



WEEKLY EPIDEMIOLOGICAL REPORT

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

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Measles Elimination Strategic Plan Part II

Complications

Complications from Measles have been reported in every organ system. Many of these complications are caused by disruption of epithelial surfaces and immunosuppression and occur in approxi-mately 30% of reported cases depending on age and predisposing conditions. Relatively common complica-tions of measles include otitis mecommon), laryngotracheobronchitis (croup), diarrhoea and pneumonia (most severe). Particularly severe complications of measles which occur in immunocompromised individuals are acute progressive encephalitis (measles inclusion -body encephalitis), and a characteristic giant cell pneu-monia. In developing countries, persistent diarrhoea with proteinlosing enteropathy may ensue, particularly in infants. In these countries, where malnutrition, particularly vitamin A deficiency, and exposure to other infectious diseases are common, the case-fatality rate for measles is usually 3-6%, but can be as high as 30%, particularly among displaced or isolated, immunologi-cally naïve populations. In developed countries death due to measles is rare and the case-fatality rate is usually 0.01-0.1%. The greatest risk of death is in chil-dren younger than 1 year and in adults older than 30 years. In HIV-infected children, the case-fatality rate has been reported to be as high as 50%.

Vitamin A deficiency contributes to delayed recovery and to the high rate of post-measles complications. In addition, measles infection may precipitate acute vita-min A deficiency and xerophthalmia. As a result, measles is an important cause of preventable childhood blind-ness, particularly in Africa.

Diagnosis

The WHO definition of suspected measles is a case with fever and maculopapular (non-vesicular) rash, or a case where a health-care worker suspects measles. Laboratory testing is necessary for definitive diagnosis as other conditions may mimic measles, including infections with rubella virus, parvovirus B19 (erythema infectiosum or Fifth disease), human herpes viruses 6 and 7 (roseola infantum), dengue virus, and Streptococcus pyogenes (scarlet fever). Laboratory confirmation of measles is based on

- Serology
 Anti Measles virus IgM
 IgG Sero conversion / 4-fold rising titre
- Virus Isolation
- Molecular assays
 PCR

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Genotypic characterization

Samples for serology

A venous blood sample of 2 - 3 ml should be collected from each case of suspected measles from the 3rd day to 28th day of the onset of signs and symptoms.

- The skin should be cleaned with 70% alcohol
- The sample should be collected into a sterile, dry, screw-capped bottle without anticoagulant
- Label and leave at room temperature for 30 minutes for clot formation
- specimen should be received by the National Reference Laboratory at MRI Colombo within 48 hours. If there is more than 48 hours delay, the specimen should be refrigerated until dispatch.
- Should be transported in a cold box with ice packs to maintain the temperature at transport at 4C
- Serum should be separated if transport takes several days

Samples for virus isolation

Samples for virus isolation should be collected within the first 5 days of the onset of signs and symptoms.

Virus isolation can be done using Naso-pharyngeal aspirates, Throat Swabs or nasal swabs

Samples should be collected into a container with virus transport medium (VTM) and should be stored immediately in a refrigerator

Should be transported in a cold box with ice packs to maintain the temperature at transport at 4C

Confirmation By reverse transcriptase polymerase chain reaction (RT-PCR) and Immuno fluorescence

Treatment

There is no specific treatment for measles. Case manage-ment of measles focuses on supportive care as well as prevention and treatment of measles complications and secondary infections. Since measles is highly contagious, patient isolation is an important interven-tion to prevent further spread of the virus. However, increasing population immunity through vaccination is the most effective way to prevent outbreaks.

Supportive care should be provided, including relieving common symptoms such as fever, cough, nasal conges-tion or rhinorrhea, conjunctivitis, and sore mouth. Nutritional support is recommended to reduce the risk of malnutrition due to diarrhoea, vomiting and poor appetite associated with measles.

Breastfeeding should be encouraged where appropriate. Oral rehydration salts should be used as needed to prevent dehydration. Antibiotics are generally not recommended for treatment of measles unless secondary bacterial complica-tions develop, such as pneumonia or otitis media.

Vitamin A should be administered to all acute cases irrespective of the timing of previous doses of vitamin A. Vitamin A oral dosage should be given immediately on diagnosis and repeated the next day; 50 000 IU should be given to infants aged <6 months, 100 000 IU to infants aged 6-11 months and 200 000 IU to children aged ≥12 months. If the child has clinical ophthalmic signs of vitamin A deficiency such as Bitot's spots, