

WEEKLY EPIDEMIOLOGICAL REPORT A publication of the Epidemiology Unit Ministry of Health, Nutrition & Indigenous Medicine 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

Chikungunya

Vol. 45 No. 51

15th- 21st December 2018

Chikungunya is a mosquito-borne viral disease, first described during an outbreak in southern Tanzania in 1952. It is an RNA virus that belongs to the alphavirus genus of the family Togaviridae. The disease was first described by Marion Robinson and W H R Lumsden in 1955, after an outbreak on the Makonde Plateau, along the border between Tanganyika and Mozambique. Chikungunya is derived from the Makonde/ Kimakonde language which means "that which bends up" or "to become contorted". Due to the stooped posture and appetence developed as a consequence of the arthritic symptoms of the disease it was called so. **Transmission**

Chikungunya is transmitted to humans by the bites of infected female mosquitoes. *Aedes aegypti* and *Aedes albopictus*, the two species of mosquitoes which can also transmit other mosquito-borne viruses, including dengue are the mostly involved in spreading the disease commonly. These mosquitoes can be found biting throughout daylight hours, though there may be peaks of activity in the early morning and late afternoon. Onset of the illness usually occurs between 4 and 8 days after the bite of an infected mosquito which can range from 2 to 12 days.

More about disease vectors

Aedes aegypti and Aedes albopictus have been involved in large outbreaks of chikungunya. Even though Ae. aegypti is limited to the tropics and sub-tropics Ae. albopictus could occur in temperate and even cold temperate regions. Ae. albopictus has spread from Asia to become established in areas of Africa, Europe and the Americas in the recent decades. Species Ae. albopictus flourishes in a wider range of water-filled breeding sites than Ae. Aegypti. Coconut husks, cocoa pods, bamboo stumps, tree holes, rock pools and artificial containers such as vehicle tyres and saucers are the major breading sites for the Ae. albopic*tus* species. Due to the diversity of habitats *Ae. albopictus* is abundantly seen in rural as well as peri-urban areas and shady city parks. *Ae. aegypti* is connected with human habitation. It mainly breeds in indoor breeding sites as flower vases, water storage vessels, concrete water tanks in bathrooms, as well as the same artificial outdoor habitats as *Ae. albopictus*. Several other mosquito vectors including species of the *A. furcifer-taylori* group and *A. luteocephalus* have been implicated in disease transmission in Africa.

Reservoirs

In times of epidemics humans serve as Chikungunya virus reservoirs. During other periods main reservoirs are other vertebrates mainly monkeys, rodents, birds and small mammals. It is reported of outbreaks in monkeys during low heard immunity.

Disease outbreaks

Chikungunya virus can be endemic as well as epidemic. Usually outbreaks begin when the vector density reaches its peak during the rainy season. The disease mainly occurs in Africa, Asia and the Indian subcontinent. Human infections in Africa have been at relatively low levels for a number of years though a large outbreak had occurred in the Democratic Republic of the Congo between 1999–2000 and in Gabon in 2007.

A major outbreak of chikungunya had occurred in the islands of the Indian Ocean from February 2005, due to which cases were imported to many countries including Europe at the peak of the outbreak. A large outbreak of chikungunya in India had occurred in 2006 and 2007. South-East Asian countries such as India, Indonesia, Maldives, Myanmar and Thailand have reported over 1.9 million cases since 2005.Small outbreaks of chikungunya were reported in late 2015 in Dakar, Senegal, and the state of Punjab, India.

Autochthonous transmission of chikungunya was reported in Argentina for the first time following an outbreak of more than 1000 suspected cases

Contents	Page
1. Leading Article – Chikungunya	1
2. Summary of selected notifiable diseases reported (07th- 14th December 2018)	3
3. Surveillance of vaccine preventable diseases & AFP (07th- 14th December 2018)	4

WER Sri Lanka - Vol. 45 No. 51

in 2016. Kenya reported an outbreak of chikungunya resulting in more than 1700 suspected cases while Pakistan continues to respond to an outbreak which started in 2016.

Symptoms

- Characterized by an abrupt onset of fever which can reach up to 39°C often associated with tremors/chills and frequently accompanied by joint pain.
- Petechial or maculopapular rash usually involving the trunk and limbs. Face, palms and sole too can be involved/observed.
- Arthralgia or arthritis affecting multiple joints. Often small joints of the hands, wrists, ankle and feet are affected with pain on movement. Even the large joints can be involved.
- Other common signs and symptoms include muscle pain, headache, nausea and fatigue.

•

Majority of the patients recover fully though for some patients joint pain may persist for several months, or even years. Occasional cases of eye, neurological, gastrointestinal and heart complications have been reported. Serious complications are not common. In older people death can occur. In areas where dengue occurs misdiagnosis or infection being unrecognized can happen.

Diagnosis

Several methods can be used for diagnosis. Serological tests, such as enzyme-linked immunosorbent assays (ELISA), may confirm the presence of IgM and IgG anti-chikungunya antibodies. Three to five weeks after the onset of the illness IgM antibody levels are highest which could persist for about 2 months. Serology and Virological methods (RT-PCR) sould be used as testing for samples collected during the first week after the onset of symptoms.

The virus may be isolated from the blood during the first few days of infection. Various reverse transcriptase–polymerase chain reaction (RT–PCR) methods are available with variable sensitivity. Some are suited to clinical diagnosis. For comparison of various samples of different geographical sources RT–PCR products from clinical samples may also be used for genotyping of the virus.

Treatment

There is no specific antiviral drug treatment for chikungunya. It is mainly directed primarily at relieving the symptoms. The use anti-pyretics, optimal analgesics and plenty of fluids is advisable. Adequate rest is essential while movement or mild exercise will improve the joint stiffness. There is no commercial chikungunya vaccine available at the moment.

Prevention and Control

The closeness of mosquito vector breeding sites to human habitation is a significant risk factor for chikungunya. Prevention and control depends on reducing the number of natural and artificial water-filled container habitats which support breeding of the mosquitoes. During outbreaks, insecticides may be sprayed to kill flying mosquitoes as well as to treat water in containers to kill the immature larvae.

Clothing which minimizes skin exposure to the day-biting vectors is advised for protection during outbreaks. Repellents can be applied to exposed skin or to clothing. Repellents should contain DEET (N, N-diethyl-3-methylbenzamide), IR3535 (3-[N -acetyl-N-butyl]-aminopropionic acid ethyl ester) or icaridin (1piperidinecarboxylic acid, 2-(2-hydroxyethyl)-1methylpropylester).

15th– 21st December 2018

For young children, sick persons or older people insecticidetreated mosquito nets afford good protection. Mosquito coils or other insecticide vaporizers may also reduce indoor biting. Use of screen in rooms to prevent mosquitoes from entering is advised.

Compiled by Dr.T.D. Haputhanthri Epdermiology Unit

Source

WHO chikungunya - <u>https://www.who.int/en/news-room/fact-sheets/detail/chikungunya</u>

Chikungunya virus net- http://www.chikungunyavirusnet.com/ history-of-chikungunya.html

Table 1 : Water Quality SurveillanceNumber of microbiological water samples November 2018									
District	MOH areas	No: Expected *	No: Received						
Colombo	15	90	95						
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	78						
Galle	20	120	39						
Matara	17	102	NR						
Hambantota	12	72	26						
Jaffna	12	72	123						
Kilinochchi	4	24	53						
Manner	5	30	NR						
Vavuniya	4	24	31						
Mullatvu	5	30	NR						
Batticaloa	14	84	0						
Ampara	7	42	40						
Trincomalee	11	66	20						
Kurunegala	29	174	103						
Puttalam	13	78	NR						
Anuradhapura	19	114	44						
Polonnaruwa	7	42	27						
Badulla	16	96	84						
Moneragala	11	66	99						
Rathnapura	18	108	71						
Kegalle	11	66	44						
Kalmunai	13	78	NR						
* No of samples ex NR = Return not n		H area / Month)							

	*J	100	100	100	100	100	100	100	100	100	93	100	100	100	66	100	100	100	100	100	95	88	100	100	100	100	100	66	
WRCD	¥-	61	63	55	59	60	24	31	72	56	36	52	36	56	26	63	63	31	61	63	44	56	48	67	46	64	50	53	
Leishmania- sis	в	ц	62	6	40	183	0	S	741	493	m	7	4	13	2	0	Μ	21	484	4	503	247	12	51	218	17	1	3128	
Leishı sis	A	0	2	0		6	0	0	12	13	0	ч	0	0	0	0	0	0	25	0	14	ч	0	2	4	m	0	87	
gitis	В	68	48	105	42	17	48	59	15	15	13	4	4	6	2	21	34	10	88	89	53	22	142	182	132	48	19	1289	Ŋ
Meningitis	A	0		2	0	0		ч	0	-	0	0	0	0	0	0	m	0	Ч	2	Ч	0	4	7	4	0	0	28	etenes
xodı	в	709	724	708	325	60	205	367	279	295	274	32	28	51	12	189	289	202	598	150	417	316	493	188	317	426	213	7867	**-Compl
Chickenpox	- 4	6	12	16	4	2	0	m	4	9		0	0	Ч	0	2	2	S	4	ъ	4	7	4	m		13	9	114	c 348 C
	B /		0	0		0	0			0	2		0			m		0	2	0	2		0	0	2	0	0	20	ent weel
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	the curr
	В	10	15	16	23	10	28	4	m	24	н	0	1	0	0	7	7	4	25	m	18	4	68	51	30	19		372	provided for
Viral Hepatitis	۲ ۲	0	ч		0	0	0	0	0	ω	0	0	0	0	0	0	0	0	2	0	2	0	2	0	2	0	0	13	its data p
SI "	В	14	10	7	109	Ŋ	142	99	92	65	358	17	12	7	8	m	0	24	29	12	23	1	93	143	29	79	1	1349	eporting un
Typhus Fever	A	0	0	0		0	~		4	2	30		0	0	Ч	0	0	0	2	Ч	0	0		2	0	2	0	55	nber of r
Leptospirosis	В	231	226	684	115	110	50	420	83	283	17	8	1	49	12	59	63	61	363	56	234	168	179	390	738	359	11	4970	iits 353 Nur
Leptos	A	7	4	21	0	9	7	9	4	11	0		0	0	0	9	10		12	Ч	11	13	m	14	24	10	2	16	orting ur
ing	В	43	179	64	30	42	159	25	8	23	224	ъ	2	16	26	33	11	15	13	10	45	20	18	4	S	97	35	1152	mber of rep
Food Poisoning	A	0	11	2	2	0	0	0		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	16	Total nu
	8	46	23	17	9	7	14	9	m	6	54	20	m	23	12	11	m	10	16	7	6	0	14	1	29	11	4	388	nber , 2018
Enteric Fever	۷	m	0	Ч	0		0	0	0	0		0	0	Ч	0	0	0	2	0	Ч	0	0	0	0	0	0	-	:	th Decer
Encephaliti E s	В	6	10	ъ	7		ŋ	14	4	9	9	H	0	4	0	ъ	9	2	19	7	8	Ŀ	10	2	43	13	4	196	r before 14
Ence s	A	0		0	0	0	0	-	0	0	0	0	0	0	0	0	0	0		0	н	0	0	0	0	0	0	4	RCD). (ed on ol
itery	В	96	77	60	118	23	60	64	28	41	214	40	26	18	8	214	82	42	148	92	92	49	144	88	203	64	64	2185	sases (WF urns receiv
Dysentery	A	m	4	1	0	0		0	0	0	ъ	-	0	0	0	16	m	1	2	4	0	0	0	m	7	2	m	56	ble Dist ars to ref
Fever	В	9548	5534	2977	3688	878	202	938	928	1072	3603	321	216	575	114	4746	241	1129	2368	1955	854	308	575	833	2184	1462	1712	48961	s of Communicable Diseases (WRCD). •1=Timeliness refers to returns received on or before 14 th December, 2018 Total number of reporting units 353 Number of reporting units data provided for the current week: 348 C**-Completeness
Dengue Fever	A	292	213	83	75	12	ω	26	33	25	270	2	14	6	S	72	S	53	62	59	13	9	11	20	52	24	30	1474	eturns of Co •T=Tin
RDHS Division		Colombo	paha	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
																												Р	age :

15th- 21st December 2018

WER Sri Lanka - Vol. 45 No. 51

Table 2: Vaccine-Preventable Diseases & AFP

15th– 21st December 2018

07th -14th Dece 2018(50th Week)

Disease	No. of	Cases b	y Provinc	e					Number of cases during current	Number of cases during same	Total num- ber of cases to date in	Total num- ber of cases to date in	Difference between the number of cases to date in		
	W	С	S	N	E	NW	NC	U	Sab	week in 2018	week in 2017	2018	2017	2018 & 2017	
AFP*	00	00	00	00	00	00	00	00	00	00	02	63	68	- 7.3 %	
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0%	
Mumps	01	01	00	00	01	01	00	01	00	05	04	351	289	10.8 %	
Measles	02	01	00	01	00	01	00	01	00	06	08	124	198	- 37.3 %	
Rubella	00	00	00	00	00	00	00	00	00	00	00	08	10	- 20 %	
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	01	0%	
Tetanus	00	00	00	00	01	00	00	00	00	01	00	20	16	25 %	
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %	
Japanese En- cephalitis	00	00	00	00	00	00	00	00	00	00	00	26	28	- 7.1 %	
Whooping Cough	01	00	02	00	01	00	00	00	00	04	00	52	22	136.3 %	
Tuberculosis	53	09	05	20	06	12	15	07	01	128	223	8465	8064	4.9 %	

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.

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													
	Human		Animal										
Month	No Total	No Positive	Infl A Infl B		Pooled samples	Serum Samples	Positives						
December	104	61	28	33	1036	454	0						
S	Sauraa Madigal Dagaarah Instituta & Vatarinary Dagaarah Instituta												

Source: Medical Research Institute & Veterinary Research Institute

PRINTING OF THIS PUBLICATION IS FUNDED BY THE WORLD HEALTH ORGANIZATION (WHO).

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. S.A.R. Dissanayake CHIEF EPIDEMIOLOGIST EPIDEMIOLOGY UNIT 231, DE SARAM PLACE COLOMBO 10

Data Sources: