



## WEEKLY EPIDEMIOLOGICAL REPORT

A publication of the Epidemiology Unit  
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### Trypanosomiasis, human African (sleeping sickness)

Human African trypanosomiasis, also known as sleeping sickness, is a vector-borne parasitic disease. It is caused by infection with protozoan parasites belonging to the genus *Trypanosoma*.

Human African trypanosomiasis, transmitted by the bite of the 'Glossina' insect, commonly known as the tsetse fly.

They are transmitted to humans by tsetse fly (*Glossina* genus) bites which have acquired their infection from human beings or from animals harbouring human pathogenic parasites.

Tsetse flies are found just in sub-Saharan Africa though only certain species transmit the disease. Rural populations living in regions where transmission occurs and which depend on agriculture, fishing, animal husbandry or hunting are the most exposed to the tsetse fly and therefore to the disease.

Human African trypanosomiasis takes 2 forms, depending on the parasite involved:

- *Trypanosoma brucei gambiense* is found in 24 countries in west and central Africa. This form currently accounts for 97% of reported cases of sleeping sickness and causes a chronic infection. A person can be infected for months or even years without major signs or symptoms of the disease. When more evident symptoms arise, the patient is often already in an advanced disease stage where the central nervous system is affected.
- *Trypanosoma brucei rhodesiense* is found in 13 countries in eastern and southern Africa. Nowadays, this form represents under 3% of reported cases and causes an acute infection. First signs and symptoms are observed a few months or weeks after infection. The disease develops rapidly and invades the central nervous system. Only Uganda presents both forms of the disease, but in separate zones.

Another form of trypanosomiasis occurs mainly in Latin America. It is known as American trypanosomiasis or Chagas disease.

#### Disease burden

Sleeping sickness threatens millions of people in 36 countries in sub-Saharan Africa. Many of the affected populations live in remote rural areas with limited access to adequate health services, which complicates the surveillance and therefore the diagnosis and treatment of cases. In addition, displacement of populations, war and poverty are important factors that facilitate transmission.

During the most recent epidemic the prevalence reached 50% in several villages in Angola, the Democratic Republic of the Congo, and South Sudan. Sleeping sickness was the first or second greatest cause of mortality in those communities, even ahead of HIV/AIDS.

#### Infection and symptoms

The disease is mostly transmitted through the bite of an infected tsetse fly but there are other ways in which people are infected:

Mother-to-child infection: the trypanosome can cross the placenta and infect the foetus.

Mechanical transmission through other

- blood-sucking insects is possible, however, it is difficult to assess its epidemiological impact.
- Accidental infections have occurred in laboratories due to pricks with contaminated needles.
- Transmission of the parasite through sexual contact has been documented.

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In the first stage, the trypanosomes multiply in subcutaneous tissues, blood and lymph. This is also called haemo-lymphatic stage, which entails bouts of fever, headaches, joint pains and itching.

In the second stage the parasites cross the blood-brain barrier to infect the central nervous system. This is known as the neurological or meningo-encephalic stage. In general this is when more obvious signs and symptoms of the disease appear: changes of behaviour, confusion, sensory disturbances and poor coordination. Disturbance of the sleep cycle, which gives the disease its name, is an important feature. Without treatment, sleeping sickness is considered fatal although cases of healthy carriers have been reported.

**Disease management: diagnosis**

Disease management is made in 3 steps:

- Screening for potential infection. This involves using serological tests (only available for *T.b.gambiense*) and checking for clinical signs - especially swollen cervical lymph nodes.
- Diagnosing by establishing whether the parasite is present in body fluids.

Staging to determine the state of disease progression. This entails examining the cerebrospinal fluid obtained by lumbar puncture. Diagnosis must be made as early as possible to avoid progressing to the neurological stage. Exhaustive screening requires a major investment in human and material resources. In Africa such resources are often scarce, particularly in remote areas where the disease is mostly found. As a result, some infected individuals may die before they can ever be diagnosed and treated.

**Treatment**

The type of treatment depends on the disease stage. The drugs used in the first stage are safer and easier to administer than those for the second stage. Also, the earlier the disease is identified, the better the prospects of a cure. The assessment of treatment outcome requires follow up of the patient up to 24 months and entails laboratory exams of body fluids including cerebrospinal fluid obtained by lumbar puncture, as parasites may remain viable for long periods and reproduce the disease months after treatment.

