**Results:** From September 2014 to September 2018, PCV13 vaccine coverage among persons ≥65 years old increased from <1% to 77%. During the same period, there was a total of 245 IPD cases. For a variety of reasons, we did not have serotype results for 57 (23%) IPD cases, which were excluded from the analysis. There were 61 (25%) PCV13-type IPD cases included in the analysis, of which 33 (14%) were serotype 3, PCV13 VE against PCV13-type serotypes was 68.0% (95% CI: 37.7%, 83.6%; P-value < 0.01), and 53.4% (95% CI: −10.0%, 80.3%; P = 0.08) against serotype 3.

**Conclusion:** During the first 4 years of PCV13 vaccination implementation in adults ≥65 years of age at KPNC, PCV13 provided significant protection against PCV13-type IPD. Further surveillance will allow for more precise estimation of PCV13 VE on overall and serotype 3 IPD over time.

**Disclosures:** All authors: No reported disclosures.

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**2712. Safety and Immunogenicity of two Doses of ExPEC4V Vaccine Against Extraintestinal Pathogenic Escherichia coli Disease in Healthy Adult Participants**

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**Session:** 277. Vaccines: Bacterial

**Saturday, October 5, 2019: 12:15 PM**

**Background:** The ExPEC4V vaccine contains 4 Escherichia coli O-antigens (O1A, O2, O6A, O25B) conjugated to exotoxin protein A and is being studied for prevention of Invasive Extraintestinal pathogenic E. coli (ExPEC) Disease (IED). This phase-2 double-blind study assessed safety and immunogenicity of ExPEC4V Clinical Trial Material (CTM), manufactured via a redesigned process (optimized O1A strain).

**Methods:** Participants (218 years) in stable health were randomized (3:1) to receive ExPEC4V dose 4:4:4:8 μg PS/serotype or placebo on Day 1 and second vaccination on Day 181 (6 months after first vaccination). Participants will be followed for safety until end of study at Day 360. Reactogenicity and immunogenicity (by ELISA, opsonophagocytic killing (OPA) assays) were evaluated pre-vaccination, and 15 days after first and second vaccinations (Day 185).

**Results:** Of 100 participants randomized (mean age 56, 48% males) and vaccinated (ExPEC4V, n = 75; placebo, n = 25), 97 completed Day 30. Solicited local AEs were higher for ExPEC4V (38.7%) than placebo (20%); most frequent was fatigue (32% vs. 12%). No serious or grade 3 solicited local AEs were reported. One participant in ExPEC4V experienced a grade 3 solicited systemic fatigue considered vaccine-related by investigator. ExPEC4V demonstrated immune responses against all serotypes at Day 15. Geometric mean titer effective concentration rank by most frequent was fatigue (32% vs. 12%). No serious or grade 3 solicited local AEs were higher in ExPEC4V (49.3%) than placebo (20%); and second vaccinations (Day 195).

**Conclusion:** ExPEC4V elicited robust and functional immune responses across all serotypes and was well tolerated with no vaccine safety findings. This study supports the development of futuremultivalent ExPEC vaccine to prevent IED.

**Disclosures:** All authors: No reported disclosures.

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**2713. Effectiveness of 23-Valent Pneumococcal Polysaccharide Vaccine in Korean National Population Cohort over 65 Years Old**

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**Background:** Twenty-three valent pneumococcal polysaccharide vaccine (PPSV) has been introduced to the National Immunization Program (NIP) for adults aged 65 years and older in Korea since 2013. We describe the effectiveness evaluation of PPSV among adults against pneumococcal infections including all-cause pneumonia (ACP), pneumococcal pneumonia (PP), and invasive pneumococcal diseases (IPD) using national population cohort.

**Methods:** Vaccination records of the national population, aged 65 years and older, from NIP registry by Korea Centers for Disease Control and Prevention (KCDC) were matched to their corresponding medical records by National Health Insurance Service (NHIS) for retrospective cohort analysis. Adults vaccinated with 1-dose PPSV between 2011 and 2016 were compared with those non-vaccinated. Primary outcomes were hospitalization due to ACP, PP, and IPD. Vaccine effectiveness (VE) adjusted for high risk and underlying conditions was calculated as one minus hazard rate ratio (HR) using Cox regression.

**Results:** Records of 6,743,002 cohort members were included. Forty-three percent were male, and median age was 75 years. Among the cohort, 3,425,949 (51%) were vaccinated during the study period. Incidence (per 100,000 person-years) of each disease in vaccinated and unvaccinated, respectively, was 2,184 and 1,584 for ACP; 8.9 and 5.4 for PP, and 1.6 and 1.9 for IPD. VE against IPD was 41.7% (95% CI 28.8–52.3) and against IPD sepsis was 55.3% (95% CI 39.8–64.0). PPSV was also protective against ACP with VE 7.2% (95% CI 6.6–7.8). When stratified by age-group, adults aged 65–74 years were better protected from ACP (VE 16.5% [95% CI 15.6–17.3]) compared with older adults aged 75 years or older (VE 0.4% [95% CI 1.2–0.4]), while VE was higher in older adults against IPD (VE 47.6% [95% CI 32.4–59.4]) and IPD sepsis (VE 54.9 [95% CI 38.4–66.9]) than in 65–74 years group (IPD VE 30.4% [95% CI 3.4–49.9]; sepsis VE 49.0% [95% CI 19.7–67.6]).

**Conclusion:** Single-dose PPSV strategy for adults in general population is protective against PCV, IPD, and IPD sepsis.

**Disclosures:** All authors: No reported disclosures.