PDF hosted at the Radboud Repository of the Radboud University Nijmegen

The following full text is a publisher's version.

For additional information about this publication click this link.
http://hdl.handle.net/2066/136364

Please be advised that this information was generated on 2020-03-13 and may be subject to change.
Nasopharyngeal Carriage of *Streptococcus pneumoniae* in Pneumonia-Prone Age Groups in Semarang, Java Island, Indonesia

Helmia Farida1*, Juliëtte A. Severin2, M. Hussein Gasem3, Monique Keuter4, Hendro Wahyono1, Peter hans van den Broek5, Peter W. M. Hermans6,7,8, Henri A. Verbrugh2

1 Department of Microbiology, Faculty of Medicine Diponegoro University - Dr. Kariadi Hospital, Semarang, Indonesia, 2 Department of Medical Microbiology and Infectious Diseases, Erasmus University Medical Centre, Rotterdam, the Netherlands, 3 Department of Internal Medicine, Dr. Kariadi Hospital - Faculty of Medicine Diponegoro University, Semarang, Indonesia, 4 Department of General Internal Medicine, Radboud University Nijmegen Medical Centre, Nijmegen, the Netherlands, 5 Department of Infectious Diseases, Leiden University Medical Centre, Leiden, the Netherlands, 6 Nijmegen Institute for Infection, Inflammation, and Immunity (N4I), Nijmegen, the Netherlands, 7 Laboratory of Paediatric Infectious Diseases, Radboud University Nijmegen Medical Centre, Nijmegen, the Netherlands, 8 Crucell - Johnson and Johnson, Leiden, the Netherlands

**Abstract**

*Streptococcus pneumoniae* is a worldwide occurring pathogen. Nasopharyngeal carriage of *Streptococcus pneumoniae* precedes pneumonia and other pneumococcal diseases in the community. Little is known about *S. pneumoniae* carriage in Indonesia, complicating strategies to control pneumococcal diseases. We investigated nasopharyngeal carriage of *S. pneumoniae* in Semarang, Indonesia.

**Methods:** A population-based survey was performed in Semarang, Indonesia. Nasopharyngeal swabs and questionnaires were taken from 496 healthy young (6–60 month-old) children and 45–70 year-old adults.

**Results:** Forty-three percent of children aged 6–60 months and 11% of adults aged 45–75 years carried *S. pneumoniae*. Determinants of carriage were being a child (OR 7.7; 95% CI = 4.5–13.0), passive smoking (OR 2.1; 95% CI = 1.3–3.4), and contact with toddler(s) at home (OR 3.0; 95% CI = 1.9–4.7). The most frequent serotypes found were 6A/B and 15B/C. The current commercially available vaccines cover <50% serotypes found in children. Twenty-four percent of *S. pneumoniae* strains were penicillin non-susceptible, and 45% were resistant to cotrimoxazol.

**Conclusions:** The limited coverage of commercially available vaccines against the serotypes found in this population, and the high proportion of non-susceptibility to penicillin and cotrimoxazol suggest the need for region-specific information and strategies to control *S. pneumoniae*.

**Introduction**

*Streptococcus pneumoniae* is a worldwide occurring pathogen [1]. Data on this species are abundantly available in developed countries, but still scarce in low-to-middle-income countries, leading to difficulties in designing national strategies to control pneumococcal diseases.

Since pneumococcal pneumonia is preceded by nasopharyngeal colonization with *S. pneumoniae* [2], it is relevant to study the nasopharyngeal carriage pattern in humans, particularly in those at higher risk of pneumonia. Pneumococcal carriage has already been extensively studied in many parts of the world, but only few data are available from Indonesia [3], the fourth most populated country in the world. We investigated the nasopharyngeal carriage of *S. pneumoniae* in an urban area of Indonesia, to study the prevalence, risk factors, serotypes, and antimicrobial susceptibility.

**Methods**

**Ethics Statement**

The study was approved by The Ethical Committee of the Faculty of Medicine, Diponegoro University, Semarang. Written informed consent was given by the subjects or their caregivers.

**Subjects**

A population-based survey was performed in Semarang, a city with 1.5 million residents in Central Java, among healthy children aged 6–60 months and healthy adults aged 45–70 years as described before [4]. Exclusion criteria were the presence of respiratory symptoms and antibiotic consumption within the last three days. Cluster random sampling was done from February to April 2010 to recruit subjects from all 16 districts of Semarang.

