP) found in most *Neisseria*, and which is involved in iron metabolism and been shown to elicit protective...
Further characterization of the meningococcal isolates has been previously published. Of the five known Neisseria species identified, with a possible novel species present in Chad, the most common were Nl and Nm. These are the species that have been observed mostly frequently in previous investigations in the African meningitis belt and they are known to have antagonistic interactions in colonization. It is possible that the isolation of some of the other species was affected by the selective media used in this and other studies. Although growth of Np, Nl17,39 and Ns42 on colistin containing agar (such as modified Thayer–Martin or New York city medium) has been reported, other species such as Neisseria perflava, Neisseria sicca, Neisseria mucosa, Neisseria cinerea may not have been identified using the culture technique employed in this study. The impact of these media on Nb is unknown. In future, the identification of NPNs could be enhanced by the use of molecular approaches, although these will have to have species-level resolution, such as the rplF assay. The Neisseria identified were highly diverse at both the strain and the species level and varied markedly over time and place. This is consistent with other carriage studies in the meningitis belt, but is different from the relatively stable carriage observed in countries that do not experience large-scale epidemic disease.

The most common NPNs recovered, Np and Nl, were isolated predominantly from young children as opposed to Nm, which was isolated most frequently from older children. Nl is known to colonize infants and young children preferentially in many settings and this is consistent with carriage of NPNs having a role in the rapid acquisition of antibody against Nm among children in the meningitis belt, although immune responses. Further characterization of the meningococcal isolates has been previously published.

Figure 3  Frequency distribution of fetA VR alleles. Frequency distribution of different alleles of the variable region of fetA. Only the 10 most common alleles are displayed; distribution by country (A), by survey (B) and by species (C).
given the age distribution of meningococcal disease these antibodies may not be protective.46

The risk factors for Nm and NPNs carriage were not the same and in some cases (eg. age, sex and season) were inversely related. Individuals were more likely to carry NPNs during the rainy season if they were a female and under 5 years old whereas they were more likely to carry

Nm during the dry season, if they were male and between 5 and 29 years old.12 These results suggest that the physical presence of an NPN in the pharynx may prevent the colonization by Nm although this may not apply to the hyperinvasive meningococci since the incidence of meningococcal disease in the African meningitis belt is highest in under five year olds. Recent vaccination with a meningitis

| Commensal strain_type | Neisseria bergeri | Neisseria lactamica | Neisseria meningitidis | Neisseria polysaccharea | Total |
|-----------------------|------------------|---------------------|------------------------|-------------------------|-------|
| 1:fnl                 |                  |                     |                        |                         | 470   |
| 2:F1-1                |                  |                     |                        |                         | 452   |
| 6:F1-31               |                  |                     |                        |                         | 205   |
| 2:F5-1                |                  |                     |                        |                         | 135   |
| 6:ND                  |                  |                     |                        |                         | 122   |
| 32:ND                 |                  |                     |                        |                         | 88    |
| 52:ND                 |                  |                     |                        |                         | 82    |
| 2:F1-3                |                  |                     |                        |                         | 74    |
| 1:ND                  |                  |                     |                        |                         | 68    |
| 9:ND                  |                  |                     |                        |                         | 66    |
| 6:F1-29               |                  |                     |                        |                         | 62    |
| 34:F1-62              |                  |                     |                        |                         | 56    |
| 6:F1-62               |                  |                     |                        |                         | 47    |
| 52:F3-60              |                  |                     |                        |                         | 46    |
| 6:F1-100              |                  |                     |                        |                         | 46    |
| 6:F1-72               |                  |                     |                        |                         | 46    |
| 1:F3-1                |                  |                     |                        |                         | 45    |
| 6:F5-133              |                  |                     |                        |                         | 45    |
| 6:F4-5                |                  |                     |                        |                         | 43    |
| 6:F4-17               |                  |                     |                        |                         | 42    |
| 9:F7-3                |                  |                     |                        |                         | 37    |
| 6:F5-84               |                  |                     |                        |                         | 36    |
| 33:F2-17              |                  |                     |                        |                         | 35    |
| 52:F1-21              |                  |                     |                        |                         | 35    |
| 62:ND                 |                  |                     |                        |                         | 35    |
| 33:ND                 |                  |                     |                        |                         | 34    |
| 88:fnl                |                  |                     |                        |                         | 32    |
| 9:F2-24               |                  |                     |                        |                         | 32    |
| 1:F5-5                |                  |                     |                        |                         | 31    |
| 34:ND                 |                  |                     |                        |                         | 31    |
| 69:ND                 |                  |                     |                        |                         | 31    |
| 6:F4-6                |                  |                     |                        |                         | 29    |
| 6:F5-12               |                  |                     |                        |                         | 29    |
| 2:ND                  |                  |                     |                        |                         | 28    |
| 6:F5-18               |                  |                     |                        |                         | 26    |
| 2:F6-3                |                  |                     |                        |                         | 25    |
| 52:F1-29              |                  |                     |                        |                         | 25    |
| 6:F1-21               |                  |                     |                        |                         | 25    |
| 9:F11-4               |                  |                     |                        |                         | 23    |
| 4:F1-7                |                  |                     |                        |                         | 23    |
| 63:F1-72              |                  |                     |                        |                         | 23    |
| 6:F6-2                |                  |                     |                        |                         | 23    |
| 1:F4-23               |                  |                     |                        |                         | 22    |
| 33:F4-6               |                  |                     |                        |                         | 22    |
| 33:F5-34              |                  |                     |                        |                         | 22    |
| 6:F2-23               |                  |                     |                        |                         | 22    |
| 6:F1-120              |                  |                     |                        |                         | 21    |
| 6:F1-101              |                  |                     |                        |                         | 20    |
Carriage of non-meningococcal Neisseria

