



# WEEKLY EPIDEMIOLOGICAL REPORT

# A publication of the Epidemiology Unit

Ministry of Health

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# Implementation of prophylactic treatment for tuberculosis in Sri Lanka

This is the second article of two in a series on "Implementation of prophylactic treatment for tuberculosis in Sri Lanka."

# The target population for Sri Lanka

For many years, Sri Lanka has been giving TPT to children under five years who are close contacts of bacteriologically confirmed PTB patients and PLHIV patients. With the expansion of the LTBI implementation, other high-risk groups recognized by WHO were also included in LTBI guidelines for Sri Lanka.

The implementation is carried out in a phasedout manner within the country according to feasibility. Initially, during 2022, all children up to 15 years and close contacts of TB patients above 50 years, were included for TPT and during 2023, this was further expanded to all close contacts among PTB patients. TPT for PLHIV continued, while certain high-risk groups such as organ transplant and dialysis categories were also addressed for TPT according to feasibility. Currently, the programme is in the process of strengthening the TPT implementation in all above stated categories.

tecting TB disease. Once the active disease is ruled out, they are tested for TBI. Children under 5 years are given TPT after excluding the disease without testing for infection. All other high-risk categories need to undergo testing for infection before initiating TPT.

Either interferon-gamma release assays (IGRAs) or tuberculin skin test (TST) is used to diagnose LTBI. The Mantoux test, the method used in Sri Lanka, is a type of TST. In addition, the IGRA facility is also established in the Na-Tuberculosis Reference Laboratory (NTRL, Welisara) and will be available for clinically high-risk groups such as PLHIV, and patients on dialysis.

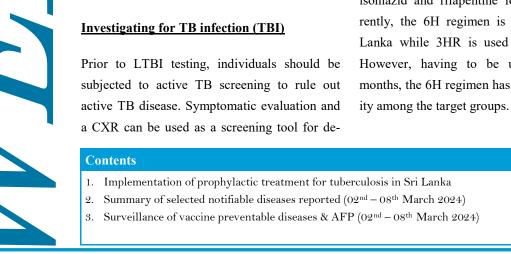
# **Tuberculosis Preventive Treatment**

In the prophylactic treatment for TBI, one or two main drugs used in the treatment of TB (Isoniazid (H), Rifampicin ®, Ethambutol €, Pyrazinamide (Z)) are used in smaller doses. The treatment regimens thus prioritized are 6H (isoniazid daily monotherapy for 6 months), 3HR (isoniazid and rifampicin daily for 3 months), and 3HP (once a week treatment with isoniazid and rifapentine for 3 months). Currently, the 6H regimen is widely used in Sri Lanka while 3HR is used to a lesser extent. However, having to be used daily for six months, the 6H regimen has reduced in popular-

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Irregular or inadequate treatment reduces the protective efficacy of the TPT regimen. Further, poor adherence or early cessation of TPT can potentially increase the risk of the individual developing TB including drug-resistant TB. It is known that the efficacy of TPT is greatest if at least 80% of the doses are taken within the duration of the recommended regimen.

Whilst currently the 6H treatment regimen is being used in the country, a more user-friendly 3HP regimen will be available within the country shortly. From a programme perspective, administering a shorter, weekly dose would greatly enhance the TPT completion rates in turn, reducing the TB caseload.

#### **Challenges**

When observing the chest clinic data on patient recruitment for TPT and follow-up has shown that retention in care has been a challenge. There are many challenges identified in relation to the diagnosis of TBI and initiation of TPT at the field level. Having to pay two visits for Mantoux testing (for administration and reading) is a drawback with many patients not following up with the second visit. Further, Mantoux testing for TB infection is currently carried out in limited healthcare settings, mainly in the district chest clinics. The main prophylactic regimen currently used for TPT is 6H which is a comparatively longer regimen which may lead to poor compliance among patients. Further, the lack of communication and counselling may lead to a higher loss to followup among the TPT population. Currently, the national programme is trying to address these barriers through the expansion of TST services within districts, the introduction of shorter regimens and capacity building among healthcare workers for counselling along with follow-up of recruited patients for TPT.

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&

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#### **References:**

- 1. National Guideline on Latent Tuberculosis Infection (LTBI) Management- 2021
- 2. GLOBAL TUBERCULOSIS 2023 REPORT <a href="https://iris.who.int/bitstream/">https://iris.who.int/bitstream/</a> <a href="handle/10665/373828/9789240083851-eng.pdf">handle/10665/373828/9789240083851-eng.pdf</a>? sequence=1
- Latent tuberculosis infection: Updated and consolidated guidelines for programmatic management- WHO https://www.who.int/publications/i/item/9789241550239
- 4. Update of WHO guidelines on the management of latent TB infection. https://www.who.int/news/item/07-01-2019-update-of-who-guidelines-on-the-management-of-latent-tb-infection

Table 1 : Water Quality Surveillance Number of microbiological water samples February 2023											
District	MOH areas	No: Expected *	No: Received								
Colombo	15	90	10								
Gampaha	15	90	NR								
Kalutara	12	72	85								
Kalutara NIHS	2	12	31								
Kandy	23	138	59								
Matale	13	78	2								
Nuwara Eliya	13	78	19								
Galle	20	120	109								
Matara	17	102	8								
Hambantota	12	72	NR								
Jaffna	12	72	175								
Kilinochchi	4	24	14								
Mannar	5	30	NR								
Vavuniya	4	24	53								
Mullatvu	5	30	7								
Batticaloa	14	84	0								
Ampara	7	42	0								
Trincomalee	11	66	NR								
Kurunegala	29	174	65								
Puttalam	13	78	NR								
Anuradhapura	19	114	2								
Polonnaruwa	7	42	0								
Badulla	16	96	0								
Moneragala	11	66	0								
Rathnapura	18	108	NR								
Kegalle	11	66	2								
Kalmunai	13	78	0								

NR = Return not received

Table 1: Selected notifiable diseases reported by Medical Officers of Health 02<sup>nd</sup> - 08<sup>th</sup> Mar 2024 (10<sup>th</sup> Week)

Table 1: Selected notifiable diseases reported by Medical Officers of Health 02 <sup>nd</sup> - 08 <sup>th</sup> Mar 2024 (10 <sup>th</sup> Week)														)															
WRCD	*5	100	66	100	100	100	100	66	100	100	93	100	100	100	100	100	100	100	98	98	100	100	66	100	100	100	100	66	
WR	<u>*</u>	100	93	92	100	92	92	92	100	100	100	100	100	100	100	100	71	100	100	91	96	100	93	91	92	100	100	96	
Tuberculosis	В	387	220	139	139	19	28	93	26	12	36	7	15	2	2	17	49	15	115	42	21	17	42	20	53	63	32	1677	
Tubero	⋖	42	27	œ	0	0	က	22	2	0	7	0	_	0	~	0	2	4	17	2	တ	0	2	7	0	2	7	159	
niasis	В	0	2	0	12	64	0	က	111	27	0	0	_	က	က	~	က	9	119	က	189	66	7	47	22	7	0	692	
Leishmaniasis	⋖	0	_	0	က	2	0	0	<u></u>	2	0	0	0	2	2	0	0	2	22	0	19	13	0	10	2	0	0	97	
	В	10	26	17	4	က	က	20	6	34	2	7	2	2	0	17	15	က	29		16	9	7	34	29	17	က	357	
Meningitis	٧	~	_	~	0	7	0	<del>-</del>	0	7	7	0	0	~	0	7	<del>-</del>	0	4	7	<del>-</del>	0	2	က	က	0	0	29	
Chickenpox	В	92	62	125	148	19	45	126	64	99	69	_	4	7	7	4	37	10	66	33	42	41	9/	23	79	166	38	1488	
Chick	⋖	12	6	00	∞	4	9	20	9	10	9	0	0	0	0	0	0	က	0	2	7	2	12	7	15		2	157	
Rabiies	В	0	0	0	0	0	0	0	0	0	~	0	0	0	0	0	0	0	2	0	0	0	0	0	<u></u>	0	0	4	
H.R.	⋖	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	0	0	
Il Hep.	В	က	_	5	2	_	_	4	2	0	က	0	0	4	0	5	က	0	2	0	9	~	5	7	7	4	0	99	
Viral	∢	0	0	0	~	0	0	0	0	0	0	0	0	~	0	~	0	0	0	0	2	0	0	0	0	0	0	5	
Typhus F.	В	0	2	2	က	0	1	29	13	9	293	9	2	_	00	_	_	5	12	4	4	_	9	13	00	2	_	450	
	∢	3 0	1	7	-	0	3	9	~	9	18	0	5 0	0	10	0	0	1	1	0	2	5 0	0	_	0	0	0	1 39	
Leptospirosis	В	7	127	142	. 62	38	62	203	203	∞	12		1,	46	45	18	06	. 70	201	105	155	6	149	331	372	142	30	2884	
	4	00	26	23	7	4	0	20	12		0	0	_	4	n	2		4	7	_	13	2	00	12	27	18	0	222	
F. Poisoning	В	က	0	0	က	4	0	16	0	4	15	_	0	~	2	6	7	_	336	0	2	2	00	0	2	2	0	427	
F. P.	⋖	0	0	0	0	0	0	0	0	2	00	0	0	7	0	9	_	0	_	0	0	0	က	0	0	2	0	24	
En. Fever	В	2	2	က	0	_	_	~	0	0	~	0	_	0	0	_	0	~	0	0	0	0	0	0	0	0	0	14	
	⋖	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	_	
Encephalitis	В	~	4	0	0	0	2	9	0	2	_	0	0	0	0	5	_	0	4	_	0	0	_	0	0	2	0	30	
Ence	∢	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	_	
Dysentery	В	5	5	6	5	~	17	4	4	2	21	_	0	0	က	38		2	5	0	က	9	7	4	18	က	00	195	
٥٨	∢	0	0	0	0	0	_	2	_	0	0	~	0	0	0	4	_	0	0	0	0	0	0	_	_	0	_	13	
Dengue Fever	В	3532	1431	867	1365	278	152	879	331	290	4516	241	165	115	162	873	106	333	1054	540	399	139	446	302	651	299	433	20366	
Deng	⋖	170	89	43	89	7	10	40	22	28	143	∞	4	2	2	29	0	13	29	14	42	<del>-</del>	7	17	38	34	27	975	
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

02<sup>nd</sup> - 08<sup>th</sup> Mar 2024 (10<sup>th</sup> Week)

Disease	No.	of Ca	ases	by P	rovin	ce		Number of cases during current	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	E	NW	NC	U	Sab	week in 2024	week in 2023	2024	2023	in 2024 & 2023
AFP*	00	00	00	00	00	00	00	00	00	00	01	14	18	-22.2 %
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %
Mumps	04	00	02	00	00	00	01	01	00	08	06	61	40	52.5 %
Measles	00	00	01	00	00	00	00	00	00	01	00	121	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

Take prophylaxis medications for leptospirosis during the paddy cultivation and harvesting seasons.

It is provided free by the MOH office / Public Health Inspectors.

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

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