Global Programme to Eliminate Lymphatic Filariasis fundamentals towards successful NTD elimination



courtesy of Nepal and GSK

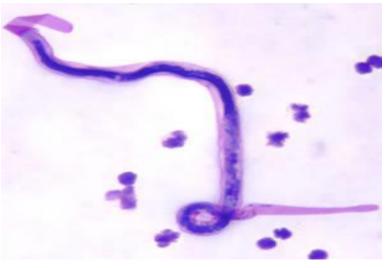


Lymphatic filariasis

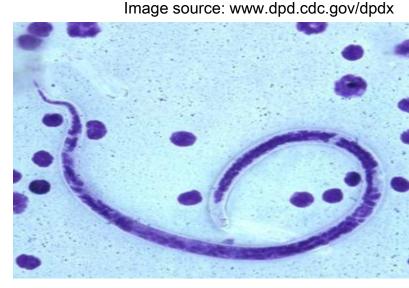
- NTD caused by three species of nematodes that nest in the lymphatic vessels
- Transmitted by mosquitoes
- Infection impairs function of lymphatic system by vessel dilatation



Wuchereria bancrofti



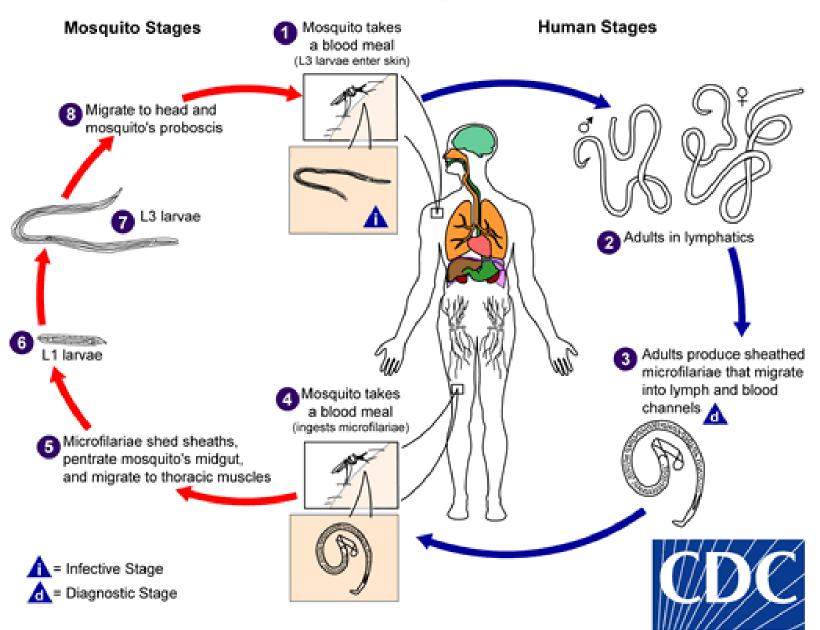
Brugia malayi



Brugia timori



Wuchereria bancrofti



LF Morbidity— a leading cause of global disability

Chronic manifestations

- Lymphoedema
- Elephantiasis
- Hydrocele

Acute manifestations

Adenolymphangitis (ADL)
 'acute attacks'



Social and Economic burden of LF

- Disfigurement, pain & disability
- Stigma
- Depressive illness
- Economic loss
- Causes and aggravates poverty