---

* E-mail: helmia_farida@yahoo.com
Specimen Collection and Laboratory Testing

Nasopharyngeal swabs were obtained using rayon-tipped swabs and transported in Amies-charcoal media (COPAN, Italy). Swabs were inoculated on 5% sheep blood agar with gentamicin (5 mg/liter) and incubated at 35°C in 5% CO₂ for 48 hours. Identification of S. pneumoniae was performed using the optochin test (Oxoid, Basingstoke, UK) and, in case of doubt, a DNA hybridization test (Accuprobe, Gen-Probe Inc., San Diego, CA, USA). Antimicrobial susceptibility tests were performed using disk diffusion method (Oxoid, UK) and E-test (bioMérieux, France) and interpreted according to EUCAST 2012. Serotyping of S. pneumoniae was done with a multiplex-PCR which covers 36 serotypes [5,6]. Control strains were included in all analyses.

Data on demography, house sanitation (crowding, smoke exposure from cigarette and mosquito coils), and water and food hygiene, were recorded using a questionnaire that was developed to identify determinants of carriage. Crowding was defined to be present when the ratio of total bedroom space to the number of family members was less than 4 m² [7]. Water hygiene was defined as poor when water other than tap or bottled water was used by the family. Food hygiene was considered poor if the family consumed street food.

Statistical Analysis

Univariate analysis was done with Chi-square or Fisher’s exact tests when appropriate, followed by backward stepwise logistic regression for variables with P value <0.2 using SPSS 17 (SPSS Inc, Chicago, USA). P value of <0.05 was considered significant.

Results

Subjects

Two hundred and fifty-three adults aged 45–70 years and 243 children aged 6–60 months participated in the study. The characteristics of the participants have been presented previously [4]. Crowding was common, as was exposure to smoke.

Carriage Prevalence and Determinants

Overall carriage of S. pneumoniae was 27% (95% CI: 20–32), 43% in children (95% CI: 32–50) and 11% in adults (95% CI: 5–15). The proportion carrying S. pneumoniae varied significantly across the districts of Semarang (P<0.05), and tended to be higher in the suburban and eastern parts of the city (Figure 1). Multivariate analysis showed that being a child (OR 7.7, 95% CI, 4.5–13.0), passive smoking (OR 2.1, 95% CI, 1.4–3.4), and contact with toddler(s) at home (OR 3.0, 95% CI, 1.9–4.7) were independent determinants of carriage.

Antimicrobial Susceptibility and Serotypes

One hundred and forty-two strains were isolated from 133 subjects. In total, 34 (24%) strains were penicillin non-susceptible (MIC ranged 0.047–1.5), including 25 (23%) from children and 9 (29%) from adults (P=0.25). Forty-five percent of the strains were resistant to cotrimoxazol, 1% to erythromycin, and 5% to tetracycline. There was no significant difference in the susceptibility pattern between isolates from children and those from adults (P>0.1). No strain was resistant to neither penicillin nor vancomycin.

Capsular type 6A/B was the most prevalent serotype in all age groups (19% in children and 39% in adults). The most common capsular serotypes in children, comprising 61% of strains, were 6A/B, 15B/C, 11A, 23F, 19F, 23A. Those in adults, were 6A/B, 15B/C, and 15A. These two serotype patterns differed significantly (P=0.029). Other serotypes were less frequently found (Table 1). Twenty percent were un-typeable with the multiplex-PCR employed.

Discussion

The carriage prevalence of S. pneumoniae among children in our study was comparable to those previously found among healthy children on Lombok island, Indonesia [3], and in the Netherlands [8]. However, it was lower than those in Gambia [9], Poland [10], Australia [11], Thailand [12], and higher than those reported from Iran [13] and Korea [14]. The carriage prevalence among adults in our study was 11%, which is higher than that found in Alaska [15], but lower than that among Australian Aboriginals of the same age [11]. The prevalence differences among populations may be related to sampling or laboratory methods (i.e. nasopharyngeal swab versus throat swab), the use of selective media), to certain characteristics of the population studied (i.e. the age of the subjects, household characteristics — especially the presence of toddlers, presence of upper respiratory tract infection, vaccination status), or to seasonal variation. Our samples were taken in the rainy season, during which the incidence of respiratory tract

Table 1. Serotype of S. pneumoniae isolated from healthy people in Semarang, Indonesia.