Treatment success in the second stage depends on drugs that cross the blood-brain barrier to reach the parasite. Such drugs are toxic and complicated to administer. In total five different drugs are used for the treatment of sleeping sickness. These drugs are donated to WHO by manufacturers and distributed free of charge to disease endemic countries.

There is no vaccine or drug for prophylaxis against African trypanosomiasis. Preventive measures are aimed at minimizing contact with tsetse flies.

Although the parasite causing human African trypanosomiasis was identified only in 1901, 'sleeping sickness' is thought to have existed on the African continent for centuries. WHO's ultimate objective is the elimination of human African trypanosomiasis as a public health problem and the implementation of sustained surveillance in all disease-endemic countries.

*Sleeping sickness occurs in 36 sub-Saharan African countries*

*where there are tsetse flies that transmit the disease. The people most exposed to the tsetse fly live in rural areas .*

*Early diagnosis and treatment can completely cure the disease. Sleeping sickness is curable with medication, but is fatal if left untreated.*

Source: Trypanosomiasis, human African (sleeping sickness) Fact Sheet .Available at :[https://www.who.int/news-room/fact-sheets/detail/trypanosomiasis-human-african-\(sleeping-sickness\)](https://www.who.int/news-room/fact-sheets/detail/trypanosomiasis-human-african-(sleeping-sickness))

CDC. African Trypanosomiasis, <https://www.cdc.gov/parasites/sleepingsickness/index.html>

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Sri Lanka

Table 1 : Water Quality Surveillance Number of microbiological water samples December 2018			
District	MOH areas	No: Expected *	No: Received
Colombo	15	90	102
Gampaha	15	90	NR
Kalutara	12	72	NR
Kalutara NIHS	2	12	NR
Kandy	23	138	NR
Matale	13	78	74
Nuwara Eliya	13	78	NR
Galle	20	120	NR
Matara	17	102	NR
Hambantota	12	72	37
Jaffna	12	72	136
Kilinochchi	4	24	1
Manner	5	30	NR
Vavuniya	4	24	NR
Mullatvu	5	30	NR
Batticaloa	14	84	109
Ampara	7	42	35
Trincomalee	11	66	NR
Kurunegala	29	174	111
Puttalam	13	78	NR
Anuradhapura	19	114	16
Polonnaruwa	7	42	40
Badulla	16	96	66
Moneragala	11	66	75
Rathnapura	18	108	NR
Kegalle	11	66	62
Kalmunai	13	78	NR

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

Table 1: Selected notifiable diseases reported by Medical Officers of Health 12<sup>th</sup> - 18<sup>th</sup> Jan 2019 (3<sup>rd</sup> Week)

