Additional reports

Australian Sentinel Practice Research Network

The Research and Health Promotion Unit of the Royal Australian College of General Practitioners operates the Australian Sentinel Practice Research Network (ASPREN). ASPREN is a national network of general practitioners who report presentations of defined medical conditions each week. The aim of ASPREN is to provide an indicator of the burden of disease in the primary health care setting and to detect trends in consultation rates.

There are currently about 50 general practitioners participating in the network from all states. Seventyfive per cent of these are in metropolitan areas and the remainder are rural based. Between 4,000 and 6,000 consultations are recorded each week.

The list of conditions is reviewed annually by the ASPREN management committee and an annual report is published.

In 2004, nine conditions are being monitored, five of which are related to communicable diseases. These include influenza, gastroenteritis, varicella and shingles. Definitions of these conditions are described in Surveillance systems reported in CDI, published in Commun Dis Intell 2004;28:99. Note that in 2004, two case definitions for influenza are being recorded in parallel.

Data from 1 July to 30 September 2004 are shown as the rate per 1,000 consultations in Figures 4, 5, 6 and 7.

Figure 4. Consultation rates for influenza-like illness, ASPREN, 1 July to 30 September 2004, by week of report



Figure 5. Consultation rates for gastroenteritis, ASPREN, 1 July to 30 September 2004, by week of report



Figure 6. Consultation rates for chickenpox, ASPREN, 1 July to 30 September 2004, by week of report







Childhood immunisation coverage

Tables 6, 7 and 8 provide the latest quarterly report on childhood immunisation coverage from the Australian Childhood Immunisation Register (ACIR).

The data show the percentage of children fully immunised at age 12 months for the cohort born between 1 April and 30 June 2003; at 24 months of age for the cohort born between 1 April and 30 June 2002; and at 6 years of age for the cohort born between 1 April and 30 June 1998, according to the Australian Standard Vaccination Schedule.

For information about the Australian Childhood Immunisation Register see Surveillance systems reported in CDI, published in Commun Dis Intell 2004;28:102 and for a full description of the methodology used by the Register see Commun Dis Intell 1998;22:36–37.

Commentary on the trends in ACIR data is provided by the National Centre for Immunisation Research and Surveillance of Vaccine Preventable Diseases (NCIRS). Telephone: +61 2 9845 1256. Email: brynleyh@chw.edu.au.

Immunisation coverage for 'fully immunised' at 12 months of age for Australia increased marginally from the last quarter by 0.4 percentage points to 91.3 per cent (Table 6). There was a substantial increase in 'fully immunised' coverage by State and Territory in one jurisdiction, the Northern Territory, with an increase of 5.3 percentage points, whilst all other jurisdictions experienced very little change in coverage. As expected, the Northern Territory also had increases in coverage for individual vaccines. Apparently large changes in coverage in jurisdictions like the Northern Territory and the Australian Capital Territory, who have relatively small populations, can result from small absolute numbers of unimmunised children and should be treated with caution.

Coverage for 'fully immunised' at 24 months of age for Australia increased marginally from the last quarter by 0.6 percentage points to 92.3 per cent (Table 7). Coverage for individual vaccines increased marginally in most jurisdictions with coverage greater than 95 per cent in almost all jurisdictions for all vaccines except Hib. HepB coverage at 24 months of age is now greater than 98 per cent in the Northern Territory.

Table 8 shows immunisation coverage estimates for 'fully immunised' and for individual vaccines at six years of age for Australia and by state/territory. 'Fully immunised' coverage at six years of age for Australia was unchanged overall, apart from increases in Tasmania (+3.1%) and in the Northern Territory (+4.1%), also reflected in individual vaccines. Coverage for vaccines assessed at six years is at or near 85 per cent in the most jurisdictions, but Western Australia remains well below the average.

Figure 8 shows the trends in vaccination coverage from the first ACIR-derived published coverage estimates in 1997 to the current estimates. There is a clear trend of increasing vaccination coverage over time for children aged 12 months, 24 months and six years, although the rate of increase has slowed over the past year for all age groups. The figure shows that there have now been four consecutive quarters where 'fully immunised' coverage at 24 months of age has been greater than 'fully immunised' coverage at 12 months of age, following the removal of the requirement for 18 month DTPa vaccine.





Acknowledgement: These figures were provided by the Health Insurance Commission (HIC), to specifications provided by the Australian Government Department of Health and Ageing. For further information on these figures or data on the Australian Childhood Immunisation Register please contact the Immunisation Section of the HIC: Telephone: +61 2 6124 6607.

Vaccine				State or	territory				
	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Australia
Number of children	959	21,653	921	12,711	4,371	1,347	15,155	6,124	63,241
Diphtheria, tetanus, pertussis (%)	94.3	92.7	91.9	92.7	92.8	93.7	93.2	90.7	92.7
Poliomyelitis (%)	94.3	92.6	91.4	92.6	92.6	93.5	93.1	90.7	92.6
<i>Haemophilus influenzae</i> type b (%)	95.6	94.6	95.7	94.8	95.7	95.7	95.1	93.8	94.8
Hepatitis B (%)	95.3	95.3	96.3	94.8	95.8	95.6	94.7	93.3	94.9
Fully immunised (%)	93.4	91.3	90.5	91.7	91.7	92.4	91.7	88.8	91.3
Change in fully immunised since last guarter (%)	+2.7	+0.8	+5.3	+0.1	+0.3	-1.0	+0.0	-0.5	+0.4

Table 6.Percentage of children immunised at 1 year of age, preliminary results by vaccine andstate or territory for the birth cohort 1 April and 30 June 2003; assessment date 30 September 2004

Table 7.Percentage of children immunised at 2 years of age, preliminary results by vaccine andstate or territory for the birth cohort 1 April and 30 June 2002, assessment date 30 September 2004*

Vaccine				State or	territory				
	ACT	NSW	NT	Qld	SA	Tas.	Vic.	WA	Australia
Number of children	1,003	21,222	873	12,724	4,274	1,444	15,217	6,184	62,941
Diphtheria, tetanus, pertussis (%)	95.7	95.4	96.9	95.0	95.4	96.3	95.7	94.4	95.3
Poliomyelitis (%)	95.6	95.2	97.1	94.9	95.4	96.2	95.7	94.2	95.2
<i>Haemophilus influenzae</i> type b (%)	94.7	93.4	95.5	94.0	94.1	94.5	94.2	92.6	93.8
Measles, mumps, rubella (%)	95.0	93.5	95.5	93.9	94.5	94.7	94.6	92.8	93.9
Hepatitis B(%)	96.1	95.7	98.2	95.6	96.2	96.4	96.3	95.4	95.9
Fully immunised (%)	92.7	91.8	93.8	92.3	93.0	93.8	93.1	90.6	92.3
Change in fully immunised since last									
quarter (%)	+2.7	+0.8	-0.7	+0.4	+0.3	-1.0	+0.8	-0.0	+0.5

Table 8.Percentage of children immunised at 6 years of age, preliminary results by vaccine andstate or territory for the birth cohort 1 April and 30 June 1998; assessment date 30 September 2004

				State or te	erritory				
	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Australia
Number of children	1,049	21,810	892	13,394	4,674	1,496	15,498	6,733	65,546
Diphtheria, tetanus, pertussis (%)	85.9	85.3	83.7	85.1	85.3	85.2	87.4	81.6	85.4
Poliomyelitis (%)	86.0	85.2	85.3	85.2	85.4	85.2	86.9	82.0	85.3
Measles, mumps, rubella (%)	84.9	84.3	85.2	84.9	85.0	84.3	87.1	81.4	84.8
Fully immunised (%)	83.8	83.1	82.7	83.7	83.9	83.4	85.7	80.1	83.6
Change in fully immunised since last									
quarter (%)	-1.1	-0.0	+4.1	+0.0	+0.6	+3.1	+0.2	-1.0	+0.1

* The 12 months age data for this cohort was published in Commun Dis Intell 2003;27:569.

Gonococcal surveillance

John Tapsall, The Prince of Wales Hospital, Randwick, NSW, 2031 for the Australian Gonococcal Surveillance Programme.

The Australian Gonococcal Surveillance Programme (AGSP) reference laboratories in the various States and Territories report data on sensitivity to an agreed 'core' group of antimicrobial agents quarterly. The antibiotics which are currently routinely surveyed are penicillin, ceftriaxone, ciprofloxacin and spectinomycin, all of which are administered as single dose regimens and currently used in Australia to treat gonorrhoea. When in vitro resistance to a recommended agent is demonstrated in 5% or more of isolates from a general population, it is usual to remove that agent from the list of recommended treatments¹. Additional data are also provided on other antibiotics from time to time. At present all laboratories also test isolates for the presence of high level (plasmid-mediated) resistance to the tetracyclines, known as TRNG. Tetracyclines are however not a recommended therapy for gonorrhoea in Australia. Comparability of data is achieved by means of a standardised system of testing and a programme-specific quality assurance process. Because of the substantial geographic differences in susceptibility patterns in Australia, regional as well as aggregated data are presented. For more information see Commun Dis Intell 2004;28:100.

Reporting period 1 July to 30 September 2004

The AGSP laboratories examined a total of 829 isolates in this quarter. The total is slightly less than the 857 isolated or referred in 2003. About 31% of the current total was from New South Wales, 24% from Victoria, 19% from Queensland, 13% from the Northern Territory, 10% from Western Australia and 3% from South Australia. Isolates from Tasmania (6) and the Australian Capital Territory (4) were few.

Quinolone antibiotics

The total number (200) and proportion (24%) of all quinolone resistant *N. gonorrhoeae* (QRNG) is at an historical high. In the first quarter of 2004 there were 188 QRNG (20.5%), and in the second quarter 172 (20.2%). The numbers here are substantially higher than the corresponding figures in the third quarter of 2003 (136 isolates, 16%). The majority of the QRNG (184 of 200, 92%) exhibited higher-level resistance. QRNG are defined as those isolates with an MIC to ciprofloxacin equal to or greater than 0.06 mg/L. QRNG are further subdivided into less sensitive (ciprofloxacin MICs 0.06 - 0.5 mg/L) or resistant (MIC => 1 mg/L) groups.

QRNG were again widely distributed. The highest number, 93, was found in New South Wales (36.6% of isolates in that State) while 64 QRNG were 33% of gonococci in Victoria. In Queensland there were 27 QRNG (17%), seven in Western Australia (9%), four in South Australia (17%), two in both the Northern Territory and Tasmania and one in the Australian Capital Territory.

Figure 9. The distribution of quinolone resistant isolates of *Neisseria gonorrhoeae* in Australia by jurisdiction, 1 July to 30 September 2004



LS QRNG	Ciprofloxacin MICs 0.06 - 0.5 mg/L
R QRNG	Ciprofloxacin MICs => 1 mg/L.

Penicillins

In this quarter 20% of all isolates examined were penicillin resistant by one or more mechanisms—10.6% penicillinase producing *Neisseria gonorrhoeae* (PPNG) and 9.4% by chromosomal mechanisms (CMRNG). The proportion of all penicillin resistant strains is little changed from the previous quarter and a slight increase from the 18% detected in the third quarter of 2003. The number of PPNG increased to 88 from the 77 seen in the same period in 2003, but the number of CMRNG was essentially unchanged (77 for this period in 2004, 76 in 2003). The proportion of all strains resistant to the penicillins by any mechanism ranged from 3.8% in the Northern Territory to 33% in Western Australia.

Figure 10 shows the proportions of gonococci fully sensitive (MIC <= 0.03 mg/L), less sensitive (MIC 0.06–0.5 mg/L), relatively resistant (MIC => 1 mg/L) or else PPNG, aggregated for Australia and by State and Territory. A high proportion those strains classified as PPNG or else resistant by chromosomal mechanisms fail to respond to treatment with penicillins (penicillin, amoxycillin, ampicillin) and early generation cephalosporins.

Figure 10. Categorisation of gonococci isolated in Australia by penicillin susceptibility and by region, 1 July to 30 September 2004



FS	fully sensitive to penicillin, MIC <,= 0.03 mg/L.
LS	less sensitive to penicillin, MIC 0.06 – 0.5 mg/L.
RR	relatively resistant to penicillin, MIC >,= 1 mg/L.
PPNG	penicillinase producing Neisseria gonorrhoeae.

The highest proportion of PPNG was found in Western Australia where the 16 PPNG were 20.3% of all isolates. Thirty-five PPNG representing 13.8% of all isolates were found in New South Wales, 15 (9.4%) in Queensland and 16 (8.2%) in Victoria. There was a single PPNG in South Australia, Tasmania and the Australian Capital Territory and three in the Northern Territory. The number of CMRNG was highest in Victoria (31, 16%) and New South Wales (32, 13%) and in Western Australia 10 CMRNG isolates were 12.6% of the total. Elsewhere CMRNG were in low numbers (Queensland, Tasmania, South Australia) or absent (Northern Territory, Australian Capital Territory).

Ceftriaxone.

An increased number of isolates (12, 4.7%) with decreased susceptibility to ceftriaxone were detected in New South Wales in this quarter, but none were seen elsewhere. Small numbers of these strains have been present for a number of years, mostly in New South Wales, but only occasionally in other jurisdictions.

Spectinomycin

All isolates susceptible to this injectable agent.

High level tetracycline resistance (TRNG)

Both the number (121) and proportion (14.6%) of TRNG continued to increase from the 2003 figures (92, 11.5%). TRNG were found in all jurisdictions

with 22 (28%) in Western Australia, 56 (22%) in New South Wales, 18 (11%) in Queensland and 19 (10%) in Victoria.

Reference

 Management of sexually transmitted diseases. World Health Organization 1997; Document WHO/GPA/ TEM94.1 Rev.1 p 37.

Meningococcal surveillance

John Tapsall, The Prince of Wales Hospital, Randwick, NSW, 2031 for the Australian Meningococcal Surveillance Programme.

The reference laboratories of the Australian Surveillance Programme report Meningococcal data on the number of laboratory confirmed cases confirmed either by culture or by non-culture based techniques. Culture positive cases, where a Neisseria meningitidis is grown from a normally sterile site or skin, and non-culture based diagnoses, derived from results of nucleic acid amplification assays and serological techniques, are defined as invasive meningococcal disease (IMD) according to Public Health Laboratory Network definitions. Data contained in the quarterly reports are restricted to a description of the number of cases per jurisdiction, and serogroup, where known. A full analysis of laboratory confirmed cases of IMD is contained in the annual reports of the Programme, published in Communicable Diseases Intelligence.

Laboratory confirmed cases of invasive meningococcal disease for the period 1 July to 30 September 2004, are included in this issue of Communicable Diseases Intelligence (Table 9).

Reporting period 1 July to 30 September 2004

Jurisdiction	Year						S	erog	roup						
			A	E	В	(С		Y	W	135	١	1D	A	AII
		Q3	ytd	Q3	ytd	Q3	ytd	Q3	ytd	Q3	ytd	Q3	ytd	Q3	ytd
Australian Capital Territory	2004			0	3	3	7							3	10
	2003			(2)	(3)	(2)	(2)							(4)	(5)
New South Wales	2004			22	60	6	15	1	3	2	4	2	12	33	94
	2003			(38)	(75)	(19)	(32)	(1)	(4)	(0)	(1)	(3)	(15)	(61)	(127)
Northern Territory	2004			0	5	0	0			0	1			0	6
	2003			(3)	(9)	(0)	(0)			(1)	(1)			(4)	(10)
Queensland	2004	0	1	13	36	8	20	0	1	1	2	0	2	22	62
	2003	(1)	(1)	(17)	(34)	(16)	(31)	(1)	(1)	(0)	(0)	(0)	(8)	(35)	(75)
South Australia	2004			2	11	1	1							3	12
	2003			(7)	(15)	(1)	(2)	(1)	(1)	(1)	(1)			(10)	(19)
Tasmania	2004			3	6	5	5			0	1	1	3	9	15
	2003			(3)	(3)	(4)	(5)							(7)	(8)
Victoria	2004			17	45	3	12	0	3	2	2	1	3	23	65
	2003			(22)	(35)	(17)	(39)	(2)	(2)	(0)	(1)	(1)	(6)	(42)	(83)
Western Australia	2004			11	23	2	4			1	1			14	28
	2003			(11)	(22)	(2)	(5)	(0)	(1)					(13)	(28)
Australia	2004	0	1	68	189	28	64	1	7	6	11	4	20	107	292
	2003	(1)	(1)	(103)	(196)	(61)	(116)	(5)	(9)	(2)	(4)	(4)	(29)	(176)	(355)

Table 9.Number of laboratory confirmed cases of invasive meningococcal disease, Australia, 1 Julyto 30 September 2004, by jurisdiction and serogroup

Numbers of laboratory confirmed diagnoses of IMD made in the same periods in 2003 are also shown in parenthesis.

