

Invasive Pneumococcal Disease Quarterly Report

January – March 2018

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by
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Introduction

Since 17 October 2008, invasive pneumococcal disease (IPD) has been notifiable to the local Medical Officer of Health under the Health Act 1956. On 1 June 2008, pneumococcal conjugate vaccine (PCV) was added to the New Zealand childhood immunisation schedule. Initially the 7-valent conjugate vaccine (PCV7), Prevenar®, was used. In July 2011, Prevenar® was replaced on the schedule with the 10-valent conjugate vaccine (PCV10), Synflorix®. In July 2014, Synflorix® was replaced by the 13-valent conjugate vaccine (PCV13), Prevenar13®. In July 2017, Prevenar13® was replaced by Synflorix®.

PCV10 includes the seven serotypes in PCV7 (4, 6B, 9V, 14, 18C, 19F and 23F) as well as serotypes 1, 5 and 7F, and cross-reactivity to serotype 19A. PCV13 includes the 10 serotypes in PCV10 as well as serotypes 3, 6A and 19A. The recommended schedule is four doses, given at 6 weeks, 3 months, 5 months and 15 months of age.

These quarterly reports are part of an enhanced surveillance programme to monitor the impact of PCV vaccination, including the changes in vaccine valency, on the epidemiology of IPD in New Zealand.

Methods

The data presented in this report (except for immunisation status) is based on the information recorded on EpiSurv, the national notifiable disease surveillance system, as at 15 June 2018. Any changes made to EpiSurv data by public health unit staff after this date will not be reflected in this report. Immunisation status of cases that were eligible for PCV vaccination was extracted from the National Immunisation Register (NIR).

Denominator data used to determine all disease rates in this report was derived from the 2016 and 2017 mid-year population estimates published by Statistics New Zealand unless otherwise specified. Rates have not been calculated where there are fewer than five notified cases in any category.

The Fisher's exact test was used to determine statistical significance. Results are considered statistically significant when the *P* value is ≤ 0.05 .

Streptococcus pneumoniae isolates are serotyped at ESR by the capsular antigen reaction (Neufeld test) using the Danish system of nomenclature and sera obtained from the Statens Serum Institut. Methods have not been established at ESR to identify the strain type when only pneumococcal DNA, rather than an isolate, is available. Therefore, the serotype can only be determined for culture-positive IPD cases. Serotype data for invasive isolates of *S. pneumoniae* was matched with the relevant IPD case notification.

Case definition

A case of invasive pneumococcal disease is defined as:

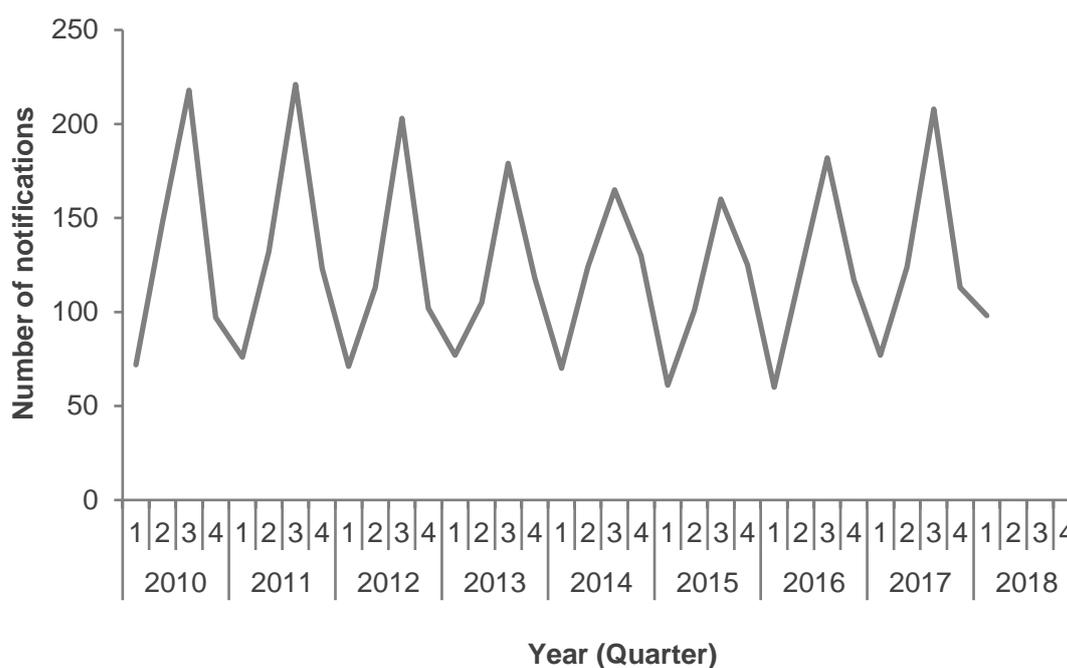
- the isolation of *S. pneumoniae* from CSF, blood or other normally sterile site; or
- the detection by nucleic acid amplification test of pneumococcal DNA in CSF, blood or other normally sterile site; or
- a positive newer-generation *S. pneumoniae* antigen test on CSF or pleural fluid.¹

