



# WEEKLY EPIDEMIOLOGICAL REPORT

# A publication of the Epidemiology Unit Ministry of Health

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## Childhood Tuberculosis - Challenges and opportunities in improving case finding

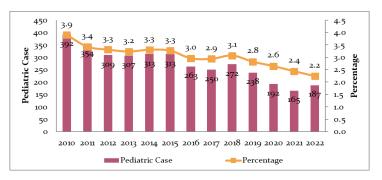
Despite taking great public health steps to control tuberculosis (TB), it remains an enormous public health challenge worldwide. Tuberculosis has been the commonest cause of death from a single infectious pathogen until 2020 when it was overtaken by COVID-19.

Sri Lanka is facing a greater challenge in TB case finding. The country usually notifies 8,000 to 9,000 TB patients, whereas the WHO estimated number is 14,000 patients, leading to the situation of missing around 5000 to 6000 TB cases annually. This is applicable to paediatric TB as well. The proportion of all forms of TB notifications among children under 0-14 years of age remained below the expected range over the years (figure 1). When compared with global estimates for low-middle income countries which should be 5-15% of total TB cases, a significant under-notification and under-reporting of child TB cases is noted in the country.

As childhood tuberculosis (TB) reflects recent transmission, its burden provides an

accurate measure of the level of TB control in a particular community. In Sri Lanka, the TB age-specific curve has been shifting towards the right over the years, emphasizing less community transmission and more vulnerability among older ages. However, considering the WHO target, the country is committed to finding out minimum of 600 childhood TB cases per year, as per the country target set in the National Strategic Plan 2021-2025. Under this backdrop, it is worth enlightening the healthcare workers in both curative and preventive health sectors on the existing challenges and opportunities in order to improve paediatric TB case finding.

Diagnosing Childhood TB carries inbuilt challenges. Globally, childhood TB is underreported due to its paucibacillary nature and the difficulty of confirming the diagnosis. The clinical and radiographic manifestations are less specific in children compared to adults and are often confused with bacterial pneumonia. Microbiologic confirmation of disease



 $Figure\ 1.\ Trend\ of\ total\ childhood\ TB\ cases\ and\ proportionate\ contribution\ to\ total\ cases\ 2010-2022$ 

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is limited by the paucibacillary nature of the disease. Collection of a good quality sputum sample from children is often a challenge and recommends induced sputum or gastric lavage which may not be carried out in all settings.

Recent TB reviews had emphasized that the efforts on diagnosing paediatric TB in the country is sub-optimum. The gaps imply a low investigation rate among children less than 5 years old, sub-optimal capacity and understanding of the childhood TB diagnostic algorithm and policies, limited capacity among health care providers to induce sputum production in children, possible low suspicion and capacity to diagnose pediatric TB among medical officers at Outpatient settings and clinics, and weak referral linkages to pediatric facilities for further evaluation, are some of these limitations leading to low childhood TB detection. On the other hand, the opportunities for momentum to improve childhood diagnosis are immense in the country. The National Programme for Tuberculosis Control and Chest Diseases (NPTCCD) has developed a paediatric TB algorithm and made both the curative and preventive sectors aware of this including the possible referral pathways (Figure 2).

In contrast to the adult TB diagnostic algorithm, the paediatric TB algorithm contains many criteria beyond respiratory symptoms. The existence of different stakeholders to gear the effective diagnosis among suspected children is what we are lacking is continuous momentum and collab an opportunity. oration among stakeholders. All healthcare workers who encounter the paediatric population, Paediatricians, Primary Care Physicians in the government and private sectors, Medical Officers of Health, Public Health Midwives, Clinical Nutritionists, Child Developmental Officers in the estates etc should suspect paediatric TB when they encounter a child with stipulated criteria in the algorithm.

