

Vol. 46 No. 19
$04^{\text {th }}-10^{\text {th }}$ May 2019

## Monkeypox-Part I

Human Monkeypox (MPX)


This is the first of a series of articles on Human Monkeypox

Asian countries are found as free from monkeypox, until the first case was found from Singapore on 9 May 2019, the case was a 38-year-old Nigerian male who arrived in Singapore on 28 April 2019 and attended a workshop from 2930 April. This is the first diagnosed case of monkeypox infection in Singapore. As a country of the Asian region, Sri Lankan should know this infectious diseases as other disease for control and prevention.

## What is monkeypox?

Monkeypox is a viral infectious disease and it was identified in human in a 9 year old boy in the Democratic Republic of Congo (Zaire) in year 1970. Monkeypox (MPXV) is a member of the Orthopoxvirus genus in the family Poxviridae. Monkeypox is a rare viral zoonosis. It is a zoonotic disease and
transmits from animal to human. This virus is an orthopoxvirus that causes a viral disease with symptoms in humans similar, but milder, to those seen in smallpox patients. At that time the case arose in Congo, where smallpox had been eliminated in 1968. Smallpox is not an infectious disease in the world anymore. It was declared eradicated in the world in 1980. Monkeypox occurs sporadically in central and western parts of Africa's tropical rainforest region.

## How a person gets infected?

Human transmission occurs with following situations; direct contact with the blood, body fluids, or cutaneous or mucosal lesions of infected animals (specially handling with infected monkeys), eating inadequately cooked meat of infected animals, close contacts such as infected respiratory tract secretions, skin lesions of an infected person or objects recently contaminated by patient fluids or lesion materials.

Gambian giant rats, squirrels, with rodents are the most likely reservoir of the virus. Though the monkeypox virus is mostly transmitted to people from various wild animals such as rodents and primates, it has limited secondary spread through human-to -human transmission. Congenital monkey-

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pox can occur, with transmissions occurred by inoculation or via the placenta. There is no evidence to date that person-to-person transmission alone can sustain monkeypox in the human population.

## What are the clinical signs and symptoms of this infection?

Interval from infection to onset of symptoms (incubation period) is usually from 6 to 16 days of monkeypox, but can range from 5 to 21 days. Signs and symptoms arise following two infectious periods. It is the invasive period and skin eruption period. Invasive period takes 0-5 days. In this period following signs and symptoms can develop; fever, intense headache, lymphadenopathy (swelling of the lymph node), back pain, myalgia (muscle ache) and an intense asthenia (lack of energy).

Skin eruption period takes 1-3 days after the onset of fever. Various stages of rash appear in this period. It begins on the face and spreads along other parts of the body. Mostly attacked place for rashes of human body is the face of affected cases ( $95 \%$ ), while palms of the hands and soles of the feet of cases are affected (75\%). This rash starts as maculopapules and then becomes vesicles and pustules followed by crusts within 10 days. Disappearance of crusts may occur within three weeks time.

Mucosal lesions can occur in various cases such as oral mucous membranes (in 70\% of cases), genitalia (30\%), and conjunctivae (eyelid) (20\%). Some cases present with affecting cornea of the eye ball. The number of the lesions varies from a few to several thousand.

It is a noticeable feature of severe lymphadenopathy (swollen lymph nodes) before the appearance of the rash, which is a distinctive feature of Monkeypox compared to other similar diseases. Complications can occur in children and adults in debilitated immune status of the body. But, monkeypox is usually a self-limited disease with the symptoms lasting from 14 to 21 days. Documentary evidence shows that case fatality rate of mon-
keypox is less than $10 \%$ and most susceptible are younger age groups.

## How do we diagnose monkeypox?

Other rashes of infectious diseases are considered as differential diagnosis of monkeypox, such as chickenpox, measles, bacterial skin infections, scabies, syphilis, and medication-associated allergies. Lymphadenopathy during the prodromal stage of illness can be a clinical feature to distinguish monkeypox from smallpox which remains eradicated today. Laboratory confirmation of the virus can be done by using optimal samples of lesions. These are vesicular swabs of lesion exudates or crusts which are stored in a dry, sterile tube (no viral transport media) and kept cold until dispatch to the laboratory.

## What is the treatment?

There is no specific treatment or vaccine available for monkeypox. It has been identified that prior smallpox vaccination was highly effective in preventing monkeypox as well. Vaccination against smallpox has been proven to be $85 \%$ effective in preventing monkeypox in the past but the vaccine is no longer available to the general public after it was discontinued following global eradication of smallpox.

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Table 1: Selected notifiable diseases reported by Medical Officers of Health $27^{\text {th }}-03^{\text {rd }}$ May 2019 (18 ${ }^{\text {th }}$ Week)


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Table 2: Vaccine-Preventable Diseases \& AFP

| 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 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 01 | 02 | 32 | 20 | 60 \% |
| Diphtheria | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 0 \% |
| Mumps | 02 | 01 | 00 | 00 | 01 | 03 | 00 | 01 | 00 | 08 | 07 | 135 | 138 | - 2.1 \% |
| Measles | 00 | 03 | 00 | 00 | 00 | 00 | 03 | 01 | 00 | 07 | 01 | 81 | 48 | 68.7 \% |
| Rubella | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 04 | 0 \% |
| 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 | 06 | 10 | - 40 \% |
| Neonatal Tetanus | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 0 \% |
| Japanese Encephalitis | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 01 | 08 | 14 | - 42.8\% |
| Whooping Cough | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 00 | 01 | 01 | 00 | 27 | 15 | 80 \% |
| Tuberculosis | 52 | 31 | 07 | 15 | 02 | 06 | 12 | 16 | 09 | 150 | 55 | 2835 | 2576 | 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

# Dengue Prevention and Control Health Messages <br> Look for plants such as bamboo, bohemia, rampe and banana in your surroundings and maintain them free of water collection. 

## 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@sItnet.lk. Prior approval should be obtained from the Epidemiology Unit before publishing data in this publication

