Invasive Pneumococcal Disease (IPD) Serotype Frequency in Iranian Patients

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1. Background

Streptococcus pneumoniae as a Gram positive diplococcic is a major worldwide causative agent of morbidity and mortality among young children and the aged (1). In addition, Streptococcus pneumoniae is a versatile human pathogen causing infectious disease ranging from mild infection (i.e. otitis media) to life threatening pneumonia and meningitis in many countries (2). The most prevalence of Invasive pneumococcal disease caused by Streptococcus pneumoniae is acute Otitis Media, Pneumonia and Meningitis (3). Among this Pneumonia is the main infection with immediate cure needed for patients (4). It has to be said that Pneumococcal account for most cases of community-acquired (5). The research reports that, 92 capsular serotypes of Streptococcus pneumoniae differ greatly in nasopharyngeal carriage prevalence, invasiveness, and disease incidence (6). Therefore, there has been some necessitate, though, regarding whether serotype independently affects the outcome of invasive pneumococcal disease (IPD).

2. Objectives

The aim of this study was to determine the frequency type of Streptococcus pneumoniae in Invasive pneumococcal disease (IPD) in Iranian patients.

3. Materials and Methods

The present study was conducted on as a prospective study to type the profile of pneumococcal strains from Invasive Pneumococcal Disease (IPD) across in Iran. During two years (2009 – 2010) all invasive Pneumococcal strains isolates from selected laboratory hospitals in Iranian central provinces were collected. The strains were isolate from culture of throat, Blood, Sputum, CSF were fluid; eye and lung infection. They were transfer to central Baqiyatallah laboratory and re-identified based on bacteriological standard methods. A total of 135 bacteria strain suspected of invasive streptococcal disease (IPD) were subjected to PCR, bacteriological methods and serotyped. All isolates were serotyped by the Quelling reaction using specific antiserum (Statens Serum Institute, Copenhagen, Denmark) based on manufacture procedure. Out of which, 134 isolated strains of S. pneumoniae was confirmed by PCR method.

Implication for health policy/practice/research/medical education:
This should be used for all of the health policy makers, and area of medical education and research as well as practice.

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3.1. Laboratory Investigation

The isolated invasive streptococcal strains were re-identified as pneumococcal according to Standardized Procedure to culture and analyze (-hemolysis, colony morphology, optochin susceptibility, and bile solubility) (7). The samples have been tested on the Blood agar and Optochin sensitivity and bile solubility as well as Gram Staining was performed too. The positive results were noted. A reaction in which anticapsular antibodies bind to the capsule of a bacterium, resulting in the capsule to swell or become more visible, was done and it was seen under the microscope. The pneumococcal isolates were then typed by immunological techniques according to Company Procedure (SSI. Denmark) (8). The demographic data including (age, sex, site of infection) was accumulated. The data was inserted in the computer and then specific Pneumococcal types were matched and then collected data analyzed by SSPS version 17.0.

4. Results

The results of this study showed some S. pneumoniae serotypes were found in both sexes and some only in one sex invasive infections. For example, serotypes 10, 14, 18 and 22 were only in female patients with infections. While serotypes 20 were only isolated from men infected. The analysis of the results had suggested that serotypes 6 and 19 are the most abundant bacterial strains isolated from patients. (See Figure 1 and Figure 2). The analysis results of serotype distribution based on site of infections were showed in Table 1.

![Figure 1. Types of S. pneumoniae and Age of Patients](image)

![Figure 2. Types of S. pneumoniae and Sex of Patients](image)

| Table 1. Serotype Distribution and Site of Isolation of Invasive Streptococcal Strains |
|---------------------------------|-----------------|------------------|
| Site of Isolation | Number of Frequency | Serotype or Serogrope |
| Wounds | 3 | 6 |
| | 3 | 5 |
| | 2 | 10 |
| Rhinit | 4 | 7 |
| Sinus | 3 | 20 |
| Eye | 9 | 6 |
| | 4 | 1 |
| | 2 | 14 |
| | 11 | 19 |
| Throat | 3 | 19 |
| | 3 | 7 |
| | 3 | 22 |
| | 5 | 20 |
| | 6 | 14 |
| | 4 | 3 |
| | 5 | 4 |
| | 3 | 1 |
| Lung | 2 | 1 |
| | 5 | 2 |
| | 3 | 7 |
| | 3 | G |
| | 4 | 5 |
| | 9 | 6 |
| | 5 | 8 |
| | 3 | 4 |
| | 3 | 17 |
| | 2 | 18 |
| blood | 1 | 17 |
| | 5 | G |
| | 1 | 1 |
| | 5 | 2 |
| | 3 | 6 |
| CSF | 3 | 20 |
| | 3 | 8 |
5. Discussions

