



# WEEKLY EPIDEMIOLOGICAL REPORT

A publication of the Epidemiology Unit Ministry of Health & Indigenous Medical Services

231, de Saram Place, Colombo 01000, Sri Lanka
Tele: + 94 11 2695112, Fax: +94 11 2696583, E mail: epidunit@sltnet.lk
Epidemiologist: +94 11 2681548, E mail: chepid@sltnet.lk
Web: http://www.epid.gov.lk

Vol. 47 No. 34

15th- 21st Aug 2020

# **Neglected Tropical Diseases - Leishmaniasis**

Leishmaniasis is a protozoal disease that affects nearly one million people worldwide each year. It

is caused by a protozoal species known as Leishmania which is transmitted to humans through the bites of infected female phlebotomine sandflies, which feed on blood to produce eggs.

Major risk factors for Leishmaniasis include poor socio-economic conditions (housing, sanitation),



population mobility and climate

change. Cutaneous leishmaniasis is increasingly seen in Sri Lanka at present.



Figure - Phlebotomine sandfly

	Occurrence	Features
Visceral Leishmania- sis (VL) also known as Kala- Azar	50,000-90,000 new cases of VL/year  Brazil, East Africa and India are the most affected	Symptoms and signs include fever, loss of weight, hepato- megaly, spleno- megaly and anae- mia.
Cutaneous leish- maniasis (CL)	commonest form of leishmaniasis  600 000 to 1 million new cases occur worldwide annually.  About 95% of CL cases occur in the Americas, the Mediterranean basin, the Middle East and Central Asia	Causes skin le- sions, mainly ulcers, on ex- posed parts of the body, leaving life- long scars and serious disability or stigma.
Mucocutaneous leishmaniasis	Over 90% of mucocutaneous leishmaniasis cases occur in Bolivia (the Plurinational State of), Brazil, Ethiopia and Peru.	leads to partial or total destruction of mucous mem- branes of the nose, mouth and throat.

Figure- Cutaneous Leishmaniasis Visceral leishmaniasis is diagnosed based on clinical signs and parasitological, or serological tests (such as rapid diagnostic tests). In cutaneous and mucocutaneous leishmaniasis clinical features along with parasitological tests confirms the diagnosis.

There are three forms of Leishmaniasis as described below.

Prevention and control involve a multi-strategic response consisting of early diagnosis and treatment of cases, vector control, heightened surveillance and community mobilization.

Contents	Page
1. Leading Article - Neglected Tropical Diseases - Leishmaniasis	1
2. Summary of selected notifiable diseases reported (08th- 14th August 2020)	3
3. Surveillance of vaccine preventable diseases & AFP (08th- 14th August 2020)	4

## Leprosy (Hansen's Disease)

Leprosy is a slow growing disease caused by a rod-shaped acid-fast bacillus known as *Mycobacterium leprae*. The incubation period on average, is about 5 years. However, symptoms may take as long as 20 years to develop. Leprosy is transmitted by droplet infection during close contact with untreated infected persons. It mainly affects the skin, peripheral nerves, mucosa of the upper respiratory tract, and the eyes; and can cause permanent damage to these tissues. Leprosy is curable with multi-drug therapy.

Elimination of leprosy as public health problem (defined as a registered prevalence of less than 1 case per 10 000 population) was achieved globally in 2000. However, the path to eradication has been tedious, and WHO launched its "Global Leprosy Strategy 2016–2020: Accelerating towards a leprosy-free world" in 2016 in order to "go the last mile".

### Lymphatic filariasis (Elephantiatis)

Lymphatic filariasis or elephantiasis, is a chronic disfiguring disease caused by a family of roundworms (nematodes) known as Filariodidea. There are 3 species responsible for the disease:

- Wuchereria bancrofti, which is responsible for 90% of the cases
- Brugia malayi, which causes most of the remainder of the cases
- Brugia timori, which also causes the disease

Transmission to humans occurs following the introduction of the microfilaria (immature larvae) into the human tissue through infected mosquitoes. Microfilaria deposited on human skin by the mosquitoes enter the human body through a break in skin/mosquito bite migrate to the lymphatic system. Here, they grow into adult worms and reside in the lymphatic vessels, disrupting the normal function of the lymphatic system. These worms live for about 6–8 years and release large numbers of microfilariae into the blood. Mosquitoes are infected during bloodmeals from infected hosts, following which the microfilariae mature into infective larvae. When these infected mosquitoes bite people, mature parasite larvae are deposited on the skin from where they can enter the body.

