

Original Article

Infectious Diseases,
Microbiology & Parasitology



Etiology of Invasive Bacterial Infections in Immunocompetent Children in Korea (2006–2010): a Retrospective Multicenter Study

Kyuyol Rhie ¹, Eun Hwa Choi ^{1,2}, Eun Young Cho ^{2,3}, Jina Lee ⁴,
Jin Han Kang ⁵, Dong Soo Kim ⁶, Yae-Jean Kim ⁷, Youngmin Ahn ⁸,
Byung Wook Eun ⁸, Sung Hee Oh ⁹, Sung-Ho Cha ¹⁰, Young Jin Hong ¹¹,
Kwang Nam Kim ¹², Nam Hee Kim ¹³, Yun-Kyung Kim ¹⁴, Jong-Hyun Kim ¹⁵,
Taekjin Lee ¹⁶, Hwang Min Kim ¹⁷, Kun Song Lee ¹⁸, Chun Soo Kim ¹⁹,
Su Eun Park ²⁰, Young Mi Kim ²⁰, Chi Eun Oh ²¹, Sang Hyuk Ma ²²,
Dae Sun Jo ²³, Young Youn Choi ²⁴ and Hoan Jong Lee ^{1,2}

OPEN ACCESS

Received: Sep 5, 2017

Accepted: Nov 22, 2017

Address for Correspondence:

Hoan Jong Lee, MD, PhD

Department of Pediatrics, Seoul National University Children's Hospital, 101 Daehak-ro, Jongno-gu, Seoul 03080, Korea.
E-mail: hoanlee@snu.ac.kr

© 2018 The Korean Academy of Medical Sciences.

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<https://creativecommons.org/licenses/by-nc/4.0/>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ORCID iDs

Kyuyol Rhie
<https://orcid.org/0000-0002-8168-4832>
Eun Hwa Choi
<https://orcid.org/0000-0002-5857-0749>
Eun Young Cho
<https://orcid.org/0000-0002-2286-4593>
Jina Lee
<https://orcid.org/0000-0002-3435-251X>
Jin Han Kang
<https://orcid.org/0000-0003-1610-6742>
Dong Soo Kim
<https://orcid.org/0000-0002-3270-506X>
Yae-Jean Kim
<https://orcid.org/0000-0002-8367-3424>
Youngmin Ahn
<https://orcid.org/0000-0002-1697-8041>

¹Department of Pediatrics, Seoul National University Children's Hospital, Seoul, Korea
²Department of Pediatrics, Seoul National University College of Medicine, Seoul, Korea
³Department of Pediatrics, Chungnam National University School of Medicine, Daejeon, Korea
⁴Department of Pediatrics, University of Ulsan College of Medicine, Asan Medical Center, Seoul, Korea
⁵Department of Pediatrics, The Catholic University of Korea College of Medicine, Seoul St. Mary's Hospital, Seoul, Korea
⁶Department of Pediatrics, Yonsei University College of Medicine, Severance Children's Hospital, Seoul, Korea
⁷Department of Pediatrics, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul, Korea
⁸Department of Pediatrics, Eulji University School of Medicine, Eulji University Hospital, Seoul, Korea
⁹Department of Pediatrics, Hanyang University College of Medicine, Seoul, Korea
¹⁰Department of Pediatrics, Kyung Hee University School of Medicine, Seoul, Korea
¹¹Department of Pediatrics, Inha University College of Medicine, Incheon, Korea
¹²Department of Pediatrics, Hallym University College of Medicine, Anyang, Korea
¹³Department of Pediatrics, Inje University Ilsan Paik Hospital, Goyang, Korea
¹⁴Department of Pediatrics, Korea University Ansan Hospital, Ansan, Korea
¹⁵Department of Pediatrics, The Catholic University of Korea College of Medicine, St. Vincent's Hospital, Suwon, Korea
¹⁶Department of Pediatrics, CHA Bundang Medical Center, CHA University, Seongnam, Korea
¹⁷Department of Pediatrics, Yonsei University Wonju College of Medicine, Wonju, Korea
¹⁸Department of Pediatrics, Dankook University College of Medicine, Cheonan, Korea
¹⁹Department of Pediatrics, Keimyung University School of Medicine, Daegu, Korea
²⁰Department of Pediatrics, Pusan National University School of Medicine, Busan, Korea
²¹Department of Pediatrics, Kosin University Gospel Hospital, Busan, Korea
²²Department of Pediatrics, Fatima Hospital, Changwon, Korea
²³Department of Pediatrics, Chonbuk National University Medical School, Jeonju, Korea
²⁴Department of Pediatrics, Chonnam National University, Gwangju, Korea

ABSTRACT

Background: Invasive bacterial infections in apparently immunocompetent children were retrospectively analyzed to figure causative bacterial organisms in Korea.

Methods: A total of 947 cases from 25 university hospitals were identified from 2006 to 2010 as a continuance of a previous 10-year period study from 1996 to 2005.

Results: *Escherichia coli* (41.3%), *Streptococcus agalactiae* (27.7%), and *Staphylococcus aureus* (27.1%) were the most common pathogens in infants < 3 months of age. *S. agalactiae* was the most prevalent cause of meningitis and pneumonia and *E. coli* was the major cause of bacteremia without localizing signs in this group. In children 3 to 59 months of age, *Streptococcus*

Byung Wook Eun 
<https://orcid.org/0000-0003-3147-9061>
 Sung Hee Oh 
<https://orcid.org/0000-0001-6943-5430>
 Sung-Ho Cha 
<https://orcid.org/0000-0003-0540-5747>
 Young Jin Hong 
<https://orcid.org/0000-0002-0878-9515>
 Kwang Nam Kim 
<https://orcid.org/0000-0003-4024-5128>
 Nam Hee Kim 
<https://orcid.org/0000-0002-1793-1392>
 Yun-Kyung Kim 
<https://orcid.org/0000-0003-4396-8671>
 Jong-Hyun Kim 
<https://orcid.org/0000-0001-8641-7904>
 Taekjin Lee 
<https://orcid.org/0000-0002-8912-6982>
 Hwang Min Kim 
<https://orcid.org/0000-0002-0329-1371>
 Kun Song Lee 
<https://orcid.org/0000-0001-7318-2296>
 Chun Soo Kim 
<https://orcid.org/0000-0003-3888-334X>
 Su Eun Park 
<https://orcid.org/0000-0001-5860-821X>
 Young Mi Kim 
<https://orcid.org/0000-0002-6853-2228>
 Chi Eun Oh 
<https://orcid.org/0000-0002-0439-8170>
 Sang Hyuk Ma 
<https://orcid.org/0000-0002-0582-9286>
 Dae Sun Jo 
<https://orcid.org/0000-0002-3141-9539>
 Young Youn Choi 
<https://orcid.org/0000-0002-8631-8241>
 Hoan Jong Lee 
<https://orcid.org/0000-0001-9643-3692>

Funding

This study was supported by Seoul National University Hospital grants (No. 06-2011-3200), which were underwritten by Pfizer Pharmaceuticals Korea Ltd.