courtesy of Nepal

and

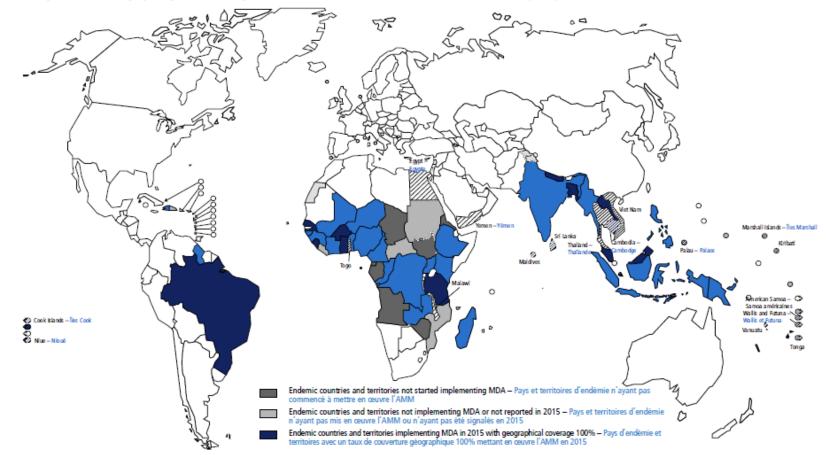
Kiribati

Distribution of lymphatic filariasis (LF)

Endemic in 72 countries; 856 million people require mass treatment

Map 1 Countries where lymphatic filariasis is endemic and status of mass drug administration (MDA) in those countries, 2015

Carte 1 Pays où la filariose lymphatique est endémique et situation de l'administration massive de médicaments (AMM) en 2015



Transmission to Elimination

What are the fundamentals needed to move from transmission towards successful elimination?

1. Global commitment – WHA 50.29 Elimination of LF as a public health problem

Global Programme to Eliminate Lymphatic Filariasis (GPELF) was launched in 2000

1. Stop transmission

Mass drug administration (MDA)

2. Reduce suffering and improve quality of life

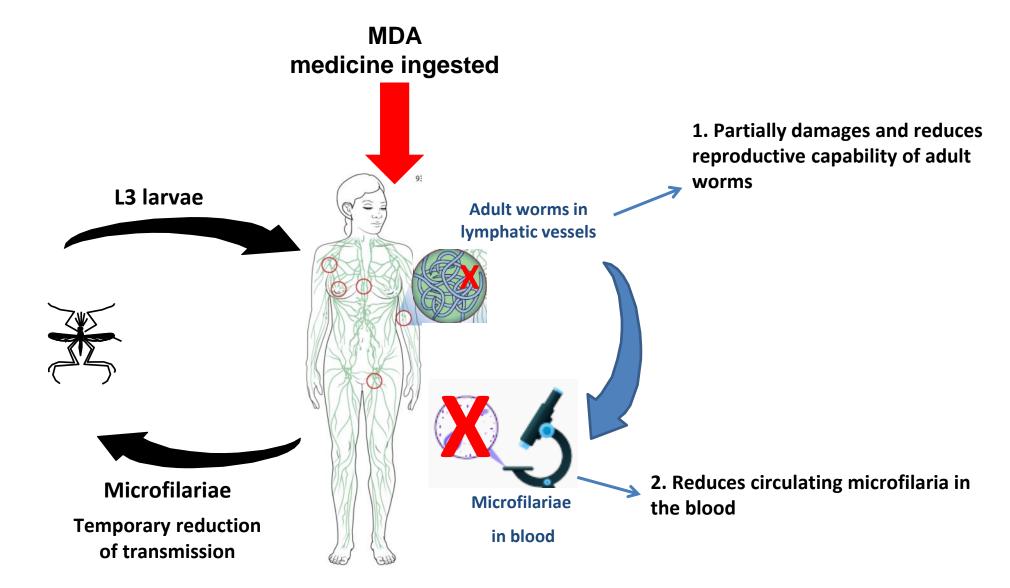
Morbidity management and disability prevention (MMDP)



2. Effective strategies – Mass Drug Administration

- Annual mass treatment with antihelminthic medicine (treating both infected and uninfected)
- 100% geographical coverage = ALL endemic districts
- Effective (≥65%) coverage of the *total* population ≥5 rounds