Most infections are asymptomatic, with no external signs of infection while contributing to transmission of the parasite. These asymptomatic infections still cause damage to the lymphatic system and the kidneys, and alter the body's immune system. When lymphatic filariasis develops into chronic conditions it leads to lymphoedema or elephantiasis of limbs and hydrocele. These body deformities often lead to social stigma and poor psychological wellbeing, loss of income-earning opportunities and increased medical expenses for patients and their caretakers.

893 million people in 49 countries worldwide remain threatened by lymphatic filariasis and require preventive chemotherapy to stop the spread of this parasitic infection. The WHO's strategy for elimination of filiariasis, which concentrates on key components:

- stopping the local spread of infection through large-scale annual treatment by mass drug administration regimes containing albendazole
- alleviating the suffering caused by lymphatic filariasis
- vector control as a supplementary strategy

#### Compiled By

**Dr. Chathurika Herath**.PG Trainee in Community Medicine, Epidemiology Unit, Ministry of Health

#### References:

World Health Organisation: Fact sheets on NTDs https://www.who.int/news-room/fact-sheets/detail/leishmaniasis https://www.who.int/news-room/fact-sheets/detail/leprosy

https://www.who.int/news-room/fact-sheets/detail/lymphatic-filariasis#:~-text=Lymphatic% 20filariasis%2C%20commonly%20known%20as.damage%20to%20the%20lymphatic% 20system.

<b>Table 1 : Water Quality Surveillance</b>	
Number of microbiological water samples	Inly 2020

District	MOH areas	No: Expected *	No: Received
Colombo	15	90	NR
Gampaha	15	90	NR
Kalutara	12	72	NR
Kalutara NIHS	2	12	NR
Kandy	23	138	NR
Matale	13	78	NR
Nuwara Eliya	13	78	12
Galle	20	120	NR
Matara	17	102	NR
Hambantota	12	72	31
Jaffna	12	72	103
Kilinochchi	4	24	69
Manner	5	30	0
Vavuniya	4	24	33
Mullatvu	5	30	NR
Batticaloa	14	84	NR
Ampara	7	42	NR
Trincomalee	11	66	NR
Kurunegala	29	174	12
Puttalam	13	78	NR
Anuradhapura	19	114	NR
Polonnaruwa	7	42	0
Badulla	16	96	NR
Moneragala	11	66	NR
Rathnapura	18	108	NR
Kegalle	11	66	21
Kalmunai	13	78	NR

<sup>\*</sup> No of samples expected (6 / MOH area / Month)

**NR** = Return not received

Table 1: Selected notifiable diseases reported by Medical Officers of Health 08th-14th Aug 2020 (33rd Week)

