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Recent Trends in Invasive Pneumococcal Disease in Korea in the Post-pneumococcal Vaccine Era

Streptococcus pneumoniae can cause invasive pneumococcal disease (IPD), including meningitis, pneumonia, and sepsis. IPD is a leading cause of high morbidity and mortality, especially in children and the elderly [1]. The distribution of pneumococcal serotypes causing IPD varies over time, by age group, and by geographic region [2]. Since the development of pneumococcal conjugate vaccines (PCVs), most developed countries have included PCVs in their national immunization program (NIP). The implementation of PCVs targeting specific serotypes has reduced the morbidity and mortality of IPD patients. However, the use of vaccines has resulted in a shift in serotype distribution, termed the “serotype replacement” phenomenon. The prevalence of IPD due to vaccine serotypes (VTs) has decreased, while IPD caused by non-vaccine serotypes (NVTs) has increased since the advent of vaccines [3-5]. Serotype replacement in the post-pneumococcal vaccine era can be accompanied by a change in antibiotic resistance; in the USA, increases in antibiotic-non-susceptible IPD caused by NVTs have been recently observed, although the overall non-susceptible IPD incidence decreased since the introduction of PCVs [6-8]. Emerging NVTs associated with different pneumococcal lineages in different countries have been reported [6, 7]. Therefore, surveillance for pneumococcal epidemiology, including serotypes and antimicrobial resistance distribution, is crucial to understand the effects of vaccination and guide vaccine development and recommendations [1, 2].

In this issue, Kim, *et al.* [9] report the serotype distribution and antimicrobial resistance of *S. pneumoniae* causing IPD in children and adults in Korea between 2017 and 2019. The authors previously surveyed IPD between 2014 and 2016 [10]. In Korea, 10- and 13-valent PCVs (PCV10 and PCV13, respectively) were introduced in the NIP for children <5 years in 2014. The NIP has been providing pneumococcal polysaccharide vaccine (PPSV23) to elderly people aged ≥65 years since May 2013. The previous study by the authors revealed significant changes in the major serotypes in the community; an increase in NVTs, especially in children, was confirmed [10]. In their present study, recent surveillance data from 16 hospitals are presented, and the serotype distribution and antimicrobial resistance changes were compared to previous data [9]. The data reflect the effects of PCV13, which is primarily used since the introduction of the NIP [11]. Data analyses based on age group, serotype, and antibiotic-specific trends in IPD in Korea reveal a remarkable increase in some serotypes as compared with the previous report [10]. A notable trend was the rapid increase in the incidence of serotype 10A in children ≤5 years. This study will be of help to readers seeking a comprehensive insight into the nationwide trends in IPD in Korea. Continuous IPD surveillance will be needed to understand the dynamics of serotype changes and guide vaccination policies.



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AUTHOR CONTRIBUTIONS

Huh HJ and Sung H contributed to manuscript writing and approved the submission of the final manuscript.

CONFLICTS OF INTEREST

None declared.

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REFERENCES

1. de Miguel S, Domenech M, González-Camacho F, Sempere J, Vicioso D, Sanz JC, et al. Nationwide trends of invasive pneumococcal disease in Spain from 2009 through 2019 in children and adults during the pneumococcal conjugate vaccine era. *Clin Infect Dis* 2021;73:e3778-87.
2. Grant LR, Slack MPE, Theilacker C, Vojcic J, Dion S, Reinert RR, et al. Distribution of serotypes causing invasive pneumococcal disease in children from high-income countries and the impact of pediatric pneumococcal vaccination. *Clin Infect Dis* 2022;ciac475.
3. Zhou M, Wang Z, Zhang L, Kudinha T, An H, Qian C, et al. Serotype distribution, antimicrobial susceptibility, multilocus sequencing type and virulence of invasive *Streptococcus pneumoniae* in China: a six-year multicenter study. *Front Microbiol* 2022;12:798750.
4. Yun KW, Rhie K, Kang JH, Kim KH, Ahn JG, Kim YJ, et al. Emergence of serotype 10A-ST11189 among pediatric invasive pneumococcal diseases, South Korea, 2014-2019. *Vaccine* 2021;39:5787-93.
5. Kakuta R, Nakano R, Yano H, Ozawa D, Ohta N, Matsuoka T, et al. First two cases of infected aortic aneurysm caused by non-vaccine *Streptococcus pneumoniae* serotype 23A. *Ann Lab Med* 2020;40:270-3.
6. Ouldali N, Levy C, Varon E, Bonacorsi S, Béchet S, Cohen R, et al. Incidence of paediatric pneumococcal meningitis and emergence of new serotypes: a time-series analysis of a 16-year French national survey. *Lancet Infect Dis* 2018;18:983-91.
7. Lo SW, Gladstone RA, van Tonder AJ, Lees JA, du Plessis M, Benisty R, et al. Pneumococcal lineages associated with serotype replacement and antibiotic resistance in childhood invasive pneumococcal disease in the post-PCV13 era: an international whole-genome sequencing study. *Lancet Infect Dis* 2019;19:759-69.
8. Bajema KL, Gierke R, Farley MM, Schaffner W, Thomas A, Reingold AL, et al. Impact of pneumococcal conjugate vaccines on antibiotic-nonsusceptible invasive pneumococcal disease in the United States. *J Infect Dis* 2022;jiac154.
9. Kim GR, Kim EY, Kim SH, Lee HK, Lee J, Shin JH, et al. Serotype distribution and antimicrobial resistance of *Streptococcus pneumoniae* causing invasive pneumococcal disease in Korea between 2017 and 2019 after introduction of the 13-valent pneumococcal conjugate vaccine. *Ann Lab Med* 2023;43:45-54.
10. Park DC, Kim SH, Yong D, Suh IB, Kim YR, Yi J, et al. Serotype distribution and antimicrobial resistance of invasive and noninvasive *Streptococcus pneumoniae* isolates in Korea between 2014 and 2016. *Ann Lab Med* 2019;39:537-44.
11. Sohn S, Hong K, Chun BC. Evaluation of the effectiveness of pneumococcal conjugate vaccine for children in Korea with high vaccine coverage using a propensity score matched national population cohort. *Int J Infect Dis* 2020;93:146-50.

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