¹ A positive *S. pneumoniae* antigen test on pleural fluid was added to the case definition in mid-September 2016.

Results

There were 98 IPD cases notified in the January–March 2018 quarter, compared with 77 cases in the same quarter in 2017. IPD displays a distinct seasonal pattern with a winter peak and summer trough (Figure 1). The notification rate for the latest 12-month period ending March 2018 (11.3 per 100,000 population, 543 cases) was higher than the rate for the previous 12-month period ending March 2017 (10.6 per 100,000, 497 cases).

Figure 1. Number of cases of invasive pneumococcal disease by quarter reported, January 2010–March 2018



The distribution of IPD cases and rates by age group is presented in Table 1. During the latest 12-month period, the highest rate was in the ≥ 65 years age group (32.3 per 100,000 population, 233 cases). Comparing the latest 12-month period with the previous 12-month period, there were no significant changes in the age-specific rates.

Table 1. Number of cases and rates of invasive pneumococcal disease by age group

Age group	Jan-Mar 2018	12 months ending Mar 2018		12 months ending Mar 2017	
	Cases	Cases	Rate ^a	Cases	Rate ^a
<2 years	11	30	24.8	25	20.9
2–4 years	5	25	13.5	21	11.3
5–64 years	42	255	6.8	248	6.7
≥ 65 years	40	233	32.3	203	29.1
Total	98	543	11.3	497	10.6

^a Rate is expressed as cases per 100,000 population.

The distribution of IPD cases and rates by region is presented in Table 2. The highest rates for the latest 12-month period were in the Midland and Central regions (13.8 per and 11.9 per 100,000 population, respectively, 126 cases each). Comparing the latest 12-month period with the previous 12-month period, there was a significant increase in the rate for the Central region. There were no significant changes for individual DHBs.

Table 2. Number of cases and rates of invasive pneumococcal disease by region

Region	Jan-Mar 2018	12 months ending Mar 2018		12 months ending Mar 2017	
	Cases	Cases	Rate ^a	Cases	Rate ^a
Northern ^b	38	200	10.8	225	12.5
Midland ^c	21	126	13.8	108	12.0
Central ^d	22	126	11.9	82	7.9
Southern ^e	17	91	9.4	82	8.6
Total	98	543	11.3	497	10.6

^a Rate is expressed as cases per 100,000 population.

^b Includes Northland, Waitemata, Auckland and Counties Manukau DHBs.

^c Includes Waikato, Lakes, Bay of Plenty, Tairāwhiti and Taranaki DHBs.

^d Includes Hawke's Bay, Whanganui, MidCentral, Hutt Valley, Capital & Coast, Wairarapa and Nelson Marlborough DHBs.

^e Includes West Coast, Canterbury, South Canterbury and Southern DHBs.

A culture was received at ESR for serotyping from 90 (91.8%) of the 98 cases notified in the January–March 2018 quarter. Table 3 shows the number of IPD cases due to each of the serotypes included in PCV7, PCV10 and PCV13, and due to non-PCV13 serotypes.

The number of IPD cases due to PCV13 serotypes decreased 10.1% between the last two 12-month periods (188 to 169 cases). In contrast, the number of IPD cases due to non-PCV13 serotypes increased 19.5% (282 to 337 cases). The increase in IPD due to non-PCV13 types occurred across all age groups (Table 3).

The four most prevalent serotypes during the last 12 months were 19A, 8, 22F and 3. Between the last two 12-month periods, cases of IPD due to type 19A decreased by 25.9% (81 to 60 cases) and cases of 22F remained relatively stable. However, there were notable increases in cases due to types 8, 3 and 12F: 43.8% (32 to 46 cases), 47.8% (23 to 34 cases) and 233.3% (9 to 30 cases), respectively (Table 3).

