

COST-EFFECTIVENESS OF THE 20-VALENT PNEUMOCOCCAL CONJUGATE VACCINE VS 15-VALENT PNEUMOCOCCAL CONJUGATE VACCINE FOR ADULTS IN ARGENTINA

Mercedes Mac Mullen¹, Abril Seyahian¹, Carolina Carballo¹
1. Pfizer Argentina

BACKGROUND

Streptococcus pneumoniae (*S. pneumoniae*) is a leading cause of morbidity and mortality among adults at a global level. The clinical spectrum of disease caused by pathogen this ranges from pneumonia to invasive pneumococcal disease (IPD) which includes bacteremia and meningitis.⁽¹⁻³⁾ Since 2017 the Ministry of Health (MoH) of Argentina recommends a sequential 13-valent pneumococcal conjugate vaccine (PCV13) - pneumococcal polysaccharide vaccine (PPSV23) regimen for adults over 65 years old and individuals up to 64 years who are at risk of severe bacterial infections.^(4, 5) Studies carried out in Argentina have determined that the PCV13-PPSV23 sequential regimen was cost-saving in the elderly population.⁽⁴⁾ With two higher value pneumococcal polysaccharide vaccines (PCVs) being developed: PCV15 and PCV20, which cover 2 and 7 additional serotypes respectively, this study aims to assess the cost-effectiveness (C-E) of these new vaccines in Argentina.

OBJECTIVE

To assess the cost-effectiveness of a single dose of PCV20 versus sequential administration PCV15 followed by PPV23 in the adult population of Argentina.

METHODS

Model structure

- A Markov-type model was used to estimate the clinical and economic impact shown on Table 1. Costs were adjusted by risk profile based on Weycker et al. of disease associated with *S. pneumoniae* infection, as well as the impact of (2016). A weighted average was then calculated based on the distribution of vaccinating the adult population, with each of the proposed strategies from the population in the different subsectors. Based on data provided by Pfizer Healthcare System perspective.
- The model uses a deterministic framework to depict lifetime risks and costs PAMI, 37,1% used the public subsector and 4,7% used the private subsector. Costs are expressed on 2021 US-Dollars.
- Utility reduction for IPD and inpatient NBP was 0.13 (annual) and reduction for outpatient NBP was 0.004 (annual).^(11,12)
- Vaccine coverage for population over 65 years was assumed to be invariant across age and risk groups and to be 65%. For population between 18 and 64 we assumed 17% of the high-risk population was vaccinated. These assumptions were made based on data provided by Pfizer.
- Costs and future benefits were discounted 3% annually.

Model parameters

- Vaccine effectiveness for PCVs against IPD and non bacteriemic pneumonia (NBP) caused by vaccine serotypes and risk adjustment were taken from published literature.⁽⁶⁻⁹⁾ PCVs were assumed to be durable for 5 years and to wane to 0% by year 16.⁽⁷⁾
- Vaccine effectiveness for PPSV23 against IPD caused by vaccine serotypes was taken from published literature.⁽¹⁰⁾ Effectiveness against NBP was assumed to be 0%. PPSV23 was assumed to wane to 0% by year 10.⁽¹⁰⁾
- Costs associated with the treatment of bacteremia, meningitis, inpatient

NBP, outpatient NBP and vaccine administration costs for each Healthcare system subsectors were estimated using a micro costing approach and are shown on Table 1. Costs were adjusted by risk profile based on Weycker et al. (2016). A weighted average was then calculated based on the distribution of the population in the different subsectors. Based on data provided by Pfizer Healthcare System perspective. PAMI, 37,1% used the public subsector and 4,7% used the private subsector. Costs are expressed on 2021 US-Dollars. Utility reduction for IPD and inpatient NBP was 0.13 (annual) and reduction for outpatient NBP was 0.004 (annual).^(11,12) Vaccine coverage for population over 65 years was assumed to be invariant across age and risk groups and to be 65%. For population between 18 and 64 we assumed 17% of the high-risk population was vaccinated. These assumptions were made based on data provided by Pfizer. Costs and future benefits were discounted 3% annually.

