

# **WEEKLY EPIDEMIOLOGICAL REPORT**

# A publication of the Epidemiology Unit Ministry of Health

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

Malaria (Part I)

# Vol. 42 No. 15

# 04<sup>th</sup> – 10<sup>th</sup> April 2015

This is the first in a series of three-article on Malaria

### Key facts

- Malaria is a life-threatening disease caused by parasites that are transmitted to people through the bites of infected mosquitoes.
- In 2013, malaria caused an estimated 584 000 deaths (with an uncertainty range of 367 000 to 755 000), mostly among African children.
- Malaria is preventable and curable.
- Increased malaria prevention and control measures are dramatically reducing the malaria burden in many places.
- Non-immune travellers from malaria-free areas are very vulnerable to the disease when they get infected.

According to the latest estimates, released in December 2014, there were about 198 million cases of malaria in 2013 (with an uncertainty range of 124 million to 283 million) and an estimated 584 000 deaths (with an uncertainty range of 367 000 to 755 000). Malaria mortality rates have fallen by 47% globally since 2000, and by 54% in the WHO African Region.

Most deaths occur among children living in Africa where a child dies every minute from malaria. Malaria mortality rates among children in Africa have been reduced by an estimated 58% since 2000. Malaria is caused by Plasmodium parasites. The parasites are spread to people through the bites of infected Anopheles mosquitoes, called "malaria vectors", which bite mainly between dusk and dawn.

There are four parasite species that cause malaria in humans:

- Plasmodium falciparum
- Plasmodium vivax
- Plasmodium malariae
- Plasmodium ovale.

Plasmodium falciparum and Plasmodium vivax are the most common. Plasmodium falciparum is the most deadly.

In recent years, some human cases of malaria have also occurred with Plasmodium knowlesi – a species that causes malaria among monkeys and occurs in certain forested areas of South-East Asia.

#### Transmission

Malaria is transmitted exclusively through the bites of Anopheles mosquitoes. The intensity of transmission depends on factors related to the parasite, the vector, the human host and the environment.

About 20 different Anopheles species are locally important around the world. All of the important vector species bite at night. Anopheles mosquitoes breed in water and each species has its own breeding preference; for example some prefer shallow collections of fresh water, such as

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puddles, rice fields and hoof prints. Transmission is more intense in places where the mosquito lifespan is longer (so that the parasite has time to complete its development inside the mosquito) and where it prefers to bite humans rather than other animals. For example, the long lifespan and strong human-biting habit of the African vector species is the main reason why about 90% of the world's malaria deaths are in Africa.

Transmission also depends on climatic conditions that may affect the number and survival of mosquitoes, such as rainfall patterns, temperature and humidity. In many places, transmission is seasonal, with the peak during and just after the rainy season. Malaria epidemics can occur when climate and other conditions suddenly favour transmission in areas where people have little or no immunity to malaria. They can also occur when people with low immunity move into areas with intense malaria transmission, for instance to find work, or as refugees.

Human immunity is another important factor, especially among adults in areas of moderate or intense transmission conditions. Partial immunity is developed over years of exposure, and while it never provides complete protection, it does reduce the risk that malaria infection will cause severe disease. For this reason, most malaria deaths in Africa occur in young children, whereas in areas with less transmission and low immunity, all age groups are at risk.

#### Symptoms

Malaria is an acute febrile illness. In a non-immune individual, symptoms appear seven days or more (usually 10–15 days) after the infective mosquito bite. The first symptoms – fever, headache, chills and vomiting – may be mild and difficult to recognize as malaria. If not treated within 24 hours, *P. falcipa-rum* malaria can progress to severe illness often leading to death. Children with severe malaria frequently develop one or more of the following symptoms: severe anemia, respiratory distress in relation to metabolic acidosis, or cerebral malaria. In adults, multi-organ involvement is also frequent. In malaria endemic areas, persons may develop partial immunity, allowing asymptomatic infections to occur.

For both P. vivax and P. ovale, clinical relapses may occur weeks to months after the first infection, even if the patient has left the malarias' area. These new episodes arise from dormant liver forms known as hypnozoites (absent in P. falciparum and P. malariae); special treatment – targeted at these liver stages – is required for a complete cure.

#### Who is at risk?

Approximately half of the world's population is at risk of malaria. Most malaria cases and deaths occur in sub-Saharan Africa. However, Asia, Latin America and to a lesser extent the Middle East and parts of Europe are also affected. In 2014, 97 countries and territories had ongoing malaria transmission.

Specific population risk groups include:

- Young children in stable transmission areas who have not yet developed protective immunity against the most severe forms of the disease
- Non-immune pregnant women as malaria causes high rates of miscarriage and can lead to maternal death
- Semi-immune pregnant women in areas of high transmission. Malaria can result in miscarriage and low birth weight, especially during first and second pregnancies
- Semi-immune HIV-infected pregnant women in stable transmission areas, during all pregnancies. Women with malaria infection of the placenta also have a higher risk of passing HIV infection to their newborns
- People with HIV/AIDS
- International travellers from non-endemic areas because they lack immunity
- Immigrants from endemic areas and their children living in non-endemic areas and returning to their home countries to visit friends and relatives are similarly at risk because of waning or absent immunity.