Disclosure

The authors have no potential conflicts of interest to disclose.

Author Contributions

Conceptualization: Rhie K, Choi EH, Cho EY, Lee HJ. Data curation: Lee J, Kang JH, Kim DS, Kim YJ, Ahn Y, Eun BW, Oh SH, Cha SH, Hong YJ, Kim KN, Kim NH, Kim YK, Kim JH, Lee T, Kim HM, Lee KS, Kim CS, Park SE, Kim YM, Oh CE, Ma SH, Jo DS, Choi YY. Writing - original draft: Rhie K. Writing - review & editing: Rhie K, Choi EH, Lee HJ.

pneumoniae (54.2%), *S. aureus* (20.5%), and *Salmonella* spp. (14.4%) were the most common pathogens. *S. pneumoniae* was the leading cause of pneumonia (86.0%), meningitis (65.0%), and bacteremia without localizing signs (49.0%) in this group. In children ≥ 5 years of age, *S. aureus* (62.8%) was the predominant pathogen, followed by *Salmonella* species (12.4%) and *S. pneumoniae* (11.5%). *Salmonella* species (43.0%) was the most common cause of bacteremia without localizing signs in this group. The relative proportion of *S. aureus* increased significantly over the 15-year period (1996–2010) in children ≥ 3 months of age ($P < 0.001$), while that of *Haemophilus influenzae* decreased significantly in both < 3 months of age group ($P = 0.036$) and ≥ 3 months of age groups ($P < 0.001$).

Conclusion: *S. agalactiae*, *E. coli*, *S. pneumoniae*, and *S. aureus* are common etiologic agents of invasive bacterial infections in Korean children.

Keywords: Bacterial Infections; Epidemiology; *Streptococcus agalactiae*; *Escherichia coli*; *Streptococcus pneumoniae*; *Staphylococcus aureus*

INTRODUCTION

Invasive bacterial infections are a major cause of morbidity and mortality in children. *Streptococcus pneumoniae*, *Haemophilus influenzae*, *Neisseria meningitidis*, *Streptococcus agalactiae*, *Streptococcus pyogenes*, *Staphylococcus aureus*, *Salmonella* species, *Listeria* species, and *Escherichia coli* are major bacterial pathogens that are responsible for invasive infections in immunocompetent children.^{1,2} However, the main organisms causing various clinical syndromes vary by age and over time. Thus, empirical antimicrobial therapy of suspected bacterial infections must be tailored to the common etiological agents of the infections in each region. In addition, countries employ immunization programs by prioritizing the burden of diseases.

To estimate the burden of invasive bacterial infections, many countries establish a nationwide surveillance system, such as Active Bacterial Core surveillance by the Centers for Disease Control and Prevention in the United States³ and Health Protection Agency surveillance in the United Kingdom.⁴ Japan and Taiwan have also nationwide surveillance systems for invasive bacterial infections.^{5,6} Although Korea launched a surveillance system for invasive bacterial infections that are preventable by the national immunization program such as diphtheria, pertussis, tetanus, and *H. influenzae* type b, this surveillance system did not include the most prevalent pathogens. Additionally, the notification rate remained low; 17.4%–43.0% for vaccine preventable diseases in the 2000s.^{7,8}

A retrospective study of invasive bacterial infections in Korean children at 18 university hospitals across the country was conducted over a ten-year period from 1996 to 2005.⁹ The study was the first multicenter, hospital-based surveillance study conducted in Korea. In the study, *S. pneumoniae* was found to be responsible for 44% (123 of 279 cases) of invasive bacterial diseases in immunocompetent children 3 to 59 months of age. Following this study, a remarkable change in immunization practices occurred to prevent invasive pneumococcal diseases (IPDs), which are the leading cause of bacterial infections in children. The pneumococcal conjugate vaccine 7 (PCV7) was introduced at the end of 2003, and the PCV7 was replaced by the PCV10 and PCV13 in 2010. This time period, after PCV7 use and before PCV13 introduction, provides data on the epidemiological changes of invasive bacterial infections influenced by PCV7 use.

Thus, we conducted a retrospective, multicenter study to analyze the relative proportion of the major bacterial agents responsible for invasive infections in apparently immunocompetent Korean children from 2006 to 2010 according to age, clinical features, and time. In addition, this study was conducted as a continuance of the previous multicenter surveillance study performed from 1996–2005,⁹ permitting a comparison of data from the two study periods.

METHODS

Participating hospitals

Twenty-five university-affiliated hospitals participated in this study: 16 hospitals are located in the national capital region (Seoul St. Mary's Hospital, Kyung Hee Medical Center, Seoul National University Children's Hospital, Severance Children's Hospital, Samsung Medical Center, Nowon Eulji Medical Center, Ewha Womans University Medical Center, Hanyang University Hospital, Inha University Hospital, Gachon University Gil Medical Center, Hallym University Medical Center, Inje University Ilsan Paik Hospital, Korea University Ansan Hospital, St. Vincent's Hospital, Seoul National University Bundang Hospital, and CHA Bundang Medical Center), and 9 hospitals are located in the regional central cities of provinces (Gangwon-do: Wonju Christian Hospital, Chungcheongnam-do: Dankook University Hospital, Jeollabuk-do: Chonbuk National University Hospital, Gwangju/Jeollanam-do: Chonnam National University Hospital, Daegu/Gyeongsangbuk-do: Keimyung University Dongsan Medical Center, and Busan/Gyeongsangnam-do: Pusan National University Hospital, Pusan National University Yangsan Hospital, Gosin University Gospel Hospital, and Changwon Fatima Hospital). The geographic distribution of the 25 university-affiliated hospitals that participated in this study is shown in Fig. 1.

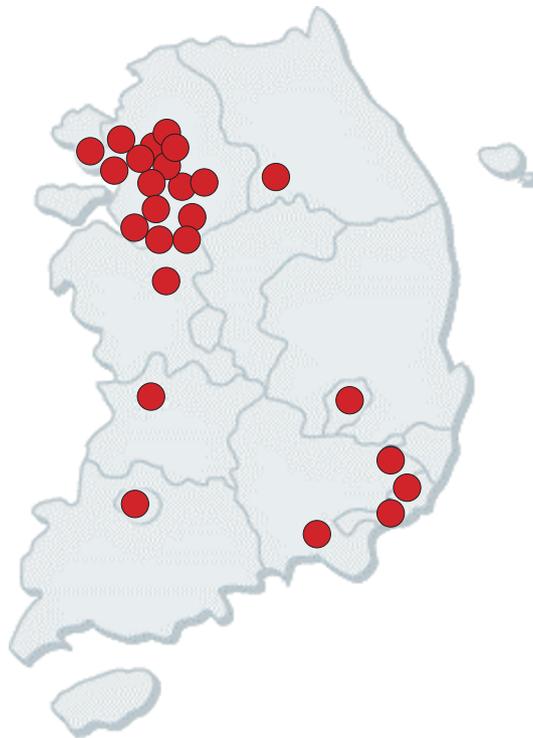


Fig. 1. Geographic distribution of the 25 university-affiliated hospitals that participated in this study.

Data collection

The survey period was from January 2006 to December 2010. The medical records of children less than 18 years of age and neonates and infants born after 37 weeks of gestation with invasive bacterial infections were reviewed retrospectively at the 25 university hospitals. Cases were selected from databases of hospital discharge records as well as from microbiology registries in departments of laboratory medicine. Initially, cases with positive results for the target organisms were selected. Investigators then reviewed the medical records of the cases and determined whether each case should be included in the study based on the case definition of the study protocol. A standard case report form was used to record the causative organisms, site(s) of bacterial isolation, clinical manifestations, laboratory findings, antibiotic susceptibility, and treatment results. The distributions of the bacterial pathogens were analyzed according to age, clinical diagnosis, and time.

Case definition

An 'invasive bacterial infection' was defined as the isolation of a bacterial organism from a normally sterile body fluid, such as blood, cerebrospinal fluid, pleural fluid, pericardial fluid, joint fluid, bone aspirate, or a deep tissue abscess.³ If the same pathogen was found in another sample taken within 30 days of the previous positive sample or during the same admission period, the episode was considered a single case. Cases with conditions that predisposed the patient to invasive bacterial infections, such as congenital or acquired immunodeficiency, steroid or cancer chemotherapy, prematurity, and infections associated with anatomic abnormalities (e.g., cerebrospinal fluid leakage), were excluded.¹⁰ Urinary tract infections without bacteremia were excluded unless a bacterial organism was isolated simultaneously from sterile body fluids other than urine, because culture results for specimens collected by the bag method, which was used in most cases, are not reliable. An invasive infection was classified on the basis of consistent clinical findings and the isolation of pathogens from the specimens. 'Bacteremia without localizing signs' was defined as cases in which no identifiable focus was identified, and the bacterium was isolated only from blood. A clinical diagnosis with a focal lesion such as 'meningitis,' 'pneumonia,' 'arthritis,' 'osteomyelitis,' 'peritonitis,' or 'deep tissue abscess' was defined if the bacterium was isolated from the corresponding body fluid, such as cerebrospinal fluid, pleural fluid, joint fluid, bone aspirate or abscess, or if the bacterial organism was isolated only from blood, but the clinical manifestations were compatible with the diagnosis.

Microbiological methods

The target organisms were limited to nine major pediatric bacterial pathogens: *S. pneumoniae*, *S. aureus*, *H. influenzae*, *N. meningitidis*, *S. agalactiae*, *S. pyogenes*, *Listeria monocytogenes*, *Salmonella* species, and *E. coli*. The current study added *E. coli* to the previous study's list of eight target organisms,⁹ and included infants younger than 3 months of age only for *E. coli*.

Clinical specimens were collected and processed using standard methods at each of the participating hospitals. Blood specimens were initially inoculated into brain-heart infusion broth and tetrathionate broth and processed using an automated culture system such as BACTEC (Becton Dickinson, Canada Inc., Mississauga, Canada) or BacT/Alert (Organon Teknika, Scarborough, Canada). The culture media was subcultured on chocolate agar and blood agar plates when turbidity was evident and, in some hospitals, before discarding after one week of incubation. Blood agar and chocolate agar plates were used for the direct plating of cerebral, pleural, or other body fluids. *Salmonella* isolates were serogrouped using serogroup-specific typing sera.

Clinical outcomes

The final outcomes were evaluated upon discharge and were categorized as follows: full recovery, recovery with sequelae, death or hopeless discharge. Cases classified as 'transferred to another hospital' and 'discharged against medical advice, not expected to die' were considered missing data because the final outcomes could not be determined. The case-fatality rate was calculated for each pathogen.

Statistical analysis

Age groups were stratified as follows: < 3 months of age, 3 to 23 months of age, 24 to 59 months of age, and ≥ 5 years of age. The temporal changes in the relative proportions of the nine bacterial organisms in infants < 3 months of age group and eight organisms (excluding *E. coli*) in infants and children ≥ 3 months of age groups were analyzed using the Cochran-Armitage test for trends. We performed statistical analyses using R version 3.3.3 (R foundation for Statistical Computing, Vienna, Austria), and considered $P < 0.05$ to be statistically significant.

Ethics statement

This study was approved by the Institutional Review Board at Seoul National University Hospital (No. H-1107-041-368), and the need for informed consent was waived.

RESULTS

Etiology

A total of 947 episodes of invasive bacterial infections and 950 bacterial organisms were identified during the study period. Co-infections were observed in 3 episodes. The male:female ratio was 0.6:1. Regardless of age and clinical diagnosis, *S. aureus* was the most frequently isolated organism (n = 317, 33.5%), followed by *E. coli* (n = 192, 20.3%), *S. pneumoniae* (n = 174, 18.4%), and *S. agalactiae* (n = 133, 14.4%). *Salmonella* species and *S. pyogenes* accounted for 67 cases (7.1%) and 25 cases (2.6%), respectively. There were very few cases of infection with *H. influenzae* (18 cases), *N. meningitidis* (14 cases), or *L. monocytogenes* (4 cases).

Of 67 *Salmonella* isolates, serogroup data were available for 62. Among these, 14 isolates were *Salmonella typhi* and 48 isolates were non-typhoidal *Salmonella* which included *Salmonella paratyphi* (n = 14), non-typhoidal group D (n = 9), serogroup A (n = 1), serogroup B (n = 4), serogroup C (n = 28), serogroup E (n = 1), and serogroup G (n = 1).

