# THE CHANGING EPIDEMIOLOGY OF INVASIVE PNEUMOCOCCAL DISEASE (IPD) IN ADULTS OF 65 YEARS OR OVER. IMPACT OF THE PAEDIATRIC **PNEUMOCOCCAL VACCINATION PROGRAMME, SPAIN, 2010-2017**

A. Navarro-Torné, C. Méndez Pfizer Spain, Madrid, Spain

Contact: dora.navarro@pfizer.com

# **INTRODUCTION**

- The paediatric heptavalent pneumococcal conjugate vaccine (PCV7) was first available in Spain in June 2001<sup>1</sup> and incorporated in the Madrid regional immunisation programme (RIP) in 2006<sup>2</sup>, remaining in the private market for other regions.
- From mid-2010 through 2016, Spanish regions introduced the 13-valent conjugate vaccine (PCV13) in their RIPs<sup>1</sup>.
- Adult vaccination with the 23-valent polysaccharide vaccine (PPV23) officially started in 2004<sup>2</sup>, and with PCV13 in 2016 for some cohorts, without expected impact for this analysis.

### **OBJECTIVES**

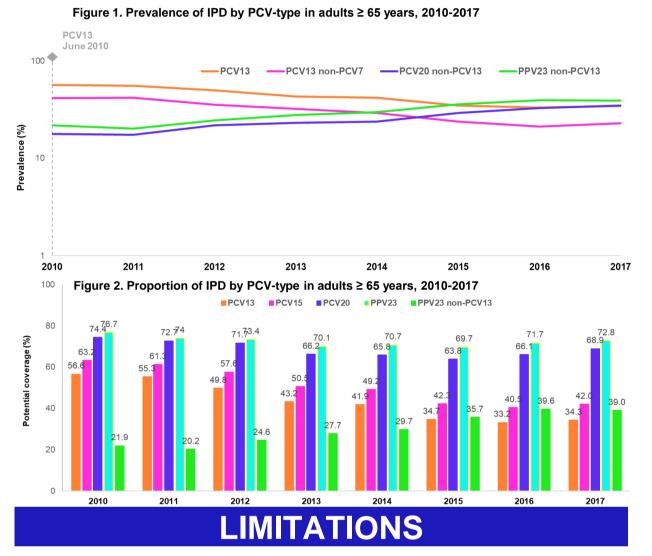
- To describe serotype distribution of invasive pneumococcal disease (IPD) in  $\geq$  65 years people for the period 2010-2017 in Spain to assess:
  - -The evolution of serotype distribution in adults since the routine paediatric use of PCV13
  - The potential coverage for PCV15 and PCV20

#### **METHODS**

- Data source: cases reported through the European Centre for Prevention and Control surveillance Disease system (available online)<sup>3</sup>
- IPD serotype specific counts were aggregated into PCV13-, PCV13 non-PCV7-, 20-valent conjugate vaccine (PCV20) non-PCV13-, PPV23-, and PPV23 non-PCV13-type groups.

#### RESULTS

- During 2010-17, a 6.1% (95%CI -11.4 to -0.6; p=0.04) and an 8.1% average annual decline (95%CI -13.7 to -2.2; p=0.02) in PCV13- and PCV13 non-PCV7-type IPD, respectively, was observed (Table 1).
- PCV20 non-PCV13-type IPD showed a 13.4% average annual increase (95%CI 7.8 to 19.2; p=0.001).
- Despite vaccination, PPV23 non-PCV13-type IPD increased an average annual 13.2% (95%CI 8.6 to 17.9; p<0.001).



- Proportions of IPD by vaccine-type groups were calculated.
- For some years ECDC reported 15B and 15C counts together as 15B/C for Spain. This approach was applied to all the period as 15B/15C strains can have reversible switching<sup>4</sup>
- The percentage change in annual number of cases was estimated using linear regression analysis of the log of the annual number of cases.
- Spanish data come from the National Reference Laboratory covering 80% of the population, thus limiting generalisability
- Roundoff in counts calculation may have introduced some bias

### **CONCLUSIONS**

- Implementation of regional paediatric PCV13 programmes in Spain has been associated with decline of PCV13-type IPD in adults  $\geq$  65 years. Further reductions may only be achieved with direct immunisation.
- In 2017, PCV20 could potentially have covered around 69% of IPD whereas PCV15 would have covered 42% (Fig 2).

Vaccine-type group	er of ca	and percent change in annual number of cases of cases of cases						Percentage change in annual number of cases (95%CI)	p value	
	2010	2011	2012	2013	2014	2015	2016	2017	-	
PCV13	531	524	525	417	373	351	304	444	-6.1 (-11.4 to -0.6)	0.04
PCV13 non-PCV7	391	399	375	311	260	241	193	295	-8.1 (-13.7 to -2.2)	0.02
PCV20 non-PCV13	168	166	231	222	213	294	301	449	13.4 (7.8 to 19.2)	0.001
PPV23	721	703	773	677	630	705	657	944	1.4 (-3.4 to 6.5)	0.5
PPV23 non-PCV13	207	193	259	268	265	361	363	506	13.2 (8.6 to 17.9)	<0.001

\*Dataset provided by ECDC based on data provided by WHO and Ministries of Health from the affected countries

## REFERENCES

1.Spanish vaccination historical and current schedules. Available at: https://www.mscbs.gob.es/profesionales/saludPublica/prevPromocion/vacunaciones/. 2. Epidemiological bulletin 2018, region of Madrid. Available at: https://www.comunidad.madrid/publicacion/ref/20209. 3. ECDC Surveillance Atlas. Available at: https .eu/en/surveillance-atlas-infectious-diseases. 4. van Selm S, et al. Infect Immun.2003;71(11):6192-8.



#### Abstract ID 470, Topic: Vaccines / C2 Impact of Vaccine programs and Serotype Replacement

Disclaimer: This poster includes data that Pfizer is not responsible for validating/storing. "The views and opinions of the authors expressed herein do not necessarily state or reflect those of the ECDC. The accuracy of the authors' statistical analysis and the findings they report are not the responsibility of ECDC. ECDC is not responsible for conclusions or opinions drawn from the data provided. ECDC is not responsible for the correctness of the data and for data management, data merging and data collation after provision of the data. ECDC shall not be held liable for improper or incorrect use of the data" Conflict of interest: All authors are Pfizer employees and may hold company stocks

12th International Symposium on Pneumococci and Pneumococcal Diseases (ISPPD), June 2020