How MDA works

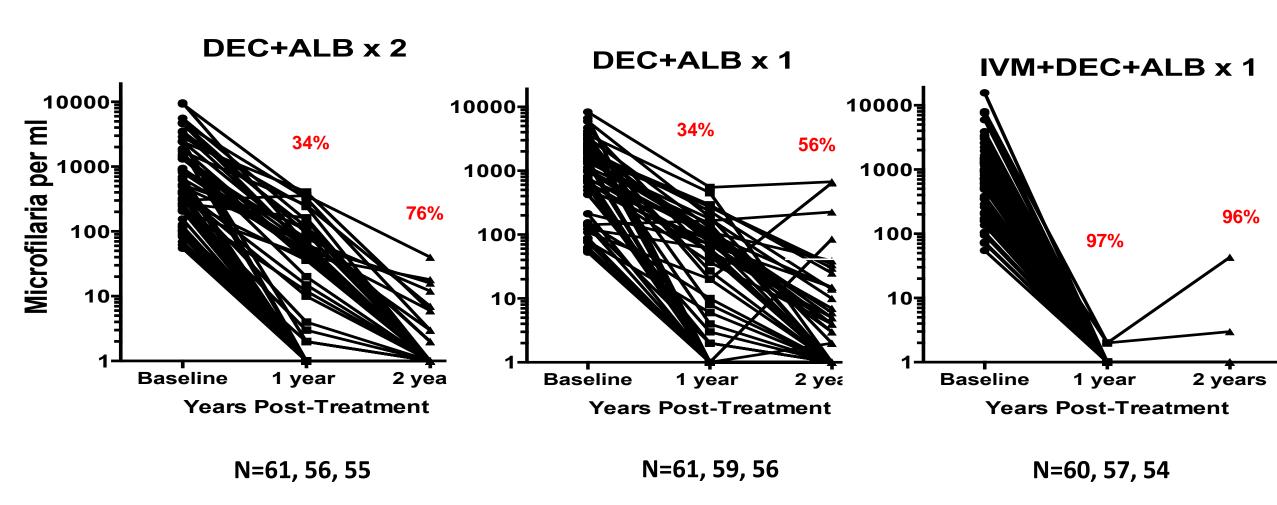


2. Effective strategies - MDA medicine

WHO recommends the following MDA regimens:

- New triple-drug therapy for annual MDA in special settings
 - Ivermectin + diethycarbamazine + albendazole (IDA or triple therapy)
- Annual two-drug therapy for all other areas
 - diethycarbamazine + albendazole
 - ivermectin + albendazole
- Biannual albendazole alone in loiasis co-endemic areas

Efficacy of IDA



Courtesy of Chris King 2017 DOLF PNG

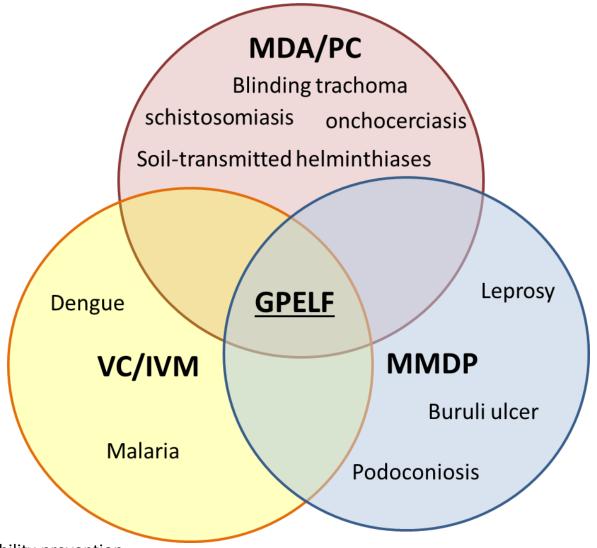
2. Effective strategies Morbidity Management and Disability Prevention



Health facilities must be able to provide minimum package of care in every district with known *patients*

- surgery for hydrocele
- management of lymphedema
- treatment of acute attacks
- treatment of patients with LF infection

3. Integration: where feasible for broad impact



PC – preventive chemotherapy

VC – vector control

IVM – integrated vector management;