| Capssular Type | Children | Adults | Total | P   |
|---------------|----------|--------|-------|-----|
|               | n (%)    | n (%)  | n (%) |     |
| 6A/6B         | (19)     | (39)   | (23)  | 0.029 |
| 15B/C         | (10)     | (13)   | (11)  |     |
| 11A           | (10)     | (3)    | (8)   |     |
| 23F           | (9)      | (3)    | (6)   |     |
| 19F           | (5)      | (0)    | (4)   |     |
| 23A           | (2)      | (10)   | (4)   |     |
| 15A           | (2)      | (10)   | (4)   |     |
| Others        | (20)     | (6)    | (26)  | 0.017 |
| Un-typeable   | (20)     | (8)    | (28)  | 0.020 |
| Total         | (111)    | (31)   | (142) |     |

doi:10.1371/journal.pone.0087431.t001

Figure 1. Distribution of nasopharyngeal carriage of S. pneumoniae among healthy population in the districts of Semarang, Indonesia.

doi:10.1371/journal.pone.0087431.g001
infection, transmission of pathogens, and thus, carriage is likely to be somewhat increased.

The prevalence of S. pneumoniae with reduced susceptibility to penicillin and cotrimoxazole was high. The national and local guidelines for empirical antibiotics for community-acquired pneumonia in children still recommend these two antibiotics as the first choices [16], and those for meningitis recommend ampicillin for the second line [17].

The commercially available 13-valent pneumococcal conjugate vaccines (PCV13) [18], which was introduced only in 2011 in Indonesia, provides approximately 45% strain coverage for the infant population in this study, varying from 13–100% across the city districts. PCV10, introduced in 2010, provides a little bit lower coverage. However, the coverage of the PCV13 over the serotype repertoire on Lombok island in Indonesia in the past [3] was 60% and in other Southeast Asian countries, the coverage ranged from 63%–97% [19]. Our results may, thus, not be taken to reflect the serotype distribution throughout Indonesia, since the study was performed in a specific geographic location.

The PCVs have not been included in the national vaccination programs since information regarding the burden of pneumococcal diseases in Indonesia is still lacking. Rather, pneumococcal vaccines have been introduced in private clinics. This vaccine is rather expensive for regular Indonesian households that it is provided such data and inform public health policies.

