

WEEKLY EPIDEMIOLOGICAL REPORT

A publication of the Epidemiology Unit Ministry of Health

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18th - 24th October 2014

Lymphatic filariasis (Part-II)

This is the second in a series of two articles on Lymphatic filariasis

Management

The medical management of a filarial infection should be specific and based on the microfilariae isolated or antigenaemia detected.

Mass drug administration reduces the transmission of filarial infection and disease morbidity by decreasing the burden of microfilaraemia, resulting in suboptimal levels for transmission by disease vectors.

For example, annual mass treatment with albendazole and ivermectin is employed to interrupt the transmission of W bancrofti. Since this species has no alternative hosts, this approach could theoretically result in eventual eradication of bancroftian filariasis.

One study evaluated the effect of higher dose and increased frequency (twice yearly) of albendazole-ivermectin therapy for W bancrofti and found that it resulted in complete microfilarial clearance, as well as a more sustained clearance than that resulting from standard-dose albendazole-ivermectin treatment.

The effects of mass treatment on filariasis have reportedly been sustained for up to 6 years. No filariasis vaccine is currently available, but efforts to develop an effective one are underway.

Surgery

Large hydroceles and scrotal elephantiasis can be managed with surgical excision. Correcting gross limb elephantiasis with surgery is less successful and may necessitate multiple procedures and skin grafting.

Diet and Activity

Fatty foods are restricted in individuals with proven chyluria that is associated with lymphatic filariasis.

Individuals with chronic lymphatic filariasis are encouraged to mobilize (with compression bandage support) the affected limb.

WHO's response

World Health Assembly Resolution 50.29 encourages Member States to eliminate lymphatic filariasis as a public health problem. In response, WHO launched its Global Programme to Eliminate Lymphatic Filariasis (GPELF) in 2000 with the aim of eliminating the disease as a public-health problem. In 2012, the WHO NTD Roadmap reconfirmed the target date for achieving elimination by 2020.

WHO's strategy is based on 2 key components:

- Stopping transmission through large-scale annual treatment of all eligible people in an area or region where infection is present;
- Alleviating the suffering caused by lymphatic filariasis through increased morbidity management and disability prevention activities

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Large-scale treatment (mass drug administration)

Prevention of lymphatic filariasis is possible by stopping the spread of the infection. Large-scale treatment involves a single dose of 2 medicines given annually to an entire at-risk population in the following way: albendazole (400 mg) together with ivermectin (150-200 mcg/kg) or with diethylcarbamazine citrate (DEC) (6 mg/kg).

These preventive chemotherapy medicines have a limited effect on adult parasites but effectively clear microfilariae from the bloodstream and prevent the spread of parasites to mosquitoes. Large-scale treatment conducted annually for 4-6 years, treating all persons living in areas where the infection is present can interrupt the transmission cycle.

By 2012, 56 countries had started implementing large-scale treatment through mass drug administration (MDA). Of the 56 countries that had implemented MDA, 13 countries have moved to the post-MDA surveillance phase.

From 2000 to 2012, more than 4.4 billion treatments were delivered to a targeted population of about 984 million individuals in 56 countries, considerably reducing transmission in many places.

Recent research data show that the transmission of lymphatic filariasis in at-risk populations has dropped by 43% since the beginning of the GPELF. The overall economic benefit of the programme during 2000-2007 is conservatively estimated at US\$ 24 billion.

Morbidity management

Morbidity management and disability prevention are vital for improving public health and should be fully integrated into the health system. Surgery can alleviate most cases of hydrocele. Clinical severity of lymphoedema and acute inflammatory episodes can be improved using simple measures of hygiene, skin care, exercise, and elevation of affected limbs.

The GPELF aims to provide access to a minimum package of care for every person with associated chronic manifestations of lymphatic filariasis in all areas where the disease is present, thus alleviating suffering and promoting improvement in their quality of life.

Vector control

Mosquito control is another supplemental strategy supported by WHO. It is used to reduce transmission of lymphatic filariasis and other mosquito-borne infections. Measures such as insecticide-treated nets or indoor residual spraying may help protect people from infection.

Sources

Lymphatic filariasis- available at <u>http://www.who.int/</u> mediacentre/factsheets/fs102/en/

Filariasis- available at http://emedicine.medscape.com/article/217776-overview#a0101

Compiled by Dr. C U D Gunasekara of the Epidemiology Unit.