Distribution of causative bacterial organisms by age

The age-related distribution of causative bacterial organisms is shown in **Table 1**. The number of cases was highest in < 3 months of age group, with 465 cases (49%), followed by ≥ 5 years of age group with 218 cases (23%), the 24–59 months of age group with 134 cases (14%), and the 3–23 months of age group with 130 cases (14%). Patients < 5 years of age accounted for 76% of all cases. The distribution of bacteria differed by age group. *E. coli* was the most prevalent organism among infants < 3 months of age and accounted for 41.3% of the cases in this age group, followed by *S. agalactiae* and *S. aureus*, which accounted for 27.7% and 27.1% of cases, respectively. These three organisms were responsible for 96.1% of all the invasive bacterial infections in infants < 3 months of age.

S. pneumoniae, *S. aureus*, and *Salmonella* species were isolated frequently from infants and children ≥ 3 months of age. *S. pneumoniae* was the most common bacterial agent responsible

Table 1. Distribution of causative bacterial organisms among 947 invasive infections from 2006 to 2010, by age group

Identified species of bacteria	Age group, No. (%)					Total
	< 3 months	3–59 months			≥ 5 years	
		3–23 months	24–59 months	Subtotal		
<i>S. aureus</i>	126 (27.1)	30 (23.1)	24 (17.9)	54 (20.5)	137 (62.8)	317 (33.5)
<i>S. pneumoniae</i>	6 (1.3)	66 (50.8)	77 (57.5)	143 (54.2)	25 (11.5)	174 (18.4)
<i>S. agalactiae</i>	129 (27.7)	6 (4.6)	0 (0.0)	6 (2.3)	1 (0.5)	136 (14.4)
<i>Salmonella</i> spp.	2 (0.4)	17 (13.1)	21 (15.7)	38 (14.4)	27 (12.4)	67 (7.1)
<i>H. influenzae</i>	2 (0.4)	5 (3.8)	7 (5.2)	12 (4.5)	4 (1.8)	18 (1.9)
<i>S. pyogenes</i>	4 (0.9)	3 (2.3)	5 (3.7)	8 (3.0)	13 (6.0)	25 (2.6)
<i>N. meningitidis</i>	1 (0.2)	3 (2.3)	0 (0.0)	3 (1.1)	10 (4.6)	14 (1.5)
<i>L. monocytogenes</i>	3 (0.6)	0 (0.0)	0 (0.0)	0 (0.0)	1 (0.5)	4 (0.4)
<i>E. coli</i>	192 (41.3)	NA ^a	NA	NA	NA	192 (20.3)
Total	465	130	134	264	218	947

NA = not applicable.

^aNA denotes data were not collected.

for invasive infections in infants and children 3 to 59 months of age, accounting for 54.2% (143/264) of cases. In contrast, *S. pneumoniae* was the third most common cause of bacterial infections in children ≥ 5 years of age (11.5% of cases). *S. aureus* was a significant cause of infections in children 3 to 59 months of age, accounting for 20.4% of invasive infections that developed in this age group, and was the leading cause of bacterial infection in children ≥ 5 years of age (62.8% of cases). *Salmonella* species accounted for 13.5% of invasive infections in children ≥ 3 months of age.

Distribution of etiologic organisms according to clinical diagnosis

Among the 947 invasive bacterial infections identified, bacteremia without localizing signs was most common, accounting for 48.9% (n = 463) of cases, followed by arthritis and osteomyelitis (n = 163, 17.2%), meningitis (n = 141, 14.9%), and pneumonia with bacteremia or empyema (n = 113, 11.9%). Other diseases not assigned to one of the above clinical entities accounted for 7% of cases.

The distributions of the causative organisms according to three age groups (< 3 months of age, 3 to 59 months of age, and ≥ 5 years of age) for the 4 most common clinical diagnoses are illustrated in Fig. 2. For meningitis, most cases were attributed to *S. agalactiae* (69.0%) in patients < 3 months of age. In patients 3 to 59 months of age, *S. pneumoniae* (65.0%) and *H. influenzae* (23.0%) were frequently identified in meningitis cases. *N. meningitidis* (43.0%) and *S. pneumoniae* (39.0%) caused most of the meningitis cases in patients ≥ 5 years of age. For bacteremia without localizing signs, *E. coli* (51%), *S. aureus* (30.0%), and *S. agalactiae* (17.0%) were the predominant causes of infection among infants < 3 months of age. In contrast, *S. pneumoniae* was the leading cause of bacteremia without localizing signs in 3 to 59 months of age group (49.0%), followed by *Salmonella* species (25.0%) and *S. aureus* (18.0%). *Salmonella* species (43.0%) and *S. aureus* (30.0%) were common causes of bacteremia in children ≥ 5 years of age.

S. aureus was the most prevalent cause of bone and joint infections in all the age groups, causing 64.0% of cases in < 3 months of age group, 80.0% of cases in the 3–59 months of age group, and 92.0% of cases in ≥ 5 years of age group. For pneumonia, *S. agalactiae* predominated in < 3 months of age group. In the other age groups, however, *S. pneumoniae* was the most frequent cause of infection, responsible for 86.0% of cases in 3 to 59 months of age group and 44.0% of cases in ≥ 5 years of age group. *S. aureus* showed the same proportion as *S. pneumoniae* in ≥ 5 years of age group.

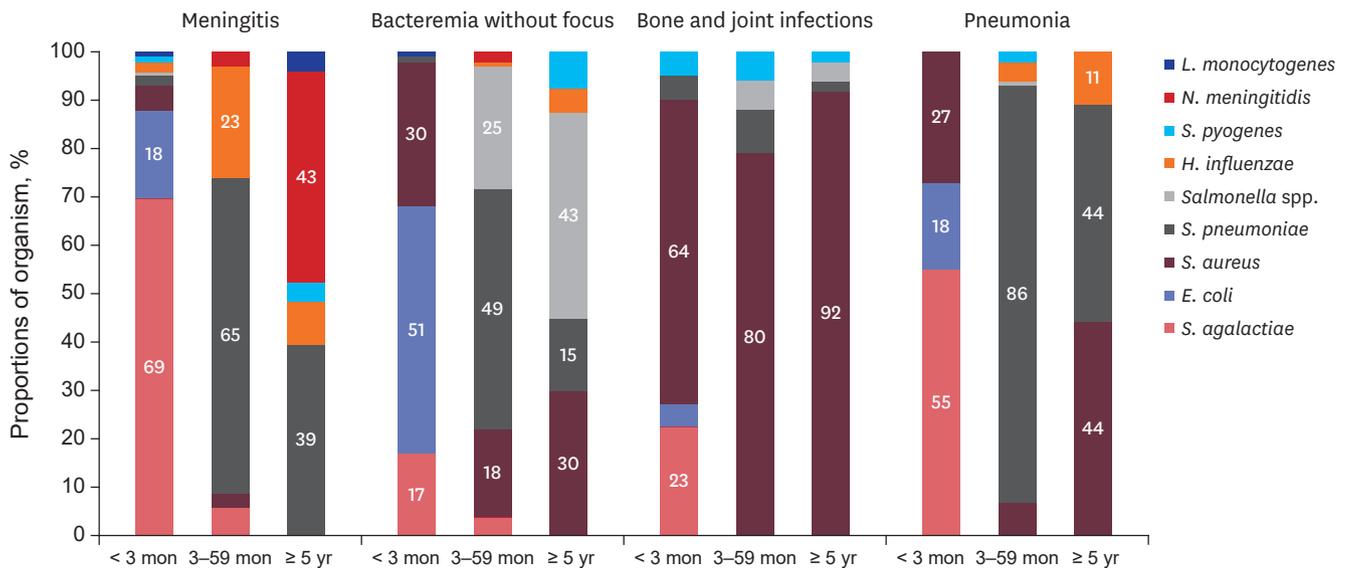


Fig. 2. Distribution of the causative bacterial organisms in invasive infections in immunocompetent Korean children between 2006 and 2010 according to age group and clinical spectrum.

Changes in the relative proportion of bacterial organisms over time

Data from the previous 10-year study of 1996–2005⁹ were reconstructed for comparison of the distributions of the bacterial organisms according to age group. Distributions of the relative proportions of the 8 bacteria that caused invasive infections in < 3 months of age group are shown in Table 2. For an exact comparison with the previous study period, *E. coli*, which was surveyed only in our study, was excluded and recalculated. The study periods were divided into 5-year increments: 1996–2000, 2001–2006, and 2006–2010. *S. aureus* and *S. agalactiae* were the two most prevalent organisms throughout the collective study period. When we performed the trend comparisons between the three five-year periods, no significant changes in the organismal trends were observed, except for *H. influenzae*. Over 15 years, there was a decreasing trend in the relative proportion of *H. influenzae* < 3 months of age group ($P = 0.036$).

The temporal changes in the relative proportions of the 8 most prevalent bacterial organisms in children ≥ 3 months of age between 1996 and 2010 were analyzed (Fig. 3). There was a significantly increasing trend of *S. aureus* infections over these 15 years ($P < 0.001$). In 1996, the relative proportion of *S. aureus* infections was 13.8%, which rose to 42.2% in 2006 and 38.6%

Table 2. Distribution in the relative proportions of the 8 causative organisms of invasive infections

Species of bacteria ^a	Period, No. (%)			P-value
	1996–2000	2001–2005	2006–2010	
<i>S. aureus</i>	35 (34.0)	71 (39.0)	126 (46.2)	0.078
<i>S. pneumoniae</i>	2 (1.9)	9 (4.9)	6 (2.2)	0.901
<i>S. agalactiae</i>	49 (47.6)	88 (48.4)	129 (47.3)	0.966
<i>Salmonella</i> spp.	4 (3.9)	1 (0.5)	2 (0.7)	0.080
<i>H. influenzae</i>	6 (5.8)	5 (2.7)	2 (0.7)	0.036
<i>S. pyogenes</i>	3 (2.9)	7 (3.8)	4 (1.5)	0.544
<i>N. meningitidis</i>	2 (1.9)	0 (0.0)	1 (0.4)	0.224
<i>L. monocytogenes</i>	2 (1.9)	1 (0.5)	3 (1.1)	0.598
Total	103	182	273	

Under 3 months of age and trend comparisons between three periods, 1996–2000, 2001–2005, and 2006–2010.

^a*E. coli* of 2006–2010 period was excluded from this analysis for the comparison with previous study periods.

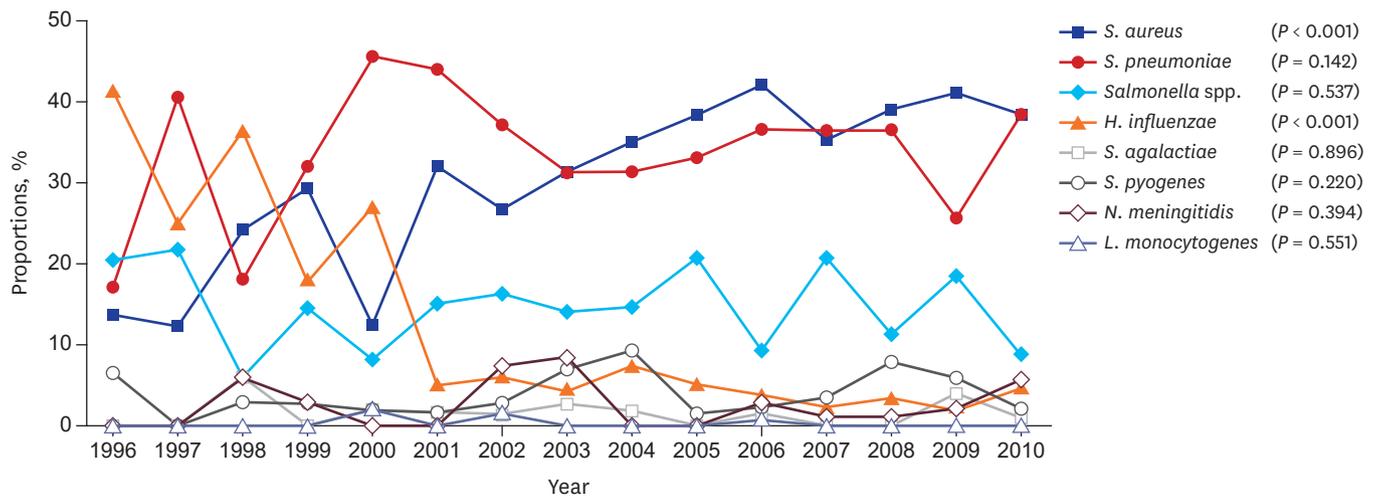


Fig. 3. Temporal changes in the relative proportions of the 8 most important bacterial organisms of invasive infections in immunocompetent Korean children 3 months of age or older between 1996 and 2010.

in 2010. *H. influenzae* infections continued to show a significantly decreasing trend ($P < 0.001$). The relative proportion of *H. influenzae* infections in 1996 was 41.4%, and in 2010, it was 4.5%. No significant changes in the relative proportions of infections with *S. pneumoniae* and *Salmonella* species were observed.

Clinical outcomes

Clinical outcomes were available for 877 (92%) of the 947 cases; 70 cases (8%) had missing data due to 'transferred to another hospital' or 'discharged against medical advice.' The overall case-fatality rate was 2.5% (22/877). When the data were analyzed according to clinical diagnosis, meningitis had the highest case-fatality rate (10.2%, 13/127), followed by bacteremia without localizing signs (2.0%, 6/301) and pneumonia (1.9%, 2/105). When the data were analyzed according to age groups, the 3–23 months of age group had the highest case-fatality rate (8.6%, 10/116), followed by the 24–59 months of age group (2.4%, 3/124), the infants < 3 months of age group (18%, 8/437), and ≥ 5 years of age group (0.5%, 1/200).

Among the bacteria studied, *N. meningitidis* was associated with the highest case-fatality rate of 9.1% (1/11); however, the overall number of *N. meningitidis* infections was small. When bacteria that were isolated in 50 or more cases were analyzed, we found that *S. pneumoniae* had the highest case-fatality rate of 8.4% (13/155), followed by *S. agalactiae* with a case fatality rate of 4.8% (6/126). *S. pyogenes* and *E. coli* had case fatality rates of 4.2% (1/24) and 0.5% (1/185), respectively. No fatalities associated with *H. influenzae* infection were identified during the study period.

In terms of the frequency of sequelae associated with meningitis cases, *H. influenzae* had a very high sequelae rate of 50% (4/8). Sequelae rates associated with meningitis caused by other common infective organisms, namely, *S. pneumoniae*, *S. agalactiae*, and *E. coli*, showed similar trends compared to their case-fatality rates and were 36.8% (7/19), 30.8% (16/52), and 13.3% (2/15), respectively. The most common sequelae were hydrocephalus, subdural effusion, and hearing loss. Long-term sequelae such as developmental delay and cognitive impairment were not evaluated due to the retrospective design of the study.

DISCUSSION

The purpose of this study was to investigate the etiology of invasive bacterial infections in a pediatric population in Korea through a retrospective review of multicenter hospital-based surveillance. This study is important not only for its purpose but also for its continuous maintenance of multicenter-based surveillance since 1996. The study period between 2006 and 2010 corresponds to the period after PCV7 use and before PCV13 introduction in Korea. Data collected during this period reflect the epidemiological changes in invasive bacterial infections that were influenced by PCV7 use. Additionally, the number of cases ($n = 947$) and bacterial organisms ($n = 950$) included in this study comprise the largest invasive infection study in Korea to date. Nationwide or regional data collected during a certain time period can be used to reliably analyze the major pathogens responsible for invasive bacterial infections. Twenty eight percent (25/89) of university hospitals in Korea participated in this study design, and they were distributed nationwide. The data in the current study will be useful for planning the empirical treatment of invasive bacterial infections as well as for establishing a national vaccine policy.

Some notable findings in this study are as follows. First, an expanded number of hospitals and clinical episodes were included in the current study compared to the 1996–2005 period: 18 hospitals and 768 episodes comprised the 1996–2005 study, while 25 hospitals and 947 episodes were included in the 2006–2010 period study. Second, we added our data to those of the previous 10 years. Thus, we were able to compare the relative proportions and trends of the 8 bacteria in both < 3 months of age group and ≥ 3 months of age groups over 15 years, from 1996 to 2010. During this period, *S. aureus* showed a significantly increasing trend in children ≥ 3 months of age. Third, *H. influenzae* continued to show a decreasing trend in both < 3 months of age group and ≥ 3 months of age groups.

In our study, the major pathogens that caused invasive bacterial infections differed according to age: for patients < 3 months of age, *E. coli*, *S. agalactiae*, and *S. aureus* predominated; for patients 3 to 59 months of age, *S. pneumoniae*, *S. aureus*, and *Salmonella* species were most prevalent; and for patients ≥ 5 years of age, *S. aureus*, *Salmonella* species, and *S. pneumoniae* dominated. In neonates and infants < 3 months of age, *S. agalactiae* was a major cause of meningitis, and *E. coli* was a major cause of bacteremia. In children ≥ 3 months of age, *S. pneumoniae* caused pneumonia, bacteremia, and meningitis. *S. aureus* was the main cause of bone and joint infections and bacteremia in this age group.

S. aureus has a diverse clinical spectrum that ranges from asymptomatic colonization, to skin and soft tissue infection, and to bone and joint infections. *S. aureus* is a major neonatal pathogen in Korea.¹¹ Additionally, *S. aureus* is becoming an increasingly frequent cause of community-acquired pneumonia in children < 5 years of age.¹² Up to 80% of bone and joint infections in children are caused by *S. aureus*. Generally, the relative proportions of the causative organisms in the different age groups and clinical diagnoses in our study were consistent with earlier studies. One notable finding was the antimicrobial sensitivity of the *S. aureus* isolates. We found that 38.7% of all the *S. aureus* isolates were oxacillin resistant, which is a higher proportion than those reported by previous studies.¹³

S. agalactiae remains one of major pathogens responsible for neonatal invasive infections in developed countries.^{14,15} In Korea, neonatal infections caused by *S. agalactiae* have increased in recent years.¹⁶ In our findings, the relative proportion of invasive infections in infants

< 3 months of age is consistent with previous studies. In a study by Lee et al.,² *S. agalactiae* accounted for 38% of the invasive infections in infants < 3 months of age. In the same age group, *S. agalactiae* accounted for 48.1% of the invasive infections in a previous study conducted from 1996 to 2005.⁹ In this study, from 2006 to 2010, *S. agalactiae* was the most prevalent organism, accounting for 47.3% of the infections. The exclusion of *E. coli* to allow for comparison with the previous study yielded this result. The previous 1996–2005 period study did not include *E. coli*; this study showed that *E. coli* was the most common pathogen to cause invasive bacterial infections in infants < 3 months of age. *E. coli* mainly caused bacteremia without localizing signs, or meningitis in this age group. Siitonen et al.¹⁷ also reported that the common infections caused by *E. coli* were bacteremia, pyelonephritis, and meningitis in a Finnish study performed during the late 1980s. The case-fatality rate caused by *E. coli* during the neonatal period was 9%. In the current study, the case-fatality rate of *E. coli* infection was 0.5% (1 of 185 identifiable cases). Although the case fatality rate was relatively low compared to the other organisms, *E. coli* is still a major invasive pathogen in neonates and infants < 3 months of age in terms of the relative proportion.