RDHS Division	Dengue Fever		Dysentery		Encephalitis		Enteric Fever		Food Poisoning		Leptospirosis		Typhus Fever		Viral Hepatitis		Human Rabies		Chickenpox		Meningitis		Leishmaniasis		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	259	884	3	3	0	1	1	4	8	1	3	1	2	0	0	8	30	1	8	0	0	44	100				
Gampaha	161	539	0	0	1	1	0	0	10	1	3	0	0	0	0	11	27	1	1	3	11	52	98				
Kalutara	78	281	0	3	1	3	0	1	5	12	46	0	1	0	0	21	69	6	10	3	3	59	100				
Kandy	70	219	0	4	0	0	0	0	2	4	12	2	8	0	1	5	19	0	4	1	1	61	100				
Matale	19	45	1	2	0	0	0	0	0	3	9	0	0	1	1	2	7	1	1	17	32	67	100				
Nuwareliya	3	22	0	1	0	0	0	0	0	3	5	1	4	0	1	1	4	0	1	0	0	23	100				
Galle	43	100	0	1	1	1	0	1	0	6	20	2	8	0	1	10	26	2	5	0	1	62	97				
Hambantota	37	111	1	1	0	0	0	0	0	1	5	7	15	0	1	6	17	0	0	35	60	69	100				
Mataru	42	155	1	1	2	2	0	0	0	1	7	4	10	1	2	6	23	0	2	14	39	63	100				
Jaffna	211	710	2	10	0	1	1	1	0	5	7	28	82	0	0	5	13	0	2	0	0	26	93				
Kilinochchi	8	19	2	3	0	0	1	1	0	5	9	2	4	1	1	0	0	0	0	0	1	42	100				
Mannar	10	24	0	0	0	0	2	3	0	0	0	2	2	0	0	0	0	0	0	0	0	53	100				
Vavuniya	5	34	0	1	0	1	2	5	0	2	3	0	2	0	0	0	8	0	0	0	0	25	100				
Mullaitivu	5	11	1	2	0	0	0	0	0	1	2	1	2	0	0	0	0	0	1	0	1	50	67				
Batticaloa	41	148	5	15	0	0	0	0	0	0	5	0	0	0	0	1	8	0	0	0	0	67	100				
Ampara	6	21	1	6	0	0	0	0	0	1	5	0	0	2	3	0	8	14	0	1	0	43	100				
Trincomalee	5	77	0	0	0	0	0	0	0	0	0	0	0	0	0	0	4	0	0	0	0	39	78				
Kurunegala	58	171	1	7	1	4	0	1	0	8	22	2	3	0	2	9	39	4	7	16	56	52	100				
Puttalam	20	63	0	4	0	0	0	0	0	1	4	1	2	0	0	3	9	0	0	0	1	59	100				
Anuradhapura	19	39	0	3	0	2	0	0	0	4	23	0	1	1	2	9	31	0	1	5	37	39	97				
Polonnaruwa	12	31	3	5	0	1	0	0	0	3	10	1	1	0	13	22	3	4	7	19	50	100					
Badulla	23	67	1	5	1	1	0	1	0	3	4	24	8	0	6	16	3	14	0	2	56	100					
Monaragala	13	48	0	2	0	0	0	0	0	5	27	1	7	0	7	22	2	8	0	0	67	100					
Ratnapura	36	129	3	6	2	7	0	0	2	2	9	38	1	0	9	30	3	12	0	1	40	100					
Kegalle	36	109	2	3	0	1	0	0	1	13	4	11	2	0	7	32	0	1	0	1	55	100					
Kalmune	36	90	4	5	0	0	0	0	0	0	0	0	0	0	4	9	0	0	0	0	51	100					
<b>SRILANKA</b>	<b>1256</b>	<b>4147</b>	<b>31</b>	<b>93</b>	<b>9</b>	<b>26</b>	<b>6</b>	<b>15</b>	<b>5</b>	<b>38</b>	<b>85</b>	<b>305</b>	<b>166</b>	<b>58</b>	<b>166</b>	<b>7</b>	<b>19</b>	<b>0</b>	<b>2</b>	<b>151</b>	<b>479</b>	<b>26</b>	<b>83</b>	<b>10</b>	<b>266</b>	<b>53</b>	<b>98</b>

Source: Weekly Returns of Communicable Diseases (WRCD).

\*T=Timeliness refers to returns received on or before 18<sup>th</sup> January, 2019 Total number of reporting units 353 Number of reporting units data provided for the current week: 344 C\*\*=Completeness  
A = Cases reported during the current week. B = Cumulative cases for the year.

**Table 2: Vaccine-Preventable Diseases & AFP**

**12<sup>th</sup> – 18<sup>th</sup> Jan 2019 (3<sup>rd</sup> Week)**

Disease	No. of Cases by Province									Number of cases during current week in 2019	Number of cases during same week in 2018	Total number of cases to date in 2019	Total number of cases to date in 2018	Difference between the number of cases to date in 2019 & 2018
	W	C	S	N	E	NW	NC	U	Sab					
AFP*	01	02	00	00	00	00	00	00	00	03	02	06	03	100 %
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %
Mumps	00	00	02	00	01	01	01	00	01	06	03	18	10	80 %
Measles	01	00	01	00	00	00	01	00	00	03	03	10	06	66.6 %
Rubella	00	00	00	00	00	00	00	00	00	00	00	00	02	-100 %
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	01	02	04	- 50 %
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %
Japanese Encephalitis	00	00	00	01	00	00	00	00	00	01	01	02	04	50 %
Whooping Cough	02	00	01	00	00	01	00	00	00	04	01	05	02	150 %
Tuberculosis	78	15	39	11	10	03	00	09	33	198	175	509	462	10.1%

**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 Surveillance in Sentinel Hospitals - ILI & SARI							
Month	Human				Animal		
	No Total	No Positive	Infl A	Infl B	Pooled samples	Serum Samples	Positives
January	95	14	10	4	2247	2001	0

Source: Medical Research Institute & Veterinary Research Institute

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**ON STATE SERVICE**

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