Hence, to improve childhood TB diagnosis, clinicians should think beyond the traditional framework of diagnosing TB when it comes to childhood Tuberculosis. It is much more important to assess the child as a whole including the risk of exposure, and epidemiological background in relation to TB to facilitate TB diagnosis among children. The other healthcare workers should adhere to the given algorithm and referral pathways to enhance diagnosis. Under the given economic crisis, there is a tendency to have nutritionally driven immune deficiency which could lead to a higher number of paediatric TB in the community than expected. Therefore, it is a timely need to "Think TB" and keep TB high on the list when examining children in the institutional and field settings as a continuous drive to eliminate childhood tuberculosis is a must to end TB by 2035.

# Children with signs and symptoms suggestive of TB<sup>1</sup> and/or Children with risk nators for active TB<sup>2</sup> History & examination including anthropometric assessment Refer for Expert opinion (Samples to be sent for Xpert MTB/RIF Positive S. Sputum for Xpert MTB/RIF Positiv

NATIONAL PROGRAMME FOR TUBERCULOSIS CONTROL AND CHEST DISEASES

### Compiled by:

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Table 1: Selected notifiable diseases reported by Medical Officers of Health 09th - 15th Mar 2024 (11th Week)

ıab	able 1: Selected notifiable diseases reported by Medical Officers of Health 1941-1541 Mar 2024 (1141 Week)																												
G	* *	100	66	100	100	100	100	100	100	100	93	100	100	100	100	100	100	100	98	100	100	100	100	100	100	100	100	99	
WRCD	<u>*</u>	92	93	93	100	100	100	92	100	100	93	100	100	100	64	100	100	100	85	54	96	100	94	82	100	100	85	93	
ulosis	В	441	247	152	144	26	29	102	34	18	46	7	16	2	∞	23	53	17	124	43	28	17	48	20	63	71	32	1882	
Tuberculosis	∢	54	27	13	2	7	6	6	ω	9	10	0	_	0	က	9	4	7	6	_	7	0	9	0	10	ω	0	205	
ania-	В	0	9	0	12	65	0	က	127	28	0	0	~	က	4	~	2	<sub>∞</sub>	134	2	212	114	7	29	99	12	0	862	
Leishmania-	⋖	0	~	0	0	~	0	0	15	_	0	0	0	0	~	0	2	2	15	_	22	15	0	12	~	~	0	90	
Meningitis	В	10	28	20	4	4	က	23	7	36	2	2	2	9	0	18	16	3	29	13	16	7	00	37	32	19	က	393	
Meni	⋖	0	2	က	0	~	0	က	7	7	0	0	0	~	0	~	~	0	00	_	0	~	~	က	က	7	0	35	
Chickenpox	В	92	62	125	148	19	45	126	64	99	69	~	4	7	2	4	37	10	66	33	42	41	9/	23	79	166	38	1488	
Chick	∢	12	0	∞	∞	4	9	20	9	10	9	0	0	0	0	0	0	က	0	2	7	2	12	7	15		5	157	
H. Rabiies	В	0	0	0	0	0	0	_	0	0	_	0	0	0	0	0	0	0	2	0	0	0	0	0	_	0	0	5	
H. Ra	⋖	0	0	0	0	0	0	~	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	_	
Hep.	В	3	_	2	2	~	က	4	2	0	က	0	0	4	0	2	က	0	2	0	9	~	5	7	00	2	_	71	
Viral	4	0	0	0	0	0	7	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	<u></u>	~	~	2	
Typhus F.	В	0	2	က	5	0	12	34	13	9	317	9	5	~	ω	~	_	7	13	5	18	~	7	13	ω	5	_	492	
Тур	⋖	0	0	~	2	0	_	4	0	0	24	0	0	0	0	0	0	2	_	_	4	0	~	0	0	0	0	41	
Leptospirosis	Ф	84	149	155	69	38	65	225	214	92	12	6	16	47	45	19	96	80	214	108	165	66	168	351	403	166	32	3121	
Lepto	⋖	7	22	13	7	0	3	20	6	3	0	0	~	_	2	_	4	10	13	3	10	4	19	20	31	24	2	233	
F. Poisoning	В	3	0	က	3	4	6	16	0	4	15	_	0	~	2	10	7	~	339	0	2	2	6	0	3	3	0	437	
F. Po	⋖	0	0	က	0	0	0	0	0	0	0	0	0	0	0	_	0	0	3	0	0	0	_	0	_	_	0	10	
Fever	В	2	2	က	0	~	_	~	0	0	2	0	_	0	0	~	0	~	0	0	0	0	0	_	0	0	0	16	
Ë	⋖	0	0	0	0	0	0	0	0	0	_	0	0	0	0	0	0	0	0	0	0	0	0	_	0	0	0	2	
Encephalitis	В	~	4	0	0	0	2	9	0	2	~	0	0	0	0	5	_	0	7	_	0	0	_	0	~	2	0	34	
Ence	٧	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	3	0	0	0	0	0	_	0	0	4	
Dysentery	В	5	5	0	5	~	19	4	5	2	21	2	0	0	က	40	12	9	5	_	4	9	ω	4	19	က	∞	207	
Dys	⋖	0	0	0	0	0	2	0	~	0	0	_	0	0	0	2	~	~	0	_	_	0	~	0	~	0	0	12	
Dengue Fever	В	3739	1521	958	1424	286	156	911	359	313	4615	249	166	120	167	928	121	356	1104	553	431	149	459	320	707	810	455	21377	
Dengu	⋖	207	90	91	29	ω	4	28	27	23	66	∞	~	2	7	22	13	23	20	0	33	10	13	19	99	44	22	1004	
RDHS		Colombo	Gampaha	Kalutara	Kandy	Matale	Nuwara Eliya	Galle	Hambantota	Matara	Jaffna	Kilinochchi	Mannar	Vavuniya	Mullaitivu	Batticaloa	Ampara	Trincomalee	Kurunegala	Puttalam	Anuradhapura	Polonnaruwa	Badulla	Monaragala	Ratnapura	Kegalle	Kalmunai	SRILANKA	

Source: Weekly Returns of Communicable Diseases (esurvillance.epid.gov.Ik). T=Timeliness refers to returns received on or before 15th Mar, 2024 Total number of reporting units 358 Number of reporting units data provided for the current week. B = Cumulative cases for the year.

# Table 2: Vaccine-Preventable Diseases & AFP

09th - 15th Mar 2024 (11th Week)

Disease	No. of Cases by Province										Number of cases during same	Total number of cases to date in	Total num- ber of cases to date in	Difference between the number of cases to date	
	W	С	S	N	Е	NW	NC	U	Sab	week in 2024	week in 2023	2024	2023	in 2024 & 2023	
AFP*	00	00	00	00	00	01	00	00	00	02	01	16	19	-15.7 %	
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %	
Mumps	01	00	00	00	01	00	00	01	00	03	02	64	44	45.4 %	
Measles	01	00	00	00	01	01	00	00	00	03	00	151	00	0 %	
Rubella	00	00	00	00	00	00	00	00	00	00	00	01	00	0 %	
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %	
Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	01	-100 %	
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %	
Japanese Encephalitis	00	00	00	00	00	00	00	00	00	00	00	01	00	0 %	
Whooping Cough	00	00	00	00	00	00	00	00	00	00	01	01	02	-50 %	

# 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

NA = Not Available

Influenza 2024													
Month		Human	Animal										
Month	No Total	No Positive	Infl A	Infl B	Pooled samples	Serum Samples	Positives						
February	435	60	46	14	2251	750	0						
Duration: from 01st of February to 28th of February													
Source: MRI , TH Karapitiya, National Cancer Institute													

Comments and contributions for publication in the WER Sri Lanka are welcome. However, the editor reserves the right to accept or reject items for publication. All correspondence should be mailed to The Editor, WER Sri Lanka, Epidemiological Unit, P.O. Box 1567, Colombo or sent by E-mail to chepid@sltnet.lk. Prior approval should be obtained from the Epidemiology Unit before publishing data in this publication

# **ON STATE SERVICE**

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