S. pneumoniae was the most common pathogen to cause invasive bacterial infections among children 3 to 59 months of age, accounting for 45.3% of infections from 1996–2006 in Korea.⁹ In our study, 54.4% of the invasive infections were caused by *S. pneumoniae*, which was the most prevalent pathogen of bacteremia without localizing signs, meningitis, and pneumonia in the same age group. The reduction of invasive infections by *H. influenzae* and *Salmonella* species resulted in this high proportion of *S. pneumoniae* in children 3 to 59 months of age. *S. pneumoniae* also had the highest case-fatality rate regardless of age in the current study. After the introduction of PCV7 in 2000 and of PCV13 in 2010 in the United States, the overall incidence of IPD declined from 100 cases per 100,000 children under 5 years of age in 1998 to 9 cases per 100,000 in 2015. IPD caused by PCV13 serotypes declined from 91 cases per 100,000 children under 5 years of age in 1998 to 2 cases per 100,000 in 2015.¹⁸ In Korea, PCV7 was offered as an optional vaccine in 2003, and vaccine coverage reached up to 70% in 2009. The proportion of IPD in children 3 to 59 months of age in this study period, 2006–2010, was 54.2%, and it was 39.6% in the 1996–2000 period and 45.7% in the 2001–2005 period. This increase in the relative proportion of IPD might have simply gained from reduction of other bacterial organisms. Another partial explanation given is that due to low PCV7 uptake rate during the first half of this study period and changes in the serotype distribution among IPD cases, a decrease in the proportion of IPD was not observed in this study period. Although PCV7 was introduced at the end of 2003, the vaccination rate was low for several years and the vaccine uptake rate of the primary 3 dose series reached around 60% in 2007.¹⁹ According to data from a study performed during the same time period, 2006–2010, the proportion of PCV7 serotypes decreased from 62.5% in 2006 to 21.4% in 2010, whereas the proportion of PCV13-specific serotypes, especially 19A, increased from 17.4% in 2006 to 47.8% in 2010.²⁰ Therefore, the proportion of PCV7 serotypes decreased while PCV13-specific serotypes increased during the 2006–2010 period, reducing the impact of PCV7 and making *S. pneumoniae* still one of the most important pathogens of invasive infections. Analysis of the relative proportion of IPD after the 2006–2010 period will be available in a follow-up study that will include the period after 2010, when the PCV10 and PCV13 were introduced, as well as the period after 2014, when PCVs were implemented in the national immunization program in Korea.

Before the introduction of conjugate vaccines, *H. influenzae* was the major pathogen that caused meningitis, pneumonia, epiglottitis, arthritis, and cellulitis worldwide.^{21,22} *H. influenzae*

infection has transitioned, however, from a major cause of childhood illness to a rare disease in every country where conjugate vaccines have been introduced into the routine immunization schedule.²³ In Korea, an *H. influenzae* vaccine was introduced in 1996 and was implemented in the routine immunization schedule in 2013. In a study performed in the pre-vaccination era, Nam and Lee¹ reported that *H. influenzae* was a major pathogen, accounting for 14% of all bacterial infections, 88% of which were reported to be meningitis. In a previous study by Lee et al.,⁹ *H. influenzae* infections accounted for 20.1% of all invasive infections and 54.2% of meningitis infections in children ≥ 3 months of age during the first half of the study period, from 1996 to 2000. However, this proportion dropped to 4.5% of all invasive infections and 19.7% of all meningitis infections in the latter half of the study period, from 2001 to 2005. In our study, we observed a significant decline in all of the age groups over the 15-year period, from 1996 to 2010.

Salmonella species are water- and food-borne pathogens in low-resource countries. The risk for infection is high in countries with poor sanitation and lack of access to safe food and water. Meanwhile, invasive *Salmonella* infections have declined in countries where the treatment of municipal water, pasteurization of dairy products, and improvement of public sanitation have been introduced.²⁴ In the current and previous studies, *Salmonella* infections have constantly declined: 23% in the 1986–1995 period, 10% in the 1996–2005 period, and 7% in the 2006–2010 period.^{1,9} However, 13.5% of the infections in children ≥ 3 months of age were attributed to *Salmonella* species. In the past, most cases of salmonellosis were associated with *S. typhi*; in the present, with the increase in the consumption of meat, infections due to nontyphoidal *Salmonella* occur consistently.²⁵ In our study, *Salmonella* species were most often isolated from cases of bacteremia without localizing signs in ≥ 3 months of age groups, which was consistent with the findings of the prior study.

This study had several limitations. We targeted only nine bacterial organisms. However, by focusing on these nine bacterial organisms, most of the invasive bacterial infections in the immunocompetent children were likely included in our analysis. These nine bacterial organisms are the most important pathogens that cause infections in children who do not have any underlying diseases or conditions related to an immune-compromised state. Different numbers of hospitals were included in the current and present study. This may have affected the result of trend analysis in making comparisons across study periods. Hospitals included in the 15-year study period, however, are university-affiliated hospitals and supposed to have similar characteristics. The retrospective design of the study may have resulted in bias. Cases of mild illness such as occult bacteremia, where the attending doctor might not perform cultures for out-patients, were excluded from this study and thus may have introduced a selection bias. Variations in laboratory protocols for the detection of bacteria in the participating hospitals could also have affected our results. For a more acute description of the distribution and etiology of invasive bacterial infections in Korea, further well-designed prospective studies are needed. This study, however, collected invasive bacterial infection cases steadily over 15 years in the absence of nationwide surveillance. The data in this study can provide epidemiological data of invasive bacterial infections in Korean children and serve as a means to judge the effects of vaccine policies.

In conclusion, the distributions of bacteria that cause invasive infections differed by age group and clinical diagnosis. In children ≥ 3 months of age, *S. pneumoniae* remained an important pathogen due to its incidence and fatality rate. The relative proportion of *S. aureus* increased significantly over the 15-year study period in children ≥ 3 months of age. Over the same

time period, the relative proportion of invasive infections caused by *H. influenzae* showed a decreasing trend in both < 3 months of age group and \geq 3 months of age groups. The national immunization policy may lead to changes in the etiology of invasive bacterial infections. Additionally, epidemics of certain bacterial infections may emerge unexpectedly. Thus, multicenter surveillance studies similar to ours should be continuously performed in the future.