Scenario analysis

- Scenario analyses were conducted using alternative input values for key parameters in one way sensitivity analysis and probabilistic sensitivity analysis.

| Table 1: Base case model input values by age risk and group | | | | | | | | | | | | | |
|-------------------------------------------------------------|-------------|----------|----------|-------------|----------|----------|-------------|----------|----------|-------------|----------|----------|-----------|
| | 18-49 years | | | 50-64 years | | | 65-74 years | | | 75-84 years | | | ≥85 years |
| | Low | Mod | High | Low | Mod | High | Low | Mod | High | Low | Mod | High | Low |
| Number of adults (100K) ¹⁴ | 161.7 | 42.5 | 6.5 | 37.7 | 23.8 | 3.3 | 14.1 | 13.2 | 3.8 | 7.0 | 6.1 | 4.1 | 2.5 |
| Incidence of bacteremia (per 100K) ¹⁵ | 6.4 | 25 | 50.1 | 23.7 | 93.8 | 145.4 | 64.6 | 203 | 275.2 | 79.1 | 189.7 | 209.4 | 109.2 |
| Incidence of meningitis (per 100K) ^{15,16} | 0.02 | 0.08 | 0.17 | 0.05 | 0.19 | 0.29 | 0.07 | 0.23 | 0.31 | 0.11 | 0.25 | 0.28 | 0.15 |
| Inpatient NBP ^{17,18} | 60 | 355 | 1,204 | 243 | 1,183 | 3,382 | 513 | 2,527 | 5,704 | 830 | 2,382 | 3,463 | 806 |
| Outpatient NBP ^{17,19} | 134 | 483 | 904 | 489 | 1,763 | 3,302 | 1,318 | 3,980 | 5,403 | 1,203 | 3,633 | 4,931 | 1,170 |
| CFR general population (per 100K) ^{20, ¥} | 0.10 | 0.10 | 0.20 | 0.50 | 0.80 | 1.00 | 1.50 | 2.30 | 3.00 | 5.60 | 8.40 | 11.10 | 5.50 |
| CFR bacteremia ^{15, 21} | 6.40 | 8.30 | 11.10 | 14.60 | 16.30 | 17.30 | 10.00 | 18.80 | 17.50 | 36.30 | 46.70 | 36.30 | 37.10 |
| CFR meningitis ²²⁻²⁴ | 12.20 | 15.80 | 21.10 | 13.70 | 15.30 | 16.20 | 8.90 | 16.80 | 15.60 | 13.30 | 17.10 | 13.30 | 13.60 |
| CFR inpatient NBP ^{17,25} | 1.80 | 2.50 | 8.00 | 1.80 | 4.40 | 9.50 | 11.20 | 16.60 | 24.60 | 15.30 | 18.90 | 20.20 | 16.20 |
| General Population Health-State Utilities ²⁶ | 0.94 | 0.94 | 0.86 | 0.88 | 0.88 | 0.75 | 0.86 | 0.86 | 0.68 | 0.80 | 0.80 | 0.61 | 0.77 |
| Medical care costs of bacteremia (per case) | \$ 4,467 | \$ 3,975 | \$ 3,923 | \$ 4,564 | \$ 4,061 | \$ 4,008 | \$ 3,513 | \$ 4,602 | \$ 3,679 | \$ 3,617 | \$ 4,739 | \$ 3,789 | \$ 3,644 |
| Medical care costs of meningitis (per case) | \$ 3,353 | \$ 2,983 | \$ 2,945 | \$ 3,426 | \$ 3,048 | \$ 3,009 | \$ 2,637 | \$ 3,454 | \$ 2,762 | \$ 2,715 | \$ 3,557 | \$ 2,844 | \$ 2,735 |
| Medical care costs of inpatient NBP (per case) | \$ 1,017 | \$ 1,217 | \$ 1,743 | \$ 1,004 | \$ 1,202 | \$ 1,721 | \$ 1,155 | \$ 1,258 | \$ 1,384 | \$ 1,153 | \$ 1,256 | \$ 1,382 | \$ 1,146 |
| Medical care costs of outpatient NBP (per case) | \$ 92 | \$ 93 | \$ 116 | \$ 91 | \$ 92 | \$ 115 | \$ 96 | \$ 99 | \$ 112 | \$ 94 | \$ 97 | \$ 110 | \$ 93 |

¥ Assumption. Mortality assumed to be 1.5 and 2.0 for at-risk and high-risk, respectively, vs. healthy.