Q3 = third quarter; ytd = year to 30 September 2004; ND = not determined.

HIV and AIDS surveillance

National surveillance for HIV disease is coordinated by the National Centre in HIV Epidemiology and Clinical Research (NCHECR), in collaboration with State and Territory health authorities and the Commonwealth of Australia. Cases of HIV infection are notified to the National HIV Database on the first occasion of diagnosis in Australia, by either the diagnosing laboratory (Australian Capital Territory, New South Wales, Tasmania, Victoria) or by a combination of laboratory and doctor sources (Northern Territory, Queensland, South Australia, Western Australia). Cases of AIDS are notified through the State and Territory health authorities to the National AIDS Registry. Diagnoses of both HIV infection and AIDS are notified with the person's date of birth and name code, to minimise duplicate notifications while maintaining confidentiality.

Tabulations of diagnoses of HIV infection and AIDS are based on data available three months after the end of the reporting interval indicated, to allow for reporting delay and to incorporate newly available information. More detailed information on diagnoses of HIV infection and AIDS is published in the quarterly Australian HIV Surveillance Report, and annually in ⁽HIV/AIDS, viral hepatitis and sexually transmissible infections in Australia, annual surveillance report'. The reports are available from the National Centre in HIV Epidemiology and Clinical Research, 376 Victoria Street, Darlinghurst NSW 2010. Internet: http://www.med.unsw.edu.au/nchecr. Telephone: +61 2 9332 4648. Facsimile: +61 2 9332 1837. For more information see Surveillance systems reported in CDI, published in Commun Dis Intell 2004;28:99.

HIV and AIDS diagnoses and deaths following AIDS reported for 1 April to 30 June 2004, as reported to 30 September 2004, are included in this issue of Communicable Diseases Intelligence (Tables 10 and 11).

				St	ate or	territo	ry				Total for	Australia	
	Sex	ACT	NSW	NT	QId	SA	Tas	Vic	WA	This period 2004	This period 2003	Year to date 2004	Year to date 2003
HIV													
diagnoses	Female	0	10	0	7	1	1	10	0	29	23	67	43
	Male	0	82	0	28	4	3	41	0	158	208	359	410
	Not reported	0	1	0	0	0	0	0	0	1	1	2	2
	Total ¹	0	93	0	35	5	4	51	0	188	232	429	455
AIDS													
diagnoses	Female	0	2	0	1	0	0	1	0	4	3	7	7
	Male	0	11	0	7	2	1	3	0	24	52	62	87
	Total ¹	0	13	0	8	2	1	4	0	28	55	70	95
AIDS													
deaths	Female	0	1	0	0	0	0	0	0	1	1	2	5
	Male	0	7	0	3	5	0	0	0	15	14	27	31
	Total ¹	0	8	0	3	5	0	0	0	16	15	29	36

Table 10.New diagnoses of HIV infection, new diagnoses of AIDS and deaths following AIDSoccurring in the period 1 April to 30 June 2004, by sex and state or territory of diagnosis

1. Persons whose sex was reported as transgender are included in the totals.

Table 11.Cumulative diagnoses of HIV infection, AIDS and deaths following AIDS since theintroduction of HIV antibody testing to 30 June 2004, by sex and state or territory

					State o	r territory				
	Sex	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Australia
HIV diagnoses		30	759	17	222	82	8	303	159	1,580
		247	12,507	121	2,373	805	88	4,664	1,049	21,854
		0	238	0	0	0	0	22	0	260
		277	13,532	138	2,604	888	96	5,008	1,215	23,758
AIDS diagnoses	Female	9	220	1	61	30	4	91	34	450
	Male	92	5,094	41	964	385	48	1,844	407	8,875
	Total ¹	101	5,329	42	1,027	416	52	1,945	443	9,355
AIDS deaths	Female	6	127	0	40	20	2	58	22	275
	Male	71	3,442	26	625	262	32	1,353	277	6,088
	Total ¹	77	3,578	26	667	282	34	1,419	300	6,383