#### Sources:

Malaria,	available	at	http://www.who.int/mediacentre/
factsheets	<u>/fs094/en/</u>		

WHO Sri Lanka country report

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

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 Table 1: Selected notifiable diseases reported by Medical Officers of Health
 28th - 03rd
 April 2015 (14th Week)

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WF	τ*	63	80	69	61	62	69	ю	75	100	83	25	60	100	80	71	57	58	78	62	84	71	82	82	67	91	46	68	
nani-	в	0	0	0		m	0	0	87	21	0	0	0	1	m	0	0	ч	27	1	75	35	4	∞	4	0	0	271	
Leishmani- asis	A	0	0	0	0	0	0	0	0		0	0	0	0		0	0			0	ы	2	0	0	0	0	0	1	
gitis	в	12	7	13	4	2	18	13	4	6	ъ	0	0	3	2	10	m	2	7	8	11	11	20	ъ	12	14	2	197	
Meningitis	A	1	1	2	0	0	0	0	0	1	0	0	0	0	0	0	0	0	7	2	0	0	4	0	0	0	0	13	
xodue	в	146	62	91	72	7	19	60	31	83	65	ø	0	9	1	12	72	29	145	27	58	45	41	33	35	70	43	1261	
Chickenpox	A	7	7	m	2	0	0		2	4	7	0	0	0	0	0	2	6	4	0	6	0	2		ω	4	m	20	
an es	в	2	0		0	0	0	0	0	0	н		0	-	0	0	0	0		0	0	0	2		0	0	0	9	
Human Rabies	A	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	
Viral Hepatitis	8	14	57	6	57	151	35	4	16	11	7	0	0	H		0	H	4	14	1	~	m	46	20	118	41	0	482	
He	A	0	З	0	m	0	m	0		0	0	0	0	0	0	0	0	0		0	0	0	m	0	2	2	0	18	
Typhus Fever	в	1	9	0	19	4	26	22	15	17	453	7	14	11	9	0	0	2	13	7	11		33	23	22	15	0	728	
Турн	A	0	ю	0	0	0	2	0	ч	0	7	0	0	0	0	0	0	0		0		0	m	0	0	ч	0	19	
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Lepto	A	2	11	4	m	Ч	0		2	6	0	0	0	0	0	0	0	1	2	0	ß	0	0	ß	2	m	0	54	
Food Poisoning	в	48	10	64	2	ω	0	9	4	4	24	25	ч	2	-	16	0	22	10	9	36	0	ъ	2	H	ω	13	348	
Fc Pois	A	1	0	0	0	0	0	0	0	0	m	0	0	0	0	16	0	0		0	2	0	0	0	0	0	0	23	
Enteric Fever	в	25	7	14	13	m	9	2	4	4	121	ω	4	24	4	9	H	14	m	1	2	ъ	m	ω	14	33	0	324	
Enteri	A	0	0	0	2	0		0	0	0	Ч	0	0	e	1	0		0	0	0	0	0	0	0	2	2	0	13	
Encephalit is	в	4	с	m	2	0		0	0	с	7	0	0	4	2	4	0	0	7	2		2	m		ω	ы	0	52	
Ence	A	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0	7	Ġ
Dysentery	В	60	22	30	40	21	86	24	10	27	206	31	4	8	10	89	16	11	57	13	21	22	48	38	96	27	39	1056	ses (WRCI
Dys	A	1	0	0	0	2	ъ	н	0	2	16	0	0	0	0	10	0	0	m	0		-	2	ε	m	0	0	50	e Disea
Dengue Fever	в	3460	1579	571	474	288	76	283	131	180	957	32	68	56	60	891	20	310	623	374	228	108	292	88	372	207	332	12060	mmunicable
Dengu	A	63	64	11	7	4	2		9	12	13	0	0	1	0	17	0	30	17	8	m	0	10		7	10	2	289	turns of Co
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	Source: Weekly Returns of Communicable Diseases (WRCD).

•T=Timeliness refers to returns received on or before 03<sup>dd</sup> April, 2015 Total number of reporting units 337 Number of reporting units data provided for the current week: 233 C\*\*-Completeness

04<sup>th</sup> April 10<sup>th</sup> 2015

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

### 28th - 03rd April 2015 (14th Week)

Disease			N	o. of Cas	es by P	rovince				Number of cases	Number of cases	Total number of	Total num- ber of	Difference between the
										during current	during same	cases to date in	cases to date in	number of cases to date
	w	С	S	N	E	NW	NC	U	Sab	week in 2015	week in 2014	2015	2014	in 2014& 2015
AFP*	00	00	00	00	00	00	00	00	00	00	02	20	25	-20%
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	%
Mumps	00	00	00	05	01	00	00	01	01	8	08	103	221	-53.4%
Measles	15	01	07	00	00	06	02	01	05	37	67	350	1312	-73.3%
Rubella	00	00	00	00	00	00	00	00	00	00	00	04	06	-33.3%
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	02	%
Tetanus	00	01	00	00	00	00	00	00	00	01	00	04	06	-33.3%
Japanese En- cephalitis	00	00	00	00	00	00	00	00	00	00	00	06	17	-65.1%
Neonatal Teta- nus	00	00	00	00	00	00	00	00	00	00	00	00	00	%
Whooping Cough	00	00	00	01	00	00	00	00	00	01	00	27	16	+69.1%
Tuberculosis	78	13	05	11	01	07	13	06	30	164	131	2591	2883	-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

AFP and all clinically confirmed Vaccine Preventable Diseases except Tuberculosis and Mumps should be investigated by the MOH

## **Dengue Prevention and Control Health Messages**

Look for plants such as bamboo, bohemia, rampe and banana in your surroundings and maintain them

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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. P. PALIHAWADANA CHIEF EPIDEMIOLOGIST EPIDEMIOLOGY UNIT 231, DE SARAM PLACE COLOMBO 10