REFERENCES

- Nam SG, Lee HJ. Etiology of invasive bacterial infections in apparently healthy children. *Korean J Infect Dis* 1998;30(3):227-34.
- Lee JH, Song EK, Lee JA, Kim NH, Kim DH, Park KW, et al. Clinical entities and etiology of invasive bacterial infections in apparently healthy children. *Korean J Pediatr* 2005;48(11):1193-200.
- Schuchat A, Hilger T, Zell E, Farley MM, Reingold A, Harrison L, et al. Active bacterial core surveillance of the emerging infections program network. *Emerg Infect Dis* 2001;7(1):92-9.
[PUBMED](#) | [CROSSREF](#)
- Zakikhany K, Degail MA, Lamagni T, Waight P, Guy R, Zhao H, et al. Increase in invasive *Streptococcus pyogenes* and *Streptococcus pneumoniae* infections in England, December 2010 to January 2011. *Euro Surveill* 2011;16(5):19785.
[PUBMED](#)
- Infectious agents surveillance report. <http://www.niid.go.jp/niid/en/iasr-e.html>. Updated 2017. Accessed April 30, 2017.
- Centers for Disease Control, R.O.C (Taiwan). <http://www.cdc.gov.tw/rwd/english>. Updated 2017. Accessed April 30, 2017.
- Kim YS, Park KH, Yoo HS, Kwon JW, Shin E. Reporting of national notifiable infectious diseases (NNIDs) and related characteristics. *Korean J Epidemiol* 2007;29(2):200-10.
- Shin E. *Evaluation and Improvement Strategy for Communicable Diseases Surveillance System*. Seoul, Korea: Korea Centers for Diseases Control and Prevention; 2003.
- Lee JH, Cho HK, Kim KH, Kim CH, Kim DS, Kim KN, et al. Etiology of invasive bacterial infections in immunocompetent children in Korea (1996–2005): a retrospective multicenter study. *J Korean Med Sci* 2011;26(2):174-83.
[PUBMED](#) | [CROSSREF](#)
- Saarinen M, Takala AK, Koskenniemi E, Kela E, Rönnerberg PR, Pekkanen E, et al. Spectrum of 2,836 cases of invasive bacterial or fungal infections in children: results of prospective nationwide five-year surveillance in Finland. *Clin Infect Dis* 1995;21(5):1134-44.
[PUBMED](#) | [CROSSREF](#)
- Kim KA, Shin SM, Choi JH. A nationwide survey on the causative organisms of neonatal sepsis in Korea. *J Korean Pediatr Soc* 2002;45(1):55-63.
- Schwartz KL, Nourse C. Pantón-Valentine leukocidin-associated *Staphylococcus aureus* necrotizing pneumonia in infants: a report of four cases and review of the literature. *Eur J Pediatr* 2012;171(4):711-7.
[PUBMED](#) | [CROSSREF](#)
- Choe YJ, Lee SY, Sung JY, Yang MA, Lee JH, Oh CE, et al. A review of *Staphylococcus aureus* infections in children with an emphasis on community-associated methicillin-resistant *S. aureus* infections. *Korean J Pediatr Infect Dis* 2009;16(2):150-61.
- Weston EJ, Pondo T, Lewis MM, Martell-Cleary P, Morin C, Jewell B, et al. The burden of invasive early-onset neonatal sepsis in the United States, 2005–2008. *Pediatr Infect Dis J* 2011;30(11):937-41.
[PUBMED](#) | [CROSSREF](#)
- Daley AJ, Isaacs DAustralasian Study Group for Neonatal Infections. Ten-year study on the effect of intrapartum antibiotic prophylaxis on early onset group B streptococcal and *Escherichia coli* neonatal sepsis in Australasia. *Pediatr Infect Dis J* 2004;23(7):630-4.
[PUBMED](#) | [CROSSREF](#)
- Park SE. Epidemiology of early and late onset neonatal sepsis. *J Korean Soc Neonatol* 2012;19(3):115-20.
[CROSSREF](#)
- Siitonen A, Takala A, Ratiner YA, Pere A, Mäkelä PH. Invasive *Escherichia coli* infections in children: bacterial characteristics in different age groups and clinical entities. *Pediatr Infect Dis J* 1993;12(7):606-12.
[PUBMED](#) | [CROSSREF](#)

18. Active bacterial core surveillance report: pneumococcal disease. <https://www.cdc.gov/pneumococcal/surveillance.html#abcs>. Updated 2017. Accessed April 10, 2017.
19. Choe YJ, Yang JJ, Park SK, Choi EH, Lee HJ. Comparative estimation of coverage between national immunization program vaccines and non-NIP vaccines in Korea. *J Korean Med Sci* 2013;28(9):1283-8.
[PUBMED](#) | [CROSSREF](#)
20. Cho EY, Lee H, Choi EH, Kim YJ, Eun BW, Cho YK, et al. Serotype distribution and antibiotic resistance of *Streptococcus pneumoniae* isolated from invasive infections after optional use of the 7-valent conjugate vaccine in Korea, 2006–2010. *Diagn Microbiol Infect Dis* 2014;78(4):481-6.
[PUBMED](#) | [CROSSREF](#)
21. Shetty S, Cohen AL, Edmond K, Ojo L, Loo J, O'Loughlin R, et al. A systematic review and critical evaluation of invasive *Haemophilus influenzae* type B disease burden studies in Asia from the last decade: lessons learned for invasive bacterial disease surveillance. *Pediatr Infect Dis J* 2010;29(7):653-61.
[PUBMED](#) | [CROSSREF](#)
22. Watt JP, Wolfson LJ, O'Brien KL, Henkle E, Deloria-Knoll M, McCall N, et al. Burden of disease caused by *Haemophilus influenzae* type B in children younger than 5 years: global estimates. *Lancet* 2009;374(9693):903-11.
[PUBMED](#) | [CROSSREF](#)
23. Morris SK, Moss WJ, Halsey N. *Haemophilus influenzae* type B conjugate vaccine use and effectiveness. *Lancet Infect Dis* 2008;8(7):435-43.
[PUBMED](#) | [CROSSREF](#)
24. Crump JA, Sjölund-Karlsson M, Gordon MA, Parry CM. Epidemiology, clinical presentation, laboratory diagnosis, antimicrobial resistance, and antimicrobial management of invasive *Salmonella* infections. *Clin Microbiol Rev* 2015;28(4):901-37.
[PUBMED](#) | [CROSSREF](#)
25. Antunes P, Mourão J, Campos J, Peixe L. Salmonellosis: the role of poultry meat. *Clin Microbiol Infect* 2016;22(2):110-21.
[PUBMED](#) | [CROSSREF](#)