1. Persons whose sex was reported as transgender are included in the totals.

National Enteric Pathogens Surveillance System

The National Enteric Pathogens Surveillance System (NEPSS) collects, analyses and disseminates data on human enteric bacterial infections diagnosed in Australia. These pathogens include Salmonella, E. coli, Vibrio, Yersinia, Plesiomonas, Aeromonas and Campylobacter. Communicable Diseases Intelligence quarterly reports include only Salmonella. Data are based on reports to NEPSS from Australian laboratories of laboratory-confirmed human infection with Salmonella. Salmonella are identified to the level of serovar and, if applicable, phagetype. Infections apparently acquired overseas are included. Multiple isolations of a single Salmonella serovar/phage-type from one or more body sites during the same episode of illness are counted once only. The date of the case is the date the primary diagnostic laboratory isolated a Salmonella from the clinical sample. Note that the historical quarterly mean counts should be interpreted with caution, and are affected by surveillance artefacts such as newly recognised (such as S. Typhimurium 197 and S. Typhimurium U290) and incompletely typed Salmonella.

Reported by Joan Powling (NEPSS Co-ordinator) and Mark Veitch (Public Health Physician), Microbiological Diagnostic Unit—Public Health Laboratory, Department of Microbiology and Immunology, University of Melbourne. NEPSS can be contacted at the above address or by telephone: +61 3 8344 5701, or facsimile: +61 3 9625 2689.

Reports to the National Enteric Pathogens Surveillance System of Salmonella infection for the period 1 July to 30 September 2004 are included in Tables 12 and 13. Data include cases reported and entered by 28 October 2004. Counts are preliminary, and subject to adjustment after completion of typing and reporting of further cases to NEPSS. For more information about NEPSS see Surveillance systems reported in CDI, published in Commun Dis Intell 2004;28:101.

Reporting period 1 July to 30 September 2004

The total number of reports to NEPSS of human Salmonella infection declined to 1,156 in the third quarter of 2004, 42 per cent fewer than in second quarter of 2004 (Table 12) but 19 per cent more than the final count for the third quarter of 2004. Case counts to 28 October 2004 are expected to comprise more than 95 per cent of the final counts for the quarter.

During the third quarter of 2004, the 25 most common Salmonella types in Australia accounted for 669 cases, 58 per cent of all reported human Salmonella infections (Table 13).

Eighteen of the 25 most common Salmonella infections in the third quarter of 2004 were among the 25 most commonly reported in the previous quarter.

Reports of common salmonellae with counts well above historical averages include S. Typhimurium phage type 197 (in the eastern mainland states), S. Virchow phage type 8 (particularly in Queensland and New South Wales), and S. Stanley (particularly in Victoria). Counts of several typically overseasacquired phage types of S. Enteritidis were also elevated. While still among the more common salmonellae, reports of S. Typhimurium phage type 170/108 declined in number and relative prominence.

We thank scientists, diagnostic and reference laboratories, State and Territory health departments, and the Australian Government Department of Health and Ageing for their contributions to NEPSS.

Acknowledgement: Thanks to contributing laboratories and scientists.

Table 12. Reports to the National Enteric Pathogens Surveillance System of Salmonella isolatedfrom humans during the period 1 July to 30 September 2004, as reported to 28 October 2004

	ACT	NSW	NT	Qld	SA	Tas	Vic	WA	Australia
Total all Salmonella for quarter	17	263	68	355	69	11	238	135	1,156
Total contributing Salmonella types	12	91	29	102	33	10	88	65	209