**References**

1. Linares J, Artanany C, Pallares R, Fenoll A (2010) Changes in antimicrobial resistance, serotypes and genotypes in *Streptococcus pneumoniae* over a 30-year period. Clin Microbiol Infect 16: 402–410.
2. Cardozo DM, Nascimento-Carvalho GM, Andrade AL, Silvany-Neto AM, Daltro GH, et al. (2006) Prevalence and risk factors for nasopharyngeal carriage of *Streptococcus pneumoniae* among adolescents. J Med Microbiol 57: 185–189.
3. Soewignjo S, Gessner BD, Sutanto A, Steinhoff M, Prijanto M, et al. (2001) *Streptococcus pneumoniae* nasopharyngeal carriage prevalence, serotype distribution, and resistance patterns among children on Lombok Island, Indonesia. Clin Infect Dis 32: 1093–1094.
4. Farhida H, Severin JA, Gassen MH, Keuter M, van den Broek PJ, et al. (2013) Nasopharyngeal carriage of *Klebsiella pneumoniae* and other Gram-negative bacteria in pneumonia-prone age groups in Semarang, Indonesia. J Clin Microbiol 51: 1614–1616.
5. Center for Disease Control and Prevention (CDC) (2011) PCR Deduction of Pneumococcal Serotypes.
6. Saha SK, Darmstadt GL, H Baqui A, Hossain B, Arifeen SE, et al. (2008) *Streptococcus pneumoniae* serotypes causing invasive and non-invasive disease in *Klebsiella pneumoniae* and other Gram-negative bacilli in pneumonia-prone age groups in Semarang, Indonesia. J Clin Microbiol 51: 1614–1616.
7. The Ministry of Health of Republic Indonesia (1999) Peraturan Menenti Kesehatan Republik Indonesia No 829/1999 tentang Persyaratan Rumah Sihat (The Decree of The Minister of Health of Indonesia Number 829/1999 regarding with Health Requirements of Housing). In: The Ministry of Health Republic Indonesia, editor.
8. Bogart D, Engelen NM, Timmers-Reker AJM, Elzaaaz KP, Peerbooms PGH, et al. (2005) Pneumococcal carriage in children in the Netherlands: A molecular epidemiological study. J Clin Microbiol 39: 3316–3320.
9. Hill PC, Townsend J, Antonio M, Akisanya B, Ebruke C, et al. (2010) Transmission of *Streptococcus pneumoniae* in rural Gambian villages: A longitudinal study. Clin Infect Dis 50: 1460–1467.
10. Koroma-Głoweniak I, Niedzielski A, Malm A (2011) Upper respiratory colonization by *Streptococcus pneumoniae* in healthy pre-school children in south-east Poland. Int J Pediatr Otorhinolaryngol 75: 1529–1534.
11. Mackenzie GA, Leach AJ, Carapetis JR, Fisher J, Morris PS (2010) Epidemiology of nasopharyngeal carriage of respiratory bacterial pathogens in children and adults: cross-sectional surveys in a population with high rates of pneumococcal disease. BMC Infect Dis 10: 304.
12. Levine S, Dejniratt S, Sangsuk L, Chanta R, Seikin DR, et al. Serotypes and antimicrobial resistance of *Streptococcus pneumoniae* in Thailand 2002–2004: The Paed Infect Disease J 25: 176–178.
13. Balsbhae M, Naderi HR, Ghazvini K, Sotoudeh K, Amali A, et al. (2012) Passive smoking and nasopharyngeal colonization by *Streptococcus pneumoniae*, *Haemophilus influenzae*, and *Moraxella catarrhalis* in daycare children. Eur Arch Otorhinolaryngol 269: 1127–1132.
14. Kuo CY, Hwang KP, Hsieh YC, Cheng CH, Huang FL, et al. (2011) Nasopharyngeal carriage of *Streptococcus pneumoniae* in Taiwan before and after the introduction of a conjugate vaccine. Vaccine 29: 5171–5177.
15. Rudolph KM, Parkinson AJ, Reasonover AL, Bulkow LR, Debra J, Parks, et al. (2000) Serotype distribution and antimicrobial resistance patterns of invasive isolates of *Streptococcus pneumoniae*: Alaska, 1991–1998. J Infect Dis 182: 490–496.
16. Direktorat Jenderal Pengendalian Penyakit dan Penyehatan Lingkungan - Kementerian Kesehatan Republik Indonesia (2011) *Streptococcus pneumoniae* serotypes causing invasive disease in Indonesia. *Jurnal Kesehatan RPJMP*.
17. Ting Adaptasi Indonesia (2008) Buku saku pelayanan kesehatan anak di rumah sakit: Pedoman bagi rumah sakit rujukan tingkat pertama di kabupaten/kota Kementerian Kesehatan RI - World Health Organization - Banten Dokter Anak Indonesia.
18. Pfister (2011) Pneumococcal 15-valent conjugate vaccine (diphtheria CRM197 protein). In: Ine WP, editor. USA.
19. Janczakiewicz E, Jewers JM, Hibberd ML, Clarke SC (2012) Prevalence of *Streptococcus pneumoniae* serotypes causing invasive and non-invasive disease in South East Asia. A review. Vaccine 30: 3503–3514.
20. Yuliarti K, Hadinegoro SR, Supriyatno B, Karuniawati A, et al. (2012) Passive smoking and nasopharyngeal colonization by *Streptococcus pneumoniae*, *Haemophilus influenzae*, and *Moraxella catarrhalis* in daycare children. Eur Arch Otorhinolaryngol 269: 1127–1132.
21. Purniti PS, Subanada IB, Kari IK, Arhana B, Iswari IS, et al. (2011) Surveilaince of *Streptococcus pneumoniae* Carriage in Indonesia using locally implementable laboratory methods.

**Author Contributions**

Conceived and designed the experiments: HAV JAS PWMH MK. Performed the experiments: HF HW JAS PWMH. Analyzed the data: HF HAV JAS PWMH. Contributed reagents/materials/analysis tools: HAV JAS PWMH MK. Wrote the paper: HF JAS PWMH MK HW.

**Acknowledgments**

We thank students from Faculty of Medicine Diponegoro University, Suisslo Pehranito from Internal Medicine Residency Program, Faculty of Medicine Diponegoro University, Christa van der Gaast-de Jongh from the Laboratory of Pediatric Infectious Diseases, Niijmegen, Mitchell Laurens and technicians from laboratory of Medical Microbiology and Infectious Diseases, Erasmus Medical Centre, Rotterdam for their excellent technical assistance and support.