

# COMMUNICABLE DISEASES REPORT, NSW, FOR OCTOBER AND NOVEMBER 2003

## TRENDS

Notifications of communicable diseases through mid-spring indicated a decline in **influenza**, **invasive pneumococcal disease** and **meningococcal disease** (Figure 2, Tables 1–2). Because the surveillance case definition for **pertussis** requires, in part, the patient to have a coughing illness for 14 days, there is an inherent delay between onset of disease and notification of the case. Recent trends in case reports of pertussis are therefore likely to be substantially underestimated, and it is possible that a further rise in case reports will occur in coming months.

A large gastroenteritis outbreak caused by **Norovirus** infection was identified in October, involving over 70 people in the Greater Murray Area Health Service. The most likely cause of this outbreak was contamination of food by a food handler. Norovirus is infectious with low doses of the virus, which can survive on surfaces and in foods for long periods. People who are ill with gastroenteritis should stay home and not prepare food for anyone until 48 hours after their symptoms have completely resolved. A report of this outbreak will be published in a future issue of the *NSW Public Health Bulletin*.

For updated information, visit [www.health.nsw.gov.au](http://www.health.nsw.gov.au) and click on the link to Infectious Diseases.

## INFLUENZA SURVEILLANCE 2003

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Enhanced surveillance for influenza in NSW indicates that the 2003 influenza season peaked in August. Most cases were caused by the influenza A virus, predominantly the A/Fujian/411/2002 strain. Little influenza B infection was reported. Preliminary analysis of emergency department data, supported by anecdotal reports from clinicians, suggests that influenza may have affected more people in 2003 than in previous recent years.

In 2003, several sources of data were included in an enhanced surveillance for influenza, including:

### Sentinel general practitioners

Up to 48 general practitioners participated in weekly reporting of influenza-like illness (ILI) activity. ILI activity peaked in mid August (34.7 per 1,000 consultations). In 2002, a similar peak (36.6 per 1000 consultations) appeared in July of that year.

### Virological surveillance

Six sentinel laboratories tested 10,391 respiratory samples for the presence of influenza virus, by either direct immunofluorescence (DIF) or culture. Influenza A was found in 831 samples, and this strain peaked in mid-to-late August (23.2 per 100 samples). In 2002, detection of

influenza A peaked at 16.9 per 100 samples. Influenza B was found in 13 samples, and this strain peaked in early September (0.5 per 100 samples).

### Serological surveillance

The same six sentinel laboratories tested 4,052 serum samples for evidence (seroconversion or rise in IgG level or high single titre) of infection with influenza. Serological diagnoses of influenza A peaked in early September, at 13.6 per 100 samples. In 2002, a similar peak (14.1 per 100 samples) appeared in late August. Serological diagnoses of influenza B were rare and no peak was identified. In 2002, the peak occurred in early July (7.8 per 100 samples).

### General practitioner direct virological surveillance

In 2003, fifteen general practitioners (GPs) volunteered to provide specimens from patients who they suspected to have influenza infection for virological testing. Three-hundred-and-nine samples were taken by the GPs, of which 51 (16.5 per cent) were positive for influenza A. No samples tested positive for influenza B.

