

LETTER TO THE EDITOR

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Nosocomial outbreak of multi-resistant *Streptococcus pneumoniae* serotype 15A in a centre for chronic pulmonary diseases

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Abstract

We report nosocomial transmission of multi-resistant serotype 15A *Streptococcus pneumoniae* (MRSP) that resulted in two lower respiratory tract infections in a centre for chronic pulmonary diseases. This outbreak highlights the potential for transmission of MRSP among vulnerable patients when laboratory turnaround time is long and patient compliance with transmission-based precautions is low.

Keywords: *Streptococcus pneumoniae*, Cross infection, Nosocomial, Antimicrobial resistance, Whole-genome sequencing

Main text

Streptococcus pneumoniae is a major cause of community-acquired pneumonia, bacteraemia and meningitis [1]. Asymptomatic nasopharyngeal colonization is a predisposing factor for pneumococcal infection [2]. Infections arising in hospitalized patients are often regarded as a result of earlier acquisition in the community [2], although hospital outbreaks of susceptible and resistant pneumococci have been reported [3–6]. Here, we describe two risk factors for nosocomial transmission of multi-resistant *Streptococcus pneumoniae* (MRSP) that resulted in two cases of hospital-acquired pneumonia.

Patient A (index case) was admitted to the Radboud University Medical Center Dekkerswald, a centre for pulmonary rehabilitation, tuberculosis and lung diseases (Groesbeek, The Netherlands) because of psychosocial issues and a mild exacerbation of chronic obstructive pulmonary disease (COPD) probably due to smoking. There were no signs of infection. This patient had been a carrier of MRSP 3.5 years ago, but as MRSP had not been identified in subsequent sputum cultures the patient was regarded as MRSP negative. On day 9 after admission a

sputum sample collected on day 3 yielded *S. pneumoniae* that was intermediately susceptible to penicillin and resistant to erythromycin, tetracycline and clindamycin (Fig. 1) indicating that patient A was an asymptomatic carrier of MRSP again. Subsequently, the patient was put on droplet and contact isolation according to protocol and discharged on day 17 (Fig. 1). Patient B was admitted to an adjacent room on the same day as patient A because of an exacerbation of COPD. Patient B was discharged after 6 days; readmitted three days later (on day 10) with a hospital acquired pneumonia, and placed on contact precautions on day 11 because of a rhino- or enterovirus positive throat swab. On day 17 (day 7 after second admission) *S. pneumoniae* with an antibiotic susceptibility pattern similar to that of patient A was isolated from a sputum sample collected on admission. Extension to droplet and contact isolation for MRSP was, however, not installed. Patient B was discharged on day 20. On day 27 patient C was admitted with an exacerbation of COPD and that same day patient A was readmitted with recommended isolation procedures but the patient's adherence to the instructions for transmission-based precautions was poor. The patients were placed in adjacent rooms. On day 32 patient C was diagnosed with a hospital acquired pneumonia and *S. pneumoniae* with an identical susceptibility profile to that of patient A was isolated from patient C's sputum, three days later. Following isolation of MRSP, patient C

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patients B and C cannot be fully ruled out, multiple arguments support nosocomial acquisition. The pneumococcal strain concerned is rarely encountered worldwide and is here proven to be communicable by WGS. Furthermore, patients B and C have been directly exposed to air droplets from patient A during regular chats, while no common source or link outside the hospital could be identified (patients were unrelated and living in different areas). Possible transmission via HCWs was investigated, yet no MRSP carriage could be identified among them, and high hand hygiene compliance (directly related to pneumococcal transmission [10]) suggested appropriate attention for hygiene precautions. Transmission to patient B probably occurred while patient A was not immediately isolated due to time needed for bacterial culture and susceptibility testing (Fig. 1). Patient C was probably infected because patient A was not adhering to droplet and contact precautions. WGS supported our suspicion that patient A was the index case, whose MRSP isolated 3.5 years ago showed 8 SNPs difference from the current isolate, compared to 2 SNPs difference between the isolates from the 3 patients involved in this outbreak.

This outbreak highlights two risk factors for nosocomial transmission of MRSP causing hospital-acquired infections (patients B and C). First, time needed for laboratory diagnosis in low-endemic regions may lead to failure to timely install adequate transmission-based precautions. In this case the delay was due to the atypical colony morphology and resistance pattern necessitating confirmatory tests. Secondly, patient compliance with transmission-based precautions may be low despite standard instructions and may require day-to-day audits.

Considering the fact that patient A still turned out to be a MRSP-carrier after 3.5 years and recent evidence suggests that *S. pneumoniae* serotype and drug-resistance are associated with carriage duration [11], additional measures to prevent nosocomial spread in low-endemic regions could include routine screening at admission for patients who were colonized with MRSP previously, and placing them on droplet and contact precautions until carriage of MRSP has been ruled out.

Abbreviations

COPD: Chronic obstructive pulmonary disease; HCWs: Healthcare workers; MLST: Multi locus sequence type.; MRSP: Multi-resistant *Streptococcus pneumoniae*; SNP: Single nucleotide polymorphism; WGS: Whole-genome sequencing

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Availability of data and materials

All data generated or analysed during this study are included in this published article. In addition, raw sequence reads are stored in the SRA database of NCBI under following accession numbers: SAMN09812500, SAMN09812501, SAMN09812502, SAMN09812503, SAMN09812504, SAMN09812505, SAMN09812506, and part of BioProject: 485675.

Authors' contributions

GJHB, MTN, MJB, JH and HFLW were involved in management of the outbreak. GJHB, MTN, AJHC and JPMC are responsible for clinical, laboratory and WGS data collection and analysis. GJHB drafted the manuscript while critical revision was provided by AJHC, JPMC, JH and HFLW. All authors read and approved the final version of the manuscript.

Ethics approval and consent to participate

The research ethics committee of the Radboudumc judged that the study does not fall within the remit of the Medical Research Involving Human Subjects Act (WMO). The study has been reviewed by the research ethics committee on the basis of the Dutch Code of conduct for health research, the Dutch Code of conduct for responsible use, the Dutch Personal Data Protection Act and the Medical Treatment Agreement Act. The ethics committee has passed a positive judgment on the study.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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