					i								•	;										į	
	Dengue Fever	Fever	Dysentery	ıtery	Encepha litis		Enteric Fever	<sub>ይ</sub>	Food Poisoning		Leptospiro sis		Typhus Fever	Viral Hepa	Viral Hepatitis	Rat Tak	Human Rabies		Chickenpox	Meningitis		Leishmani- asis	ni- WRCD	9	
	A	В	⋖	В	A B	∢	В	⋖	Ф	⋖	æ	⋖	В	∢	В	⋖	Ф	⋖	В	A	∢	a	<u>*</u>	*	
	22	3640	0	22	0	6	0	2	0	14	10	236	0	2 0		3	0 0	Т	186	П	35	0	7	26	100
	20	2187	0	8	0	2	0	9	0	19	9	179	0	4		6 1	_	0	228	7	24	0	40	42	86
	23	1542	0	10	0	2	0	4	0	4	12	205	0	13 1		9	0 0	2	256	0	33	0	0	66	66
	101	2595	7	23	0	П	0	6	0	13	2	168	T1	83 0		4	0 0	0	142	П	22	0	22	63	100
	7	529	0	7	0	е	0	Ŋ	0	9	0	87	0	0 9		)	0 1	7	49	7	4	11 2	232	. 29	100
	m	153	m	26	0	П	2	m	0	6	9	93	2	70 0		3	0	0	69	2	12	0	0	23	100
	70	1505	7	29	0	17	0	4	0	14	22	522	2	20 0		3	0 0	7	281	m	46	0	4	31	66
	8	325	0	7	0	4	0	7	ъ	47	6	178	2,	51 2		)	0 1	4	159	П	39	19	514	69	100
	7	471	Н	22	<b>+</b>	15	0	П	0	m	<b>'</b>	389	0	10 0		7	0	m	117	0	17	7	295	16	100
	ъ	1975	m	75	0	0	0	20	0	23	0	20	3 4	499 0		0	0 1	1	95	0	6	П	7	31	93
	0	124	0	37	0	7	0	11	0	13	0	18	-	29 0		1	0 0	0	12	0	10	0	13	64	100
	М	133	0	0	0	0	0		0	7	0	9	0	2 0		0	0 0	0	2	1	8	0	0	39	100
	0	248	0	11	0	0	0	2	0	m	0	40	0	1 0		0	0 0	0	29	0	4	0	-1	64	100
	П	80	П	6	0	0	0	9	0	7	0	20		10 0		3	0 2	0	6	П	2	0	9	43	93
	14	2297	П	29	0	4	0		н	46	-	27	0	0 0		2	0 1	2	83	0	18	0		25	100
	7	305	П	16	0	4	0	0	0	0	4	83	0	0 0		7	0 0	9	105	0	15	н	2	89	100
	0	2268	0	12	0	0	0	0	0	7	0	28	0	0 6		0	0 0	2	83	0	<sub>∞</sub>	0	0	46	91
	19	823	7	21	-1	6	0	m	0	36	∞	165	0	25 0		2	0	m	287	7	78	20	324	47	66
	10	437	0	6	0	4	0	က	0	Н	7	22	Η.	15 0		0	0 1	1	72	П	43	m	6	22	100
Anuradhapur	m	386	0	16	0		0	4	-	53	9	216	0	20 0		12 (	0 1	0	164	m	45	2	160	42	92
Polonnaruwa	7	223	0	5	0	0	0	0	0	2	 H	117	0	1 0		17 (	0 1	1	118	0	14	8	187	62	83
	7	430	0	15	0	2	0	e	0	4	13	272	m	26 0		13 (	0 0	0	127	0	30	0	17	29	93
	0	0	0	0	0	0	0	0	0	0	0	0	0	0		0	0 0	0	0	0	0	0	0		
	20	1708	4	73	m	56	0	2	2	34	37 1:	1172	2	38 1	1	15 (	0 0	7	157	ю	88	9	66	49	100
	16	089	П	18	0	8	0	e	0	17	22	350	7	39 0		6	0 0	7	143	9	49	П	25	22	100
	Ŋ	878	ю	48	0	е	1	1	0	9	0	16	0	2 0		3	0 0	1	268	П	35	0	0	69	100
	401	25942	74	286	5 17	126	3 105	Σ.	6	352 1	16 49	4959 2	23 105	55	128	8	13	35	3241	30	641 7	79 19	1991	49	95

\*T=Timeliness refers to returns received on or before 14th Aug, 2020 Total number of reporting units 356 Number of reporting units data provided for the current week: 318 C\*\*-Completeness Source: Weekly Returns of Communicable Diseases (WRCD).

# Table 2: Vaccine-Preventable Diseases & AFP

# 08th-14th Aug 2020 (33rd Week)

Disease	No. of	Cases b	y Provinc	е						Number of cases during current	Number of cases during same	Total num- ber of cases to	Total num- ber of cases to date in	Difference between the number of cases to date in
	W	С	S	N	Е	NW	NC	U	Sab	week in 2020	week in 2019	date in 2020	2019	2020 & 2019
AFP*	00	00	00	00	00	00	00	00	01	01	03	26	50	- 46.8 %
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %
Mumps	00	00	02	00	00	00	00	00	01	03	09	119	232	- 48.7 %
Measles	01	00	00	00	00	00	00	00	00	01	02	36	225	- 84 %
Rubella	00	00	00	00	00	00	00	00	00	00	00	00	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	03	14	- 76.9 %
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	31	10	210 %
Whooping Cough	00	00	00	00	00	00	00	00	00	00	00	05	36	- 86.1 %
Tuberculosis	84	23	00	17	14	26	00	34	05	203	229	3884	5404	- 28.1 %

## 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 Surveil	lance in Sentinel	Hospitals - ILI & SARI					
N 1	Human				Animal		
Month	No Total	No Positive	Infl A	Infl B	Pooled samples	Serum Samples	Positives
May							
Source: Medical	Research Institut	e & Veterinary Research 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

Dr. Sudath Samaraweera CHIEF EPIDEMIOLOGIST EPIDEMIOLOGY UNIT 231, DE SARAM PLACE COLOMBO 10