MMDP – morbidity management and disability prevention

4. Standardized tools: Recommended diagnostic tests

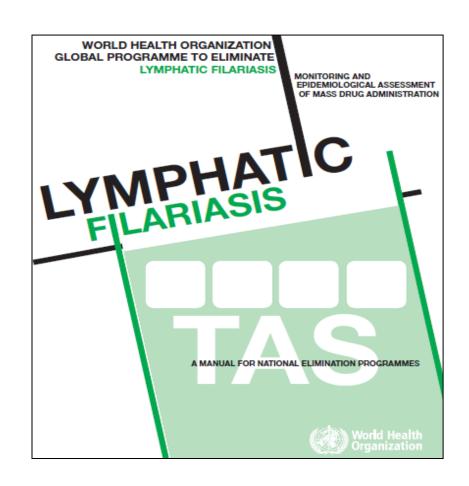
Field assay	Detection target	Recommended for use during
Blood film	Microfilariae	Mapping, Sentinel site and Spot- check site monitoring
Alere Filariasis Test Strip (FTS)	Filarial antigen (Ag)	Mapping, Sentinel site and Spotcheck site monitoring, and Transmission Assessment Survey (TAS)
Brugia Rapid [™] test	Antifilarial antibody (Ab)	TAS



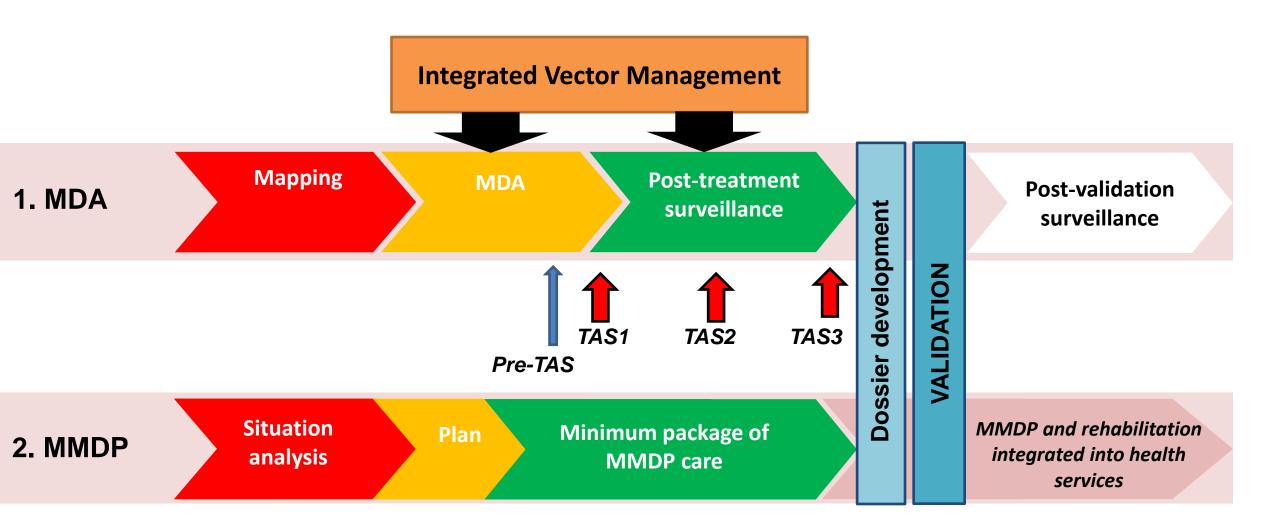


4. Standardized tools: Transmission Assessment Survey

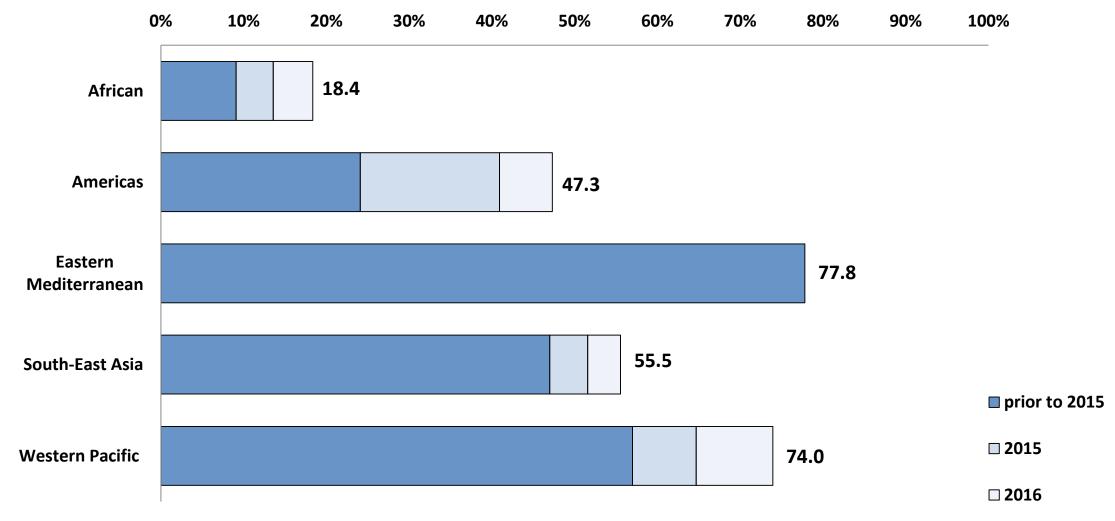
- Decision making tool, tells when to stop MDA
- Standardized survey with robust, yet practical statistical design
- Uses children as an indicator of incident infection
- Measures whether prevalence of infection is below a threshold (critical cut-off) at which transmission is assumed no longer sustainable, even in the absence of MDA



5. Strategic Framework



6. Measurable progress: proportion of implementation units that have completed TAS and no longer require MDA*



