



WEEKLY EPIDEMIOLOGICAL REPORT

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Ministry of Health

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Enteric Fever (Part I)

This is the first in a series of two articles on Enteric Fever

Background

Enteric fever is a systemic infection caused by the human-adapted pathogens *Salmonella enterica* serotype Typhi (*S. Typhi*) and *S. enterica* serotype Paratyphi (*S. Paratyphi*) A, B and C. These organisms are important causes of febrile illness in crowded populations with inadequate sanitation that are exposed to unsafe water and food and also pose a risk to travelers visiting Enteric fever endemic regions.

Epidemiology

Burden of illness and death

In 2000, typhoid fever caused an estimated 21.7 million illnesses and 217,000 deaths, and paratyphoid fever caused an estimated 5.4 million illnesses worldwide. Infants, children, and adolescents in south-central and South-eastern Asia experience the greatest burden of illness. Typhoid and paratyphoid fever most often present as clinically similar acute febrile illnesses and accurate diagnosis relies on laboratory confirmation. Bone marrow culture remains the gold standard diagnostic test for enteric fever. Efforts to develop serologic methods for the diagnosis of typhoid fever that improve on the poor performance of the Widal test still suffer from substantial limitations of both sensitivity and specificity. Serological approaches to the diagnosis of *S. Paratyphi* A, B, and C have been developed but have not been evaluated or adapted for field use. Consequently, blood culture, a less sensitive method than bone marrow culture, is often the practical first choice test for both patient diagnosis and epidemiologic evaluation of *S. Typhi* and *S. Paratyphi* burden. However, most enteric fever occurs in low and middle-income countries where blood cultures are often unavailable, unaffordable or inconsistently applied. The

most robust approach to the measurement of incidence of typhoid and paratyphoid fever is by regular, community-wide household visits to identify persons with febrile illness from whom blood samples for culture confirmation may be obtained. Alternatively, the results of surveys of health-seeking behaviour and sentinel health care facility-based surveillance may be combined to estimate incidence. Because of the limited availability of blood culture services and the logistic challenges of enteric fever surveillance techniques capable of measuring disease incidence, the burden of typhoid and paratyphoid fever is poorly characterized in much of the world, particularly in sub-Saharan Africa. Furthermore, accurate estimates of rates of complications and death at the population level are not available. To reduce gaps in the current understanding of typhoid fever incidence, complications and case-fatality rate, large population-based studies using blood culture confirmation of cases are needed in representative sites, especially in low and medium human development index countries outside Asia.

Epidemiologic trends

Despite the limitations of currently available epidemiologic data, a number of recent trends in enteric disease epidemiology have emerged in the African, Asian and Latin American regions. In sub-Saharan Africa, where the burden of enteric fever is the least well characterized, hospital-based studies indicate that non-Typhi serotypes of *Salmonella*, particularly *S. enterica* serotype Enteritidis and *S. enterica* serotype Typhimurium, greatly outnumber *S. Typhi* and *S. Paratyphi* as causes of bloodstream infection. Nonetheless, outbreaks of typhoid fever are frequently reported from sub-Saharan Africa, often with large numbers of patients presenting with intestinal perforations leaving open important questions about the epidemiology of enteric fever in the region. In Asia, a large

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population-based prospective study using standardized surveillance methods estimated typhoid fever incidence in China, India, Indonesia, Pakistan and Vietnam to gather data for typhoid fever vaccine policy. This study confirmed the high incidence of typhoid fever in the region, particularly among children and adolescents, but also demonstrated that substantial variation in incidence occurs between surveillance sites in the same region. Simultaneously, *S. Paratyphi A* was responsible for a growing proportion of enteric fever in a number of Asian countries, sometimes accounting for 50% of *Salmonella* bloodstream isolates among patients with enteric fever. This trend raises important concerns about the impact of typhoid fever vaccine on enteric fever rates.

Prevention and Control Strategies

Contaminated water and food are important vehicles for transmission of typhoid fever. Historical surveillance data suggest that enteric fever was endemic in Western Europe and North America and this rate decreased in parallel with the introduction of water treatment facilities to municipal water supply schemes, pasteurization of dairy products and the exclusion of human faeces from food production. At present, enteric fever prevention focuses on improving sanitation, ensuring the safety of food and water supplies, identification and treatment of chronic carriers of *S. Typhi* and use of typhoid vaccines to reduce the susceptibility of hosts to infection.

Non-vaccine measures

Extending the benefits of improved sanitation and the availability of safe water and food that was achieved in industrialized countries a century ago to low and middle-income countries has proved to be a challenge. United Nations Millennium Development Goal 7 sets a target to halve, by 2015, the proportion of the population without sustainable access to safe drinking water and basic sanitation. Recent evidence suggests that interventions to improve the quality of drinking water may be relatively more important for the prevention of enteric infection relative to sanitation measures than was previously thought. Although centrally treated reticulated water for all is an important goal, a growing body of research suggests that improving water quality at the household level, as well as at the source, can significantly reduce diarrhoea. Although not formally evaluated with enteric fever as an outcome, it is likely that interventions that reduce the rate of diarrhoeal diseases transmitted through contaminated water, food and poor hygiene would have similar effects on rates of enteric fever.

The identification and treatment of *S. Typhi* carriers, particularly those involved with food production, has proven to be an important strategy for the control of typhoid fever in low-incidence settings.

Source - Global Trends in Typhoid and Paratyphoid Fever

Available from <http://cid.oxfordjournals.org/content/50/2/241.full>

Compiled by
Dr. Madhava Gunasekera of the Epidemiology Unit

Invasive Bacterial Disease surveillance in Sentinel Sites- 3rd quarter 2013

No. of suspected meningitis cases	26
No. of probable meningitis cases	4
Percentage (%) of CSF samples tested positive for organisms	0%
No. of children who met the pneumonia case definition	117
Percentage (%) of Pneumonia cases with positive blood cultures	0%
No. of sepsis cases	31
Percentage (%) of Sepsis cases with positive blood cultures	0%
Source-LRH, Epidemiology Unit	

Rota virus surveillance in Sentinel Sites - 3rd quarter 2013

Number of acute diarrhoea hospitalizations in children <5 years	362
Number of stool specimen collected	135
Number of stool specimen tested positive for rotavirus	42
Percentage (%) of stool specimen tested positive for rotavirus	33 %
Source-MRI, Epidemiology Unit	

**Table 3 : Water Quality Surveillance
 Number of microbiological water samples - Sept / 2013**

District	MOH areas	No: Expected *	No: Received
Colombo	12	72	12
Gampaha	15	90	34
Kalutara	12	72	NR
Kalutara NISH	2	12	19
Kandy	23	138	NR
Matale	12	72	36
Nuwara Eliya	13	78	03
Galle	19	114	71
Matara	17	102	NR
Hambantota	12	72	55
Jaffna	11	66	47
Kilinochchi	4	24	88
Manner	5	30	28
Vavuniya	4	24	33
Mullatvu	4	24	18
Batticaloa	14	84	55
Ampara	7	42	NR
Trincomalee	11	66	19
Kurunegala	23	138	NR
Puttalam	9	84	7
Anuradhapura	19	114	43
Polonnaruwa	7	42	27
Badulla	15	90	53
Moneragala	11	66	70
Rathnapura	18	108	NR
Kegalle	11	66	29
Kalmunai	13	78	NR

* No of samples expected (6 / MOH area / Month)
 NR = Return not received

Table 4: Selected notifiable diseases reported by Medical Officers of Health 12th - 18th October (42nd Week)

RDHS	Dengue Fever		Dysentery		Encephalitis		E Fever		F Poisoning		Leptospirosis		T Fever		V Hepatitis		H Rabies		Chickenpox		Meningitis		Leishmani-			WRCD %	
	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	T*	C**	
Colombo	48	7868	1	173	0	17	1	129	0	53	1	181	0	7	0	71	0	1	6	371	0	58	0	0	62	38	
Gampaha	40	3023	2	175	2	18	0	46	0	31	5	347	0	18	0	167	0	0	7	149	1	83	0	5	47	53	
Kalutara	22	1502	2	156	0	20	3	74	0	23	5	351	0	5	1	24	0	0	3	232	0	64	0	0	85	15	
Kandy	11	1528	0	134	0	11	0	25	0	10	0	68	0	94	0	104	0	0	1	121	0	14	0	4	74	26	
Matale	4	404	0	84	0	4	0	24	1	8	1	58	0	4	1	46	0	0	0	43	1	34	0	11	62	38	
NuwaraEliya	2	217	1	142	0	2	1	14	0	217	2	26	1	58	2	22	0	0	0	107	0	12	0	0	85	15	
Galle	3	751	3	103	0	19	0	6	8	89	0	194	1	53	1	14	0	2	6	286	0	45	0	0	84	16	
Hambantota	0	283	1	52	0	3	0	15	0	32	1	160	0	62	0	84	0	0	3	95	1	46	1	279	67	33	
Matara	3	409	3	74	1	13	0	28	0	27	0	136	2	80	2	138	0	2	2	237	0	70	5	83	94	6	
Jaffna	9	603	18	346	0	10	3	302	0	96	0	8	2	331	0	17	0	1	2	137	1	55	0	0	75	25	
Kilinochchi	1	58	0	35	0	0	0	14	0	5	0	9	0	16	0	0	0	2	0	2	0	7	0	11	50	50	
Mannar	0	65	0	72	0	3	3	62	0	36	0	14	0	19	0	2	0	0	0	11	0	5	0	4	80	20	
Vavuniya	1	65	0	55	0	13	0	12	0	20	0	50	1	3	0	3	0	2	0	22	0	33	0	10	100	0	
Mullaitivu	2	112	0	19	0	2	1	9	0	43	0	37	1	7	0	1	0	2	0	8	0	6	0	14	80	20	
Batticaloa	3	507	5	278	0	5	0	10	0	73	0	33	0	2	0	14	0	3	1	41	0	7	0	0	64	36	
Ampara	0	170	4	153	0	1	0	5	0	10	0	33	0	1	0	8	0	0	0	81	1	18	0	3	43	57	
Trincomalee	0	183	0	60	0	3	0	6	0	3	0	59	0	15	0	3	0	1	0	37	0	4	0	28	67	33	
Kurunegala	15	2524	1	159	0	36	0	38	0	26	1	286	0	41	0	52	0	1	3	323	1	95	4	48	78	22	
Puttalam	2	806	5	74	0	7	0	16	0	36	0	42	0	12	0	7	0	1	0	78	0	32	1	9	62	38	
Anuradhapur	2	465	4	100	0	16	0	3	19	59	1	303	0	23	0	25	0	2	0	159	2	92	10	368	68	32	
Polonnaruwa	2	396	3	74	1	2	0	14	2	64	2	160	0	3	0	29	0	2	0	121	0	17	3	148	86	14	
Badulla	5	450	5	179	0	5	1	18	1	11	0	54	4	81	0	44	0	0	3	116	2	64	0	7	82	18	
Monaragala	5	222	1	106	0	4	0	23	0	25	0	196	0	57	3	159	0	1	0	48	0	23	0	10	73	27	
Ratnapura	4	1575	6	340	0	83	0	38	0	17	5	320	1	65	17	437	0	1	4	156	1	76	0	13	72	28	
Kegalle	16	985	3	114	0	16	0	28	0	11	7	202	0	71	6	208	0	0	7	291	0	99	0	2	100	0	
Kalmune	0	492	0	138	0	2	0	3	0	117	1	9	0	2	0	5	0	0	0	83	0	9	0	1	38	62	
SRI LANKA	200	25663	68	3395	04	315	13	962	31	1142	32	3336	19	1130	39	1684	00	24	48	335	11	1068	24	1058	72	28	