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			ACT	NSN	NT	QLD	SA	Tas	Vic	WA		quarter		
	~	S Typhimurium 135	0	20	0	31	4	0	20	26	101	27	450	559
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	9	S Typhimurium 170	4	13	0	7	0	~	10	0	35	19	413	340
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9S Stanley11414101142613593610S Enteritidis Ga03031118102632501611S Typhimurium RDNC0130311183250363613S Aberdeen052111012632503614S Weltwroden05211301284616S Weltwroden0328011916716S Weltwroden0111111916717S Typhimurium 1260111111916718S Typhimurium 1201111111916718S Typhimurium 1201111111916718S Typhimurium 1201111111111111117S Typhimurium 12001111111111111111111111111	Ø	S Chester	0	7	4	Ø	5	0	0	က	27	20	156	174
	0	S Stanley	~	4	~	4	~	0	5	4	26	13	59	35
11S Typhimurium RDNC0130330502415845612S Infantis05211103222111316613S Aberdeen0521130121916714S Weltevreden032211301916715S Weltevreden032521142911916716S Weltevreden0101111916716S Weltevreden011111916716S Weltevreden01111916716S Typhimurium 126011111916717S Typhimurium 1260111111916718S Ball0111111191675818S Typhimurium 1260010111191675818S Typhimurium 120010011111120S Typhimurium 1203011111111<	10	S Enteritidis 6a	0	с	0	က	~	~	œ	10	26	3.2	50	15
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14S Weltevreden03280051197593415S Muenchen0425201418169010216S Entertitidis 4b010111418169010217S Typhimurium 1260101111419585618S Typhimurium 12600140011419585618S Typhimurium 12600140011419585619S Typhimurium 1200140011419585620S Litchfield0010011419585621S Entertitidis 1001000134.32027422S Entertitidis 1001000134.32027423S Agona11111111366111113661423S Agona11111111113671524S Typhimurium mutypable0201011111111 </td <td>13</td> <td>S Aberdeen</td> <td>0</td> <td>2</td> <td>~</td> <td>13</td> <td>0</td> <td>~</td> <td>2</td> <td>0</td> <td>19</td> <td>1</td> <td>91</td> <td>67</td>	13	S Aberdeen	0	2	~	13	0	~	2	0	19	1	91	67
15SMuenchen0425201418169010316SEnteritidis 4b0101011421844291117S Typhimurium 126050111121844291118S Typhimurium 1260501400141958585619S Typhimurium 1200140014106495619S Typhimurium 120014001419585810S Typhimurium 120014001419585820S Litchfield0140014101419585821S Enteritidis 103013141958343622S Agona111111126321123S Agona111111111121421423S Agona1111111111111111111111111111111 </td <td>14</td> <td>S Weltevreden</td> <td>0</td> <td>c</td> <td>2</td> <td>œ</td> <td>0</td> <td>0</td> <td>5</td> <td>-</td> <td>19</td> <td>7</td> <td>59</td> <td>34</td>	14	S Weltevreden	0	c	2	œ	0	0	5	-	19	7	59	34
16S Enteritidis 4b0101001142184.4291117S Typhimurium 126050111111419585618S Ball00140014001419585619S Typhimurium 12600140014061419585610S Typhimurium 120014001401419585620S Typhimurium 1200140011419585621S Typhimurium 120544100134.32027421S Enteritidis 10544100133.6343622S Enteritidis 10601000133.6343623S Enteritidis 10601000133.6343623S Enteritidis 111111112141916321523S Agona111111111111111216321624S Typhimurium	15	S Muenchen	0	4	2	5	2	0	~	4	18	16	06	109
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18 SBall 0 14 0 14 0 14 6 49 38 19 STyphimurum 12 0 9 0 1 0 14 6 49 38 20 STyphimurum 12 0 9 0 1 0 13 4.3 202 74 20 SLitchfield 0 5 4 4 0 0 13 4.3 202 74 21 SEnteritidis 1 0 5 4 4 0 0 1 12 12 12 12 12 12 12 12 16 32 55 56 16 32 15 16 32 15 12 12 12 12 12 12 16 32 15 16 32 15 15 16 32 15 15 16 32 15 12 12 16 16 16<	17	S Typhimurium 126	0	5	0	-	-	0	9	-	14	19	58	55
19S Typhimurum 120901030134.32027420S Litchfield054400133.6343621S Entertitidis 1030712126321222S Entertitidis 1060102312121623S Agona1111102312121624S Typhimurium untypable0402011111221125S Hvittingfoss020710191177	18	S Ball	0	0	14	0	0	0	0	0	14	9	49	39
20 SLitchfield 0 5 4 4 0 0 13 3.6 34 30 21 SEnteritidis 1 0 3 0 2 3 1 12 6 32 12 22 SEnteritidis 1b 0 6 0 1 0 2 3 12 12 6 32 12 23 SAgona 1 1 1 1 1 1 13 63 55 24 STyphimurium untypable 0 4 1 0 1 11 11 11 22 14 25 SHvittingfoss 0 2 0 7 1 0 11 9 117 7	19	S Typhimurium 12	0	0	0	~	0	0	ო	0	13	4.3	202	74
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25 SHvittingfoss 0 2 0 7 1 0 1 0 11 9 117 73	24	S Typhimurium untypable	0	4	0	2	0	0	4	-	11	11	22	18
	25	S Hvittingfoss	0	2	0	7	-	0	~	0	11	6	117	73