^{*}Percent of all known endemic implementation units (IU) in countries by region that have completed TAS1 or previous stop-MDA surveys and reported meeting criterion for stopping MDA. IUs where endemicity is unknown have not been included.



6. Measurable progress: MDA status of countries

MDA not started	

MDA started but not at scale

MDA scaled to all endemic IUs

Post-MDA Surveillance

Elimination as a Public Health Problem

Equatorial Guinea Gabon

Sao Tome and Principe

South Sudan

New Caledonia

Angola Cameroon

Central African Republic

Chad

Congo

Democratic Republic Congo

Ethiopia

Guinea-Bissau

Nigeria

Sudan

Comoros

Eritrea

Madagascar

Guyana

Indonesia

Papua New Guinea

16 (22%)

Benin. Burkina Faso Côte d'Ivoire, Ghana, Guinea, Liberia, Mali Mozambique, Niger Senegal, Sierra-Leone Tanzania, Uganda,

Kenya, Zambia, Zimbabwe

Dominican Republic Haiti

India, Myanmar Nepal, Timor-Leste

French Polynesia Fiji, FSM, Lao PDR Malaysia, Samoa Brunei Darussalam Philippines, Tuvalu

31 (43%)

Malawi

Brazil

Egypt

Yemen

Bangladesh

American Samoa

Kiribati

Palau

Vietnam

Wallis and Futuna

10 (14%)

HARAGA

Togo

Maldives Sri Lanka Thailand

Cambodia
Cook Islands
Marshall Islands
Niue
Tonga
Vanuatu

10 (14%)

5 (7%)