Source: Weekly Returns of Communicable Diseases (WRCD).

*T=Illness refers to returns received on or before 18th October, 2013. Total number of reporting units 339. Number of reporting units data provided for the current week:244 C** Completeness

A = Cases reported during the current week. B = Cumulative cases for the year. H Rabies = Human Rabies, E Fever = Enteric Fever, F Poison = Typhus Fever, V Hepatitis = Viral Hepatitis

Table 1: Vaccine-Preventable Diseases & AFP **12th - 18th October 2013 (42nd Week)**

Disease	No. of Cases by Province									Number of cases during current week in 2013	Number of cases during same week in 2012	Total number of cases to date in 2013	Total number of cases to date in 2012	Difference between the number of cases to date in 2013 & 2012
	W	C	S	N	E	NW	NC	U	Sab					
AFP*	00	00	00	00	00	00	00	01	00	01	01	76	65	+ 16.9%
Diphtheria	00	00	00	00	00	00	00	00	00	-	-	-	-	-
Mumps	01	01	02	00	02	04	00	00	02	12	33	1274	3904	- 67.3 %
Measles	09	01	08	00	00	01	03	01	07	30	01	3215	52	+ 6082.7 %
Rubella	00	01	00	00	00	00	00	00	00	01	-	26	-	-
CRS**	00	00	00	00	00	00	00	00	00	00	-	06	-	-
Tetanus	00	00	00	00	00	00	00	00	00	00	01	19	10	+ 90.0 %
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	-	00	-	-
Japanese Encephalitis	01	00	00	00	00	00	00	00	00	00	-	68	-	-
Whooping Cough	01	00	00	00	00	00	00	00	00	01	00	70	87	- 19.5 %
Tuberculosis	02	00	02	05	00	06	05	00	00	20	327	6617	7150	- 04.5 %

Key to Table 1 & 2

Provinces: W: Western, C: Central, S: Southern, N: North, E: East, NC: North Central, NW: North Western, U: Uva, Sab: Sabaragamuwa.
RDHS Divisions: CB: Colombo, GM: Gampaha, KL: Kalutara, KD: Kandy, ML: Matale, NE: Nuwara Eliya, GL: Galle, HB: Hambantota, MT: Matara, JF: Jaffna, KN: Killinochchi, MN: Mannar, VA: Vavuniya, MU: Mullaitivu, BT: Batticaloa, AM: Ampara, TR: Trincomalee, KM: Kalmunai, KR: Kurunegala, PU: Puttalam, AP: Anuradhapura, PO: Polonnaruwa, BD: Badulla, MO: Moneragala, RP: Ratnapura, KG: Kegalle.

Data Sources:

Weekly Return of Communicable Diseases: Diphtheria, Measles, Tetanus, Neonatal Tetanus, Whooping Cough, Chickenpox, Meningitis, Mumps., Rubella, CRS,

Special Surveillance: AFP* (Acute Flaccid Paralysis), Japanese Encephalitis

CRS** =Congenital Rubella Syndrome

AFP and all clinically confirmed Vaccine Preventable Diseases except Tuberculosis and Mumps should be investigated by the MOH

Influenza Surveillance in Sentinel Hospitals - ILI & SARI								
Month	Human				Animal			
	No Received	Infl A untyped	Infl B	A(H1N1)pdm09	A(H3N2)	Pooled samples	Serum Samples	Positives
September	233	9	9	26	8	190	460	0

Source: Medical Research Institute & Veterinary Research Institute

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ON STATE SERVICE

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