The shared protein variants could create a cross-reactive immunity at the subscapular level. The inclusion of \textit{fetA} in the meningococcal typing system also increased discrimination between strains (supplemental Table 6). Although there was correlation between \textit{porA} and \textit{fetA} alleles for some \textit{Nm} strains (e.g. \textit{cnf1:P1.18-11,42-1:fnl}), in others the \textit{fetA} \textit{VR} (e.g. W:P1.5,2:F1-1 and W:P1.5,2:F6-3) or both OMPs (e.g. W:P15-1,2-36:F5-1) sequences varied, suggesting that the serogroup \textit{W Nm} strains are antigenically diverse. This has potential implications for using proteins such as \textit{porA} or \textit{fetA} in vaccine formulations.\textsuperscript{48} The Neighbor Joining phylogeny (supplemental Fig. 3) shows the clustering of all of the isolates with the appropriate species except for one, which has the \textit{f_rplF} allele 58 defined previously as \textit{Ns}\textsuperscript{2} but clusters more closely, in this study, with the \textit{Nb} species which was not discovered at the time of the previous study. This particular isolate may also be a \textit{Ns} with a \textit{f_rplF} allele similar to \textit{Nb}. More sequence data from this isolate will clarify this issue.

This study has demonstrated the dynamic nature and high diversity of the genus \textit{Neisseria} in pharyngeal carriage in countries of the African meningitis belt. The results are consistent with the idea that the carriage of NPNs may influence invasive meningococcal disease epidemiology in the African meningitis belt. Although more research is needed to elucidate such effects, understanding these organisms could potentially contribute to meningococcal disease control. This is of particular importance given the absence of comprehensive vaccines against all meningococcal serogroups and the continuing interest in the development of protein based vaccines.

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### Conflict of interest

Caroline Trotter reports that she received a Consulting payment in 2013 for a critical review of health economic model of meningococcal ACWY vaccine by GlaxoSmithKline (GSK); Ray Borrow reports that he performed contract researches on behalf of Public Health England for Novartis Vaccines and Diagnostics, Baxter Biosciences, Sanofi Pasteur, Serum Institute of India and GSK. All other authors report no potential conflicts of interest.

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### Appendix A. Supplementary data

Supplementary data related to this article can be found at http://dx.doi.org/10.1016/j.jinf.2016.03.010.

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