499,400,000

Population no longer requiring MDA in 44 countries

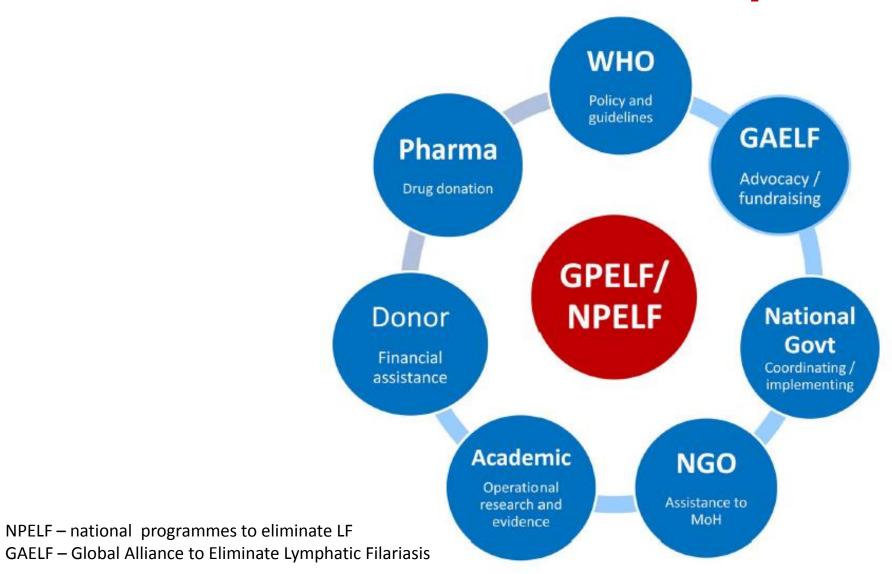
Prevented or cured more than 97 million cases

(Ramaiah, Ottesen 2014 PLoS NTD)

Forecasted to avert >US \$100 billion in economic loss

(Turner et al 2016 Infect Dis Pov)

7. Partnerships



7. Partnerships: Government Leadership







7. Partnerships: Donations - medicines

	Ivermectin	Diethylcarbamazine (DEC)	Albendazole
Dosage	150-200 μg/kg	6mg/kg	400mg
Manufacturer	Merck Sharp & Dohme (MSD)	Eisai	GSK
Commitment	a. Since 1997b. 2017-2025	2014-2020	Since 2009 until elimination is achieved
Donation	 a. Needed amount for LF elimination in onchocoendemic countries b. Up to 280 million tablets per year for IDA 	Up to 2.2 billion tablets	Up to 600 million tablets annually

7. Partnerships: Operational research

- Research that directly contributes to the achievement of GPELF aims
 - -raises level of awareness and directs investments
 - develops more effective strategies
 - identifies programme challenges and solutions
 - -facilitates costing, forecasting and planning

Conclusion

What are the fundamentals driving GPELF towards success?

- 1. Global commitment
- 2. Effective strategies
- 3. Integrated approach
- 4. Standardized tools
- 5. Strategic framework
- 6. Measurable progress
- 7. Partnerships





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Associated Institute of the University of Basel

Swiss Centre for International Health Health Systems Support Unit Data-Evaluation-Evidence-Policy (DEEP) Group

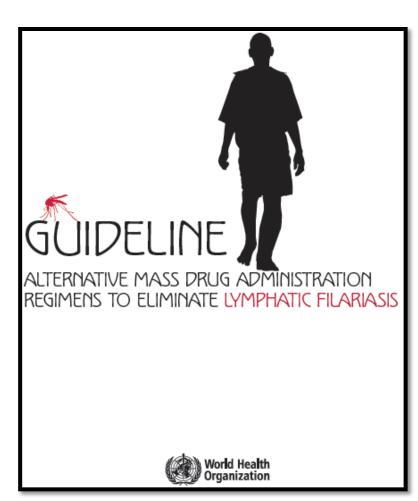
WHO Guideline development group meeting

Systematic Review: alternative MDA strategies for the elimination of lymphatic filariasis



Xavier Bosch-Capblanch, Peter Steinmann, Amanda Ross, Heather Ames, Meike Zuske

Geneva, 17-19 May 2017



Current obstacles to success

- MDA has not started in all endemic districts
- Sustaining effective coverage for ≥5 MDA rounds
- Sub-optimal results of impact surveys
 - 10% of TAS have failed across 14 countries
- Persistent transmission in certain settings
 - Aedes vector
 - Zoonotic Brugia malayi
- Need more sensitive tools/responsive targets for measuring impact of better regimens



Thank you

Swiss TPH!

courtesy of Zanzibar and GSK