Table 3. Number of invasive pneumococcal disease cases by serotype and age group

Serotypes	Age group											
	<2 years			2–4 years			≥5 years			Total		
	Q1 2018 ^a	2018 ^b	2017 ^c	Q1 2018 ^a	2018 ^b	2017 ^c	Q1 2018 ^a	2018 ^b	2017 ^c	Q1 2018 ^a	2018 ^b	2017 ^c
4	0	0	1	0	0	0	0	15	16	0	15	17
6B	0	0	0	0	0	0	1	1	2	1	1	2
9V	0	0	0	0	0	1	1	2	4	1	2	5
14	0	0	0	0	1	0	0	0	8	0	1	8
18C	0	0	0	0	0	0	0	1	2	0	1	2
19F	0	0	0	0	0	0	0	14	10	0	14	10
23F	0	0	0	0	0	0	1	1	3	1	1	3
Total PCV7	0	0	1	0	1	1	3	34	45	3	35	47
1	1	1	0	0	0	0	3	5	1	4	6	1
5	0	0	0	0	0	0	0	0	2	0	0	2
7F	0	0	1	0	0	0	3	31	32	3	31	33
Total PCV10	1	1	2	0	1	1	9	70	80	10	72	83
3	0	3	1	0	1	2	3	30	20	3	34	23
6A	0	0	0	0	0	0	1	3	1	1	3	1
19A ^d	0	0	6	0	4	8	10	56	67	10	60	81
Total PCV13	1	4	9	0	6	11	23	159	168	24	169	188
6C	0	0	0	1	1	1	4	14	17	5	15	18
8	0	0	1	0	0	0	7	46	31	7	46	32
9N	0	0	2	0	0	0	2	11	13	2	11	15
10A	2	4	0	0	0	0	5	13	9	7	17	9
11A	0	0	0	0	0	0	3	19	11	3	19	11
12F	2	5	0	1	1	0	8	24	9	11	30	9
15A	1	1	0	0	2	0	3	20	13	4	23	13
15B	0	0	3	0	0	1	0	12	6	0	12	10
16F	0	0	0	0	0	0	1	14	11	1	14	11
17F	0	1	0	0	0	0	1	5	9	1	6	9
22F	0	1	1	0	3	1	5	32	38	5	36	40
23A	1	1	0	0	1	0	1	11	8	2	13	8
23B	0	1	0	1	1	0	5	19	19	6	21	19
33F	0	1	3	0	1	1	3	15	17	3	17	21
35B	0	0	0	0	0	1	0	5	6	0	5	7
38	2	4	0	0	0	1	2	12	6	4	16	7
Other types ^e	0	2	3	0	1	1	5	33	39	5	36	43
Total non-PCV13	8	21	13	3	11	7	55	305	262	66	337	282

^a Cases reported in the first quarter of 2018 (January–March 2018).

^b Cases reported in the 12 months ending 31 March 2018.

^c Cases reported in the 12 months ending 31 March 2017.

^d The indications for PCV10 include cross-protection against 19A disease.

^e Any of these other types accounted for <5 IPD cases during the 12 months ending 31 March 2018.

Table 4 shows the immunisation status for cases notified in the January–March 2018 quarter who were age-eligible for PCV (ie, cases born after 1 January 2008 and aged ≥ 6 weeks). Of the 16 cases that were age-eligible for PCV, one case did not have a record in the NIR. Of the remaining 15 cases, 12 were due to non-PCV13 serotypes and serotype information was not available for the other three cases.

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Table 4. Immunisation status of the invasive pneumococcal disease cases notified in the January - March 2018 quarter and who were eligible for PCV

Number of doses received ^a	Cases due to PCV7 serotypes: 4, 6B, 9V, 14, 18C, 19F or 23F ^b	Cases due to additional PCV10 serotypes: 1, 5, 7F ^b	Cases due to additional PCV13 serotypes: 3, 6A, 19A ^b	Cases due to non-PCV13 serotypes ^b	Total ^{b,c}
	Number	Number	Number	Number	Number
0	0	0	0	0	0
1	0	0	0	0	0
2	0	0	0	0	1
3	0	0	0	5	6
4	0	0	0	7	8
Total	0	0	0	12	15

^a Number of doses received prior to 14 days before onset of IPD. Onset of IPD was determined using the earliest episode date available from onset of illness date, hospitalised date or date case notified to the public health unit.

^b Only IPD cases eligible for PCV as part of the childhood immunisation schedule (ie, cases born after 1 January 2008 and aged ≥ 6 weeks) are presented.

^c The total number of cases includes three cases where serotype information was not available.

Note: Immunisation status is based on information recorded in the National Immunisation Register (NIR). There was no NIR record for one case who was eligible for PCV. This case is not included in the Table.