### The WHO Influenza Collaborating Centre

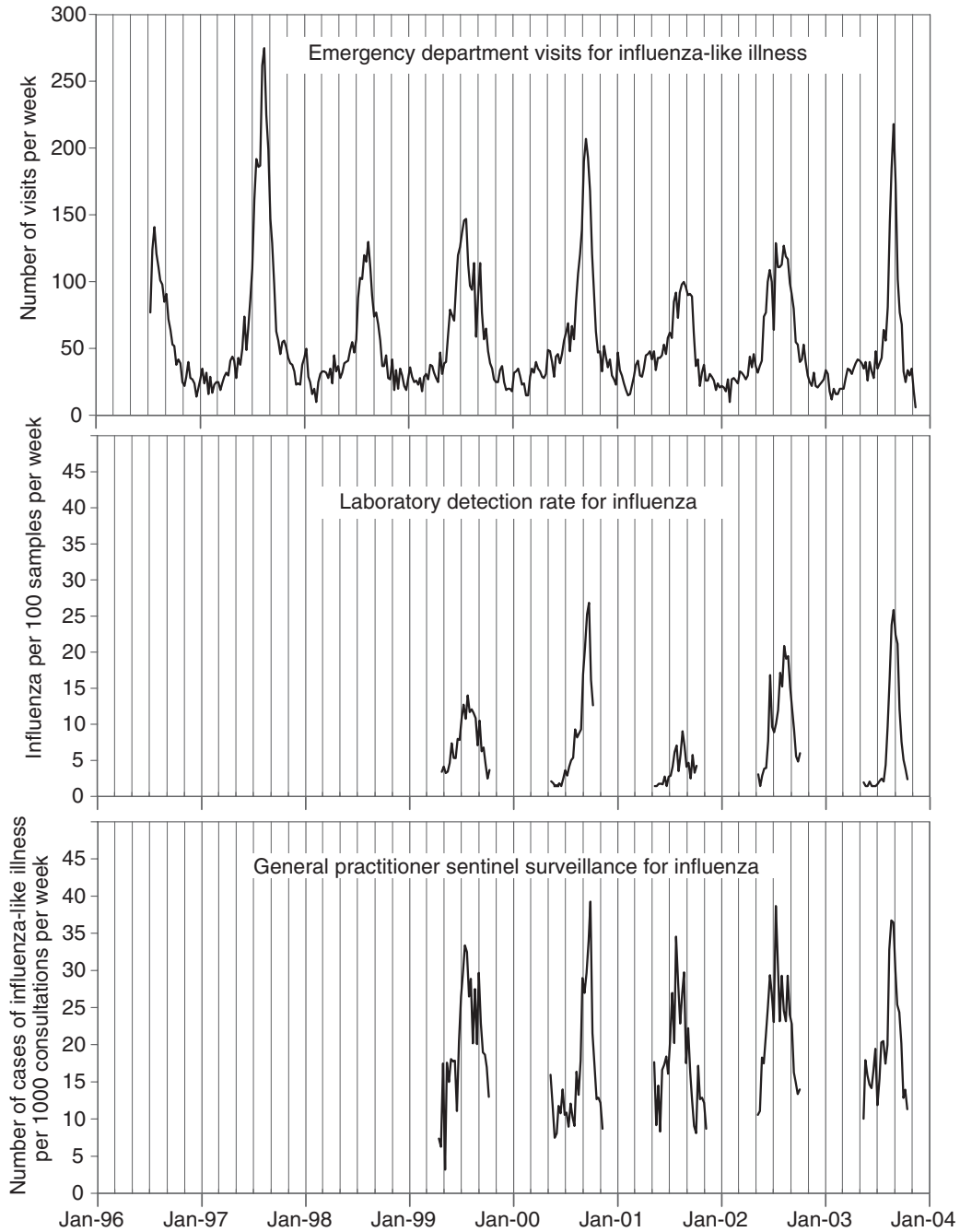
The WHO Influenza Collaborating Centre for Reference and Research on Influenza, located in Melbourne, reports that the majority of influenza A isolates identified during the peak period were A(H3) viruses of the A/Fujian/411/2002 type. This year, some antigenic drift has been detected in the virus strains circulating in Australia and New Zealand. The A/Fujian-like viruses are related to the A/Moscow-like strain included in the 2003 vaccine, and the vaccine has been demonstrated to induce antibodies to the A/Fujian-like strains but generally at a reduced level. In the last few years, dominant strains of influenza A have included A/Nanchang/95, A/Sydney/97, A/Moscow/99, and A/New Caledonia/99 (the last part of the name of each strain represents the year in which it was first identified).

### Emergency department surveillance

Information on visits to NSW Emergency Departments (EDs) collected routinely by hospitals is currently being evaluated as a monitoring tool for influenza surveillance. Figure 1 compares the number of ED visits assigned a provisional diagnosis of influenza in hospitals participating in the NSW Emergency Department Data Collection with other influenza indicators currently used in NSW. Only EDs providing reasonably-complete provisional diagnosis information using the International Classification of Diseases for the period July 1996 to October 2003 were included. The collection captures approximately two-thirds of NSW Emergency Department visits. Peaks in the number of visits to EDs assigned a provisional diagnosis of influenza corresponded to peaks in reports from laboratory virology and GP sentinel surveillance. The highest peaks in the years 1997, 2000,

**FIGURE 1**

**COMPARISON OF EMERGENCY DEPARTMENT VISITS FOR INFLUENZA-LIKE ILLNESS WITH LABORATORY-BASED DETECTION RATES AND GENERAL PRACTITIONER SENTINEL SURVEILLANCE FOR INFLUENZA, NSW, 1996–2003**



Notes: Influenza-like illness in emergency departments was based on unplanned visits assigned a principal provisional diagnosis of influenza. Laboratory and general practitioner data were only available from May 1999.

Source: NSW Emergency Department Data Collection (HOIST), Centre for Epidemiology and Research, NSW Department of Health; and NSW Influenza Surveillance Program, Communicable Diseases Branch, NSW Department of Health.

and 2003 coincided with a predominance of newly emergent strains of influenza A virus among laboratory samples that had strain identification performed (Sydney/97, Moscow/99 and Fujian/2002 in each of those peak years respectively).

#### **COMMENT**

The data and information collected from these sources indicate that influenza peaks each year in winter, usually between mid-July and mid-September. In 2003, the peak was in August, and influenza activity may have been more widespread than in recent previous years.

There are several limitations to these data. First, none of the surveillance systems mentioned here are very sensitive: all collect data on only a very small proportion of people infected with influenza in NSW, and this proportion may vary over time, rendering comparisons open to bias. Second, none of the systems provide a very representative sample of influenza cases either by the

demographics of the affected people, their place of residence, or severity of illness. Laboratory surveillance is based in urban hospitals, and is more likely to include very sick children (who tend to present to hospital for testing) than the GP systems. Participating GPs are not located randomly across the state. Third, apart from the laboratory-based systems, the diagnosis of influenza-like illness is not specific, and the systems are likely to pick up a range of other respiratory conditions not caused by influenza viruses.

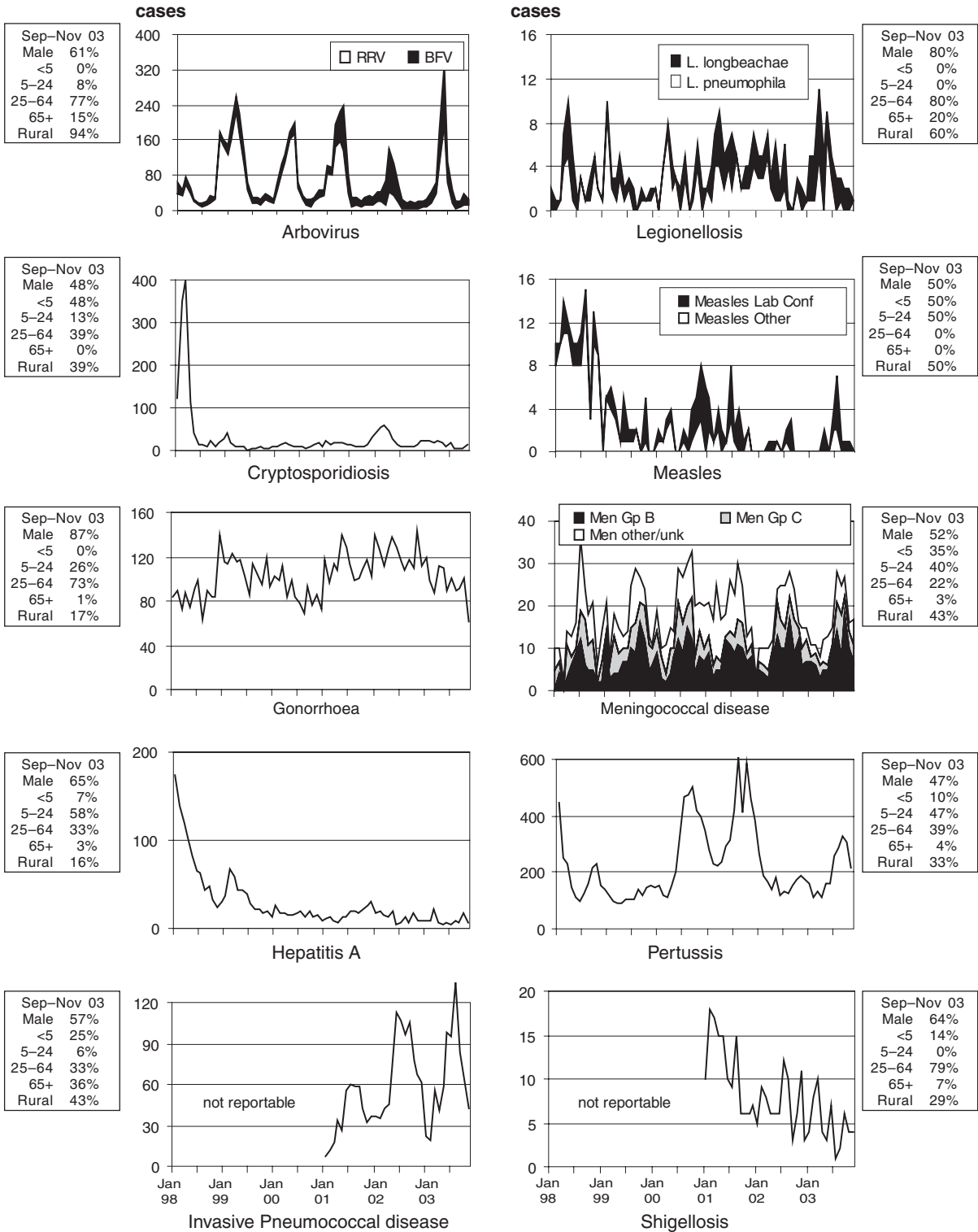
The apparent triennial variation in the magnitude of the influenza peaks found in Emergency Department visits appears to be temporally associated with the predominance of a newly emergent A strain among circulating strains of the influenza virus for the year. These data suggest that ED surveillance could be a useful tool for monitoring not only the occurrence of influenza epidemics in NSW but also their extent. The Centre for Epidemiology and Research has developed methods for the rapid transfer and analysis of these data for surveillance purposes. ☒

**FIGURE 2**

**REPORTS OF SELECTED COMMUNICABLE DISEASES, NSW, JANUARY 1996 TO NOVEMBER 2003, BY MONTH OF ONSET**

These are preliminary data: case counts for recent months may increase because of reporting delays. Laboratory-confirmed cases, except for measles, meningococcal disease and pertussis.

NSW population	
Male	50%
<5	7%
5-24	28%
25-64	52%
65+	13%
Rural*	42%



**TABLE 1 REPORTS OF NOTIFIABLE CONDITIONS RECEIVED IN OCTOBER 2003 BY AREA HEALTH SERVICES**

Condition	Area Health Service													for Oct'	Total To date'				
	CSA	NSA	WSA	WEN	SWS	CCA	HUN	ILL	SES	NRA	MNC	NEA	MAC			MWA	FWA	GMA	SA
<b>Blood-borne and sexually transmitted</b>																			
Chancroid*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Chlamydia (genital)*	82	76	56	21	33	14	55	18	123	31	23	31	8	15	9	22	17	4	642
Gonorrhoea*	21	10	2	3	5	1	6	-	41	6	4	1	1	-	-	1	-	-	102
Hepatitis B - acute viral*	-	1	1	-	1	-	1	-	2	-	-	-	-	-	1	-	-	-	7
Hepatitis B - other*	36	26	31	6	35	3	5	6	39	1	3	2	-	-	3	-	-	2	198
Hepatitis C - acute viral*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Hepatitis C - other*	46	37	9	17	57	22	29	29	47	13	19	11	6	20	8	19	16	31	441
Hepatitis D - unspecified*	-	-	-	-	1	-	-	1	-	-	-	-	-	-	-	-	-	-	2
Syphilis	10	3	7	2	8	3	-	2	29	4	1	2	3	-	-	1	2	-	78
<b>Vector-borne</b>																			
Barmah Forest virus*	-	-	-	-	-	-	-	-	-	9	6	1	-	-	-	-	-	-	16
Ross River virus*	-	-	-	-	-	-	-	-	-	4	1	-	1	-	-	2	3	-	11
Arboviral infection (Other)*	-	2	-	-	1	-	1	-	-	-	-	-	-	-	-	-	-	-	2
Malaria*	-	2	2	-	-	-	-	-	1	-	-	-	-	-	-	-	-	-	5
<b>Zoonoses</b>																			
Anthrax*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Brucellosis*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Leptospirosis*	-	-	-	-	-	2	-	-	-	-	-	2	-	-	-	-	-	-	4
Lyssavirus*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Psittacosis*	-	1	-	1	-	-	2	-	-	-	-	3	-	-	1	-	1	-	9
Q fever*	1	-	-	-	-	-	4	-	-	-	2	5	3	-	2	-	1	-	18
<b>Respiratory and other</b>																			
Blood lead level*	-	5	-	-	2	-	4	-	1	-	1	-	-	-	7	-	1	-	21
Influenza*	2	7	19	5	9	2	1	2	7	2	3	4	-	-	-	-	-	-	63
Invasive pneumococcal infection*	6	6	6	1	3	12	13	4	8	2	2	-	1	3	-	2	-	-	69
<i>Legionella longbeachae</i> infection*	-	-	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	1
<i>Legionella pneumophila</i> infection*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	31
Legionnaires' disease (Other)*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	19
Leprosy	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Meningococcal infection (invasive)*	1	1	1	-	3	1	1	3	1	-	-	1	-	-	-	3	1	-	1
Tuberculosis	4	-	6	2	1	-	-	-	6	1	1	-	-	-	1	-	-	-	18
-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	23
<b>Vaccine-preventable</b>																			
Adverse event after immunisation	-	-	2	-	3	-	3	-	-	-	-	-	2	-	-	5	-	-	15
<i>H. Influenzae b</i> infection (invasive)*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Measles	1	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	1
Mumps*	2	-	-	-	1	-	-	-	1	-	-	-	-	-	-	-	-	-	4
Pertussis	32	45	50	9	40	6	16	9	31	2	6	2	1	7	-	20	9	-	285
Rubella*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	1,926
Tetanus	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	22
-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	1
<b>Enteric</b>																			
Botulism	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Cholera*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Cryptosporidiosis*	-	-	1	-	-	-	-	-	2	-	-	1	-	-	-	1	-	-	5
Giardiasis*	-	12	12	6	7	1	7	2	6	-	3	3	2	9	-	1	-	-	71
Haemolytic uraemic syndrome	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	3
Hepatitis A*	3	1	2	-	7	-	-	-	1	-	-	-	-	-	-	-	-	-	14
Hepatitis E*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	1
Listeriosis*	-	1	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	6
Salmonellosis *	8	12	10	5	12	8	5	7	8	12	4	5	-	7	-	1	2	-	24
Shigellosis*	-	2	-	-	1	-	-	-	1	1	-	-	-	1	-	-	-	-	1,596
Typhoid and paratyphoid*	-	-	-	-	2	-	-	-	2	-	-	-	-	-	-	-	-	-	5
Verotoxin producing <i>E. coli</i> *	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	5

\* lab-confirmed cases only + includes cases with unknown postcode \*\* HIV and AIDS data are reported separately in the NSW Public Health Bulletin each quarter

CSA = Central Sydney Area	WEN = Wentworth Area	HUN = Hunter Area	NRA = Northern Rivers Area	MAC = Macquarie Area	GMA = Greater Murray Area
NSA = Northern Sydney Area	SWS = South Western Sydney Area	ILL = Illawarra Area	MNC = North Coast Area	MWA = Mid Western Area	SA = Southern Area
WSA = Western Sydney Area	CCA = Central Coast Area	SES = South Eastern Sydney Area	NEA = New England Area	FWA = Far West Area	CHS = Corrections Health Service