Our data is different from other research. For example, some studies were shown that, the serotypes 1 and 5 were commonly causing IPD (<9-13>). While in this study, the serotypes 6 and 19 were the most common involved in invasive streptococcal infections (Table 1). Recently, in national level the results of a report was published has been shown similar data. They were serotyped only 75 S. pneumoniae strains and due to limited sample have claimed that heptavalent pneumococcal conjugate vaccine (PCV7) are covering 80% of the serotypes (14). Based on the results of this study, the determine serotypes involved in invasive infections are equal the 23 valent pneumococcal vaccine (PPV23). However, the compare data but a few studies may be similar typing results in different country. In Denmark overall, 92% (93% blood, 87% CSF) of isolates and 94% of all childhood isolates belonged to the 23 vaccine types. In addition, In Denmark, the ten most frequently occurring types from children were (6A + 6B, 18C, 14, 1, 7F, 19F, 9V, 4, and 23F) (15). Pneumococcal types may be different country by country for this reason different vaccine (Heptavalent Pneumococcal Conjugate Vaccine, 7 valant or 13-valent pneumococcal conjugate vaccine or even 23 valents has been applied in different countries in order to prevent the disease in different age groups (16-18). Hence, the only vaccine that is able coverage serotypes our country is the 23-valent vaccine. Molecular typing must be applied in the further be helped the details (19).

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Authors’ Contribution

None declared.

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References

1. Centers for Disease Control and Prevention. Preventing Pneumococcal Disease Among Infants and Young Children. Morb Mortal Wkly Rep. 2000;49(2):3-8.
2. Holmlund E. Mother-Infant Antibodies to Pneumococcal Polysaccharides and Proteins: Transfer and Persistence of Maternal Antibodies and Development of Vaccine-Induced and Naturally Acquired Antibodies in Infants. National Institute for Health and Welfare. 2011.
3. Roche PW, Krause V, Cook H, David Coleman JB, Sweeny A, Fielding J, et al. Invasive pneumococcal disease in Australia, 2006. 32. 2006.
4. Henrichsen J. Six newly recognized types of Streptococcus pneumoniae. J Clin Microbiol. 1995;33(10):2759-62.
5. Brueggemann AB, Pai R, Crook DW, Beall B. Vaccine escape recombinants emerge after pneumococcal vaccination in the United States. PLoS Pathog. 2007;3(1).
6. Weinberger DM, Harboe ZB, Sanders EA, Ndritu M, Klugman KP, Ruckinger S, et al. Association of serotype with risk of death due to pneumococcal pneumonia: a meta-analysis. Clin Infect Dis. 2010;51(6):692-9.
7. Murray PR, Baron EJ. Manual of Clinical Microbiology. Oxford: Blackwell Distributer; 2007.
8. Sorensen UB. Typing of pneumococci by using 12 pooled antisera. J Clin Microbiol. 1993;31(8):2097-100.
9. Scott JA, Hall AJ, Hannington A, Edwards R, Mwarumba S, Lowe B, et al. Serotype distribution and prevalence of resistance to benzylpenicillin in three representative populations of Streptococcus pneumoniae isolates from the coast of Kenya. Clin Infect Dis. 1998;27(6):1442-50.
10. Brueggemann AB, Peto TE, Crook DW, Butler JC, Kristinsson KG, Spratt BG. Temporal and geographic stability of the serogroup-specific invasive disease potential of Streptococcus pneumoniae in children. Infect Dis. 2004;190(7):2320-31.
11. Chiba N, Morozumi M, Sunaoshi K, Takahashi S, Takano M, Komori T, et al. Serotype and antibiotic resistance of isolates from patients with invasive pneumococcal disease in Japan. Epidemiol Infect. 2010;138(10):1668.
12. Hausdorff WP, Hajhej R, Al-Mazrou A, Shibil A, Soriano-Gabarro M. The epidemiology of pneumococcal, meningococcal, and Haemophilus disease in the Middle East and North Africa (MENA) region–current status and needs. Vaccine. 2007;25(10):1935-44.
13. Vestrheim DF, Hobly EA, Aasberge IS, Caugant DA. Impact of a pneumococcal conjugate vaccination program on carriage among children in Norway. Clin Vaccine Immunol. 2010;17(5):525-34.
14. Bokaria M, Khazaie HA, Javadimehr M. Serotype distribution and antimicrobial resistance of invasive Streptococcus pneumoniae isolates from children in Zahedan, Iran. Afr J Microbiol Res. 2012;6(1):28-33.
15. Nielsen SW, Henriksen J. Incidence of invasive pneumococcal disease and distribution of capsular types of pneumococci in Denmark, 1989–94. Epidemiol & Infection. 1996;117(3):417-25.
16. Streptococcus laboratory: PCR deduction of pneumococcal serotypes. PCR Deduction of Pneumococcal Serotypes. J Infect Dis. 2006;194(5):682-8.
17. Bradford DGessner, Judith EMueller, Seydou Yaro. African meningitis belt pneumococcal disease epidemiology indicates a need for an effective serotype 1 containing vaccine, including for older children and adults. BMC Infect Dis. 2010;10(22)30.
18. Enright Mark, Spratt Brian. A multisecuence typing scheme for Streptococcus pneumoniae: identification of clones associated with serious invasive disease. Microbiology. 1998;144(1):3049-3060.