Table 1 : Water Quality SurveillanceNumber of microbiological water samples - September/2014									
District	MOH areas	No: Expected *	No: Received						
Colombo	12	72	51						
Gampaha	15	90	92						
Kalutara	12	72	NR						
Kalutara NIHS	2	12	13						
Kandy	23	138	0						
Matale	12	72	52						
Nuwara Eliya	13	78	27						
Galle	19	114	76						
Matara	17	102	13						
Hambantota	12	72	49						
Jaffna	11	66	15						
Kilinochchi	4	24	0						
Manner	5	30	0						
Vavuniya	4	24	7						
Mullatvu	4	24	5						
Batticaloa	14	84	0						
Ampara	7	42	0						
Trincomalee	11	66	NR						
Kurunegala	23	138	39						
Puttalam	9	54	55						
Anuradhapura	19	114	6						
Polonnaruwa	7	42	0						
Badulla	15	90	48						
Moneragala	11	66	42						
Rathnapura	18	108	84						
Kegalle	11	66	11						
Kalmunai	13	78	0						
* No of samples ex NR = Return not re	pected (6 / MOI eceived	I area / Month)							

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Table 1: Selected notifiable diseases reported by Medical Officers of	f Health í
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11th - 17th Oct 2014 (42nd Week)

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kenpox	В	352	249	213	165	48	110	365	128	160	121	15	10	12	5	54	89	94	354	75	200	138	69	LΓ	171	224	89	3587
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Viral epatitis	8	44	224	19	166	129	31	8	16	36	6	0	-	2	0	7	5	2	53	4	11	8	125	110	389	220	0	162
	A	2	10	0	14	9	-	2	0	2	-	0	0	0	0	0	0	0	-	0	-	0	2	3	13	-	0	59
nus Fever	B	3	20	m	74	2	55	85	65	53	274	20	24	9	1	2	12	21	43	22	27	8	96	145	96	53	0	1220
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entery	в	120	126	142	80	58	226	105	44	88	525	86	36	49	53	269	64	43	116	62	149	40	149	59	195	96	108	3088
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e Fever	в	119	6036	2179	1405	393	246	887	535	557	666	47	128	108	87	680	129	508	1741	559	442	445	590	240	2532	1362	146	34043
Dengue	A	176	119	36	09	11	6	5	5	32	50	-	15	0	0	7	-	ę	32	11	6	10	21	3	45	12	Ъ	675
RDHS Division		Colombo	Gampaha	Kalutara	Kandy	Matale	NuwaraEliya	Galle	Hambantota	Matara	Jaffna	Kilinochchi	Mannar	Vavuniya	Mullaitivu	Batticaloa	Ampara	Trincomalee	Kurunegala	Puttalam	Anuradhapura	Polonnaruwa	Badulla	Monaragala	Ratnapura	Kegalle	Kalmune	SRILANKA

•T=Timeliness refers to returns received on or before 17th October, 2014 Total number of reporting units 337 Number of reporting units data provided for the current week: 257 C**-Completeness

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Table 2: Vaccine-Preventable Diseases & AFP

18th – 24th October 2014

11th - 17th Oct 2014 (42nd Week)

Disease			٦	lo. of Cas	ses by P	rovince	1	Number of cases during current	Number of cases during same	Total number of cases to date in	Total number of cases to date in	Difference between the number of cases to date			
	W	С	S	N	E	NW	NC	U	Sab	week in 2014	week in 2013	2014	2013	in 2013& 2014	
AFP*	00	00	00	00	01	00	01	00	00	02	01	67	76	-11.9%	
Diphtheria	00	00	00	00	00	00	00	00	00	00	-	00	-	%	
Mumps	01	00	02	02	02	01	00	01	00	09	12	567	1274	-55.5%	
Measles	11	02	04	00	01	03	00	01	02	24	30	2826	3215	-12.1%	
Rubella	00	00	00	00	00	00	00	00	00	00	01	17	26	-34.6%	
CRS**	00	00	00	00	00	00	00	00	00	00	00	04	06	-33%	
Tetanus	00	00	00	00	00	00	00	00	00	00	00	12	19	-36.9%	
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	00	%	
Japanese Encephalitis	00	00	00	00	00	00	00	00	00	00	00	22	68	-67.6%	
Whooping Cough	00	00	00	01	00	00	01	01	01	04	01	61	70	-12.9%	
Tuberculosis	89	41	15	12	03	06	08	08	31	213	20	7962	6617	+20.3%	

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: Mullattivu, 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	ILI	SARI	Infl A	Infl B	Pooled samples	Serum Samples	Positives				
September	2249	100	19	4	1	600	666	0				

Source: Medical Research Institute & Veterinary Research Institute

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