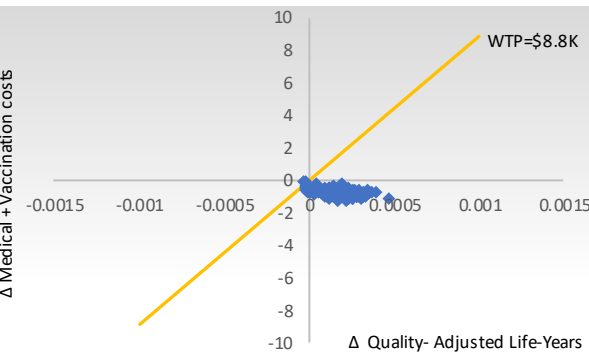
RESULTS

From a payer's perspective, PCV20 results in a dominant strategy compared to PCV15+PPV23 (WTP = \$8,441/QALY). The model estimated that PCV20 reduces bacteremia cases by 681, all-cause NBP by 7,305 and deaths due to acute disease by 780. Medical costs were reduced by \$6.96M and vaccination costs by \$19.83M for the PCV20 strategy.

Scenario analysis

- PCV20 was cost-saving in all scenario analyses.
- In the PSA 97% of the replications PCV20 was dominant and in 98% was below the WTP (1 GDP per capita = \$ 8,842).

Figure 1. probabilistic sensitivity análisis – ICER scatterplot

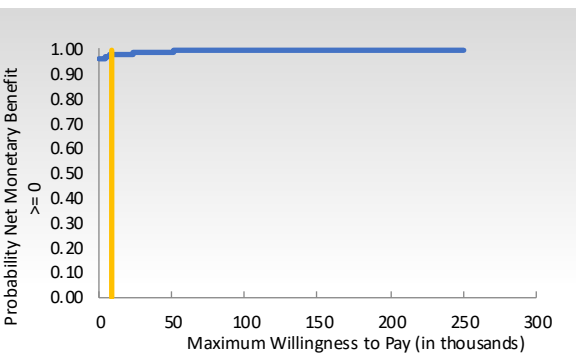


CONCLUSIONS

From a healthcare perspective PCV20 could be a dominant strategy compared to PCV15+PPV23 in Argentina. PCV20 would avert more cases of IPD, all-cause NBP and deaths with a higher number of LYG and QALYs at a lower cost.

Results were robust to alternative parameter estimates considered in scenario and sensitivity analyses

Figure 2. Acceptability curve – Cost per Quality-Adjusted Life-Year gained



REFERENCES

- Huang SS, et al. Vaccine. 2011;29(18):3398-412.
- Said MA, et al. PLoS One. 2013;8(4):e60273.
- Blasi F, et al. Clin Microbiol Infect. 2012;18 Suppl 5:7-14.
- Giglio ND, et al. Value Health Reg Issues. 2022;28:76-81.
- Argentina Strategy 2017-2018: pneumococcal vaccination, technical guidelines and vaccinator manual. Argentine MoH.
- Bonten MJ, et al. N Engl J Med. 2015;372(12):1114-25.
- Mangen MJ, et al. Eur Respir J. 2015;46(5):1407-16.
- Klugman KP, et al. N Engl J Med. 2003;349(14):1341-8.
- French N, et al. N Engl J Med. 2010;362(9):812-22.
- Djennad A, et al. EClinicalMedicine. 2018;6:42-50.
- Mangen MJ, et al. BMC Infect Dis. 2017;17(1):208.
- Melegaro A, et al. Vaccine. 2004;22(31-32):4203-14.
- Ochoa-Gondar O, et al. BMC Public Health. 2017;17(1):610.
- Proyecciones Provinciales de Población por sexo y grupo de edad 2010-2040. INDEC; 2013.
- Active Bacterial Core Surveillance (ABCs) Report Emerging Infections Program Network Streptococcus pneumoniae. CDC 2018 2020.
- Información sobre la vigilancia de las neumonías y meningitis bacterianas. SIREVA II. OPS. 2019.
- Lopardo GD, et al. BMJ Open. 2018;8(4):e019439.
- Pelton SI, et al. Clin Infect Dis. 2019;68(11):1831-8.
- Weycker D, et al. BMC Health Serv Res. 2016;16:182.
- Tasa de mortalidad por mil habitantes, según grupo de edad y sexo. Total del país. Años 2012-2018. INDEC.
- Gentile JH, et al. The University of Louisville Journal of Respiratory Infections. 2018;2(1).
- Factsheet about pneumococcal disease. ECDC.
- Gierke R, et al. CDC 2021.
- Active Bacterial Core Surveillance (ABCs) Report Emerging Infections Program Network Streptococcus pneumoniae, 2018. CDC 2020.
- Averin A, et al. Respir Med. 2021;185:106476.
- Self-Reported Population Health: An International Perspective based on EQ-5D: Springer Open; 2014.