**TABLE 2 REPORTS OF NOTIFIABLE CONDITIONS RECEIVED IN NOVEMBER 2003 BY AREA HEALTH SERVICES**

Condition	Area Health Service													Total for Nov <sup>1</sup>	Total To date <sup>2</sup>				
	CSA	NSA	WSA	WEN	SWS	CCA	HUN	ILL	SES	NRA	MNC	NEA	MAC			MWA	FWA	GMA	SA
<b>Blood-borne and sexually transmitted</b>																			
Chancroid*	71	73	79	17	42	-	-	63	33	138	26	34	24	11	15	1	17	15	2
Chlamydia (genital)*	20	1	4	1	3	-	3	1	-	33	4	2	1	-	-	-	2	-	-
Gonorrhoea*	-	1	-	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-
Hepatitis B - acute viral*	42	36	-	4	53	-	7	1	37	3	2	7	3	1	-	2	1	5	
Hepatitis B - other*	-	-	1	-	-	-	4	-	-	-	-	-	-	-	-	-	-	-	
Hepatitis C - acute viral*	72	32	14	20	83	-	34	27	79	32	28	9	8	11	-	26	13	37	
Hepatitis C - other*	-	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	
Hepatitis D - unspecified*	28	2	7	1	20	-	-	-	36	6	-	3	-	-	-	-	-	-	
Syphilis	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
<b>Vector-borne</b>																			
Barmah Forest virus*	-	-	-	-	-	-	1	-	-	-	10	6	-	-	-	-	-	-	
Ross River virus*	-	-	-	-	-	-	2	-	-	-	5	3	-	1	-	-	-	1	
Arboviral infection (Other)*	-	-	-	-	1	-	-	-	-	-	-	-	-	1	-	-	-	-	
Malaria*	1	-	2	1	1	-	1	-	2	2	-	1	2	-	-	1	-	14	
<b>Zoonoses</b>																			
Anthrax*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Brucellosis*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Leptospirosis*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Lyssavirus*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Psittacosis*	-	-	-	-	-	-	5	-	-	1	1	2	-	-	-	-	-	9	
Q fever*	-	-	-	-	-	-	-	-	1	1	-	8	7	-	-	-	-	17	
<b>Respiratory and other</b>																			
Blood lead level*	1	-	-	-	4	-	2	1	-	-	-	2	1	-	-	1	-	-	
Influenza*	1	1	7	-	5	-	-	-	8	2	-	1	-	-	-	1	-	12	
Invasive pneumococcal infection*	1	8	5	5	13	-	4	4	10	1	8	-	2	1	-	-	4	26	
<i>Legionella longbeachae</i> infection*	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	67	
<i>Legionella pneumophila</i> infection*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	1	
Legionnaires' disease (Other)*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	32	
Leprosy	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	19	
Meningococcal infection (invasive)*	-	1	1	1	2	-	-	1	5	1	-	-	-	1	-	-	-	1	
Tuberculosis	4	4	7	-	4	-	1	2	9	-	-	-	1	-	-	-	-	13	
<b>Vaccine-preventable</b>																			
Adverse event after immunisation	-	-	-	-	1	-	3	-	2	-	-	-	-	1	2	-	-	9	
<i>H. Influenzae b</i> infection (invasive)*	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	1	
Measles	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	16	
Mumps*	2	2	1	-	-	-	-	-	1	-	-	-	-	-	-	-	-	4	
Pertussis	28	50	60	11	51	-	19	24	70	7	9	5	3	4	-	31	26	398	
Rubella*	-	-	-	-	-	-	-	-	1	-	-	-	-	-	-	-	-	1	
Tetanus	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	23	
<b>Enteric</b>																			
Botulism	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Cholera*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
Cryptosporidiosis*	-	-	2	-	2	-	2	-	3	-	-	2	-	1	-	-	-	12	
Giardiasis*	4	14	11	5	4	-	5	2	19	-	2	3	4	3	-	1	2	79	
Haemolytic uraemic syndrome	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	3	
Hepatitis A*	3	-	3	-	5	-	-	-	-	2	-	-	-	1	-	-	-	14	
Hepatitis E*	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	6	
Listeriosis*	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	1	
Salmonellosis (not otherwise specified)*	8	15	14	4	9	-	12	5	16	11	1	1	1	3	-	1	1	105	
Shigellosis*	-	1	1	-	1	-	-	-	3	-	-	-	-	-	-	-	-	7	
Typhoid and paratyphoid*	-	-	-	-	-	-	-	-	2	-	-	-	-	-	-	-	-	3	
Verotoxin producing <i>E. coli</i> *	-	-	-	-	-	-	1	-	-	-	-	-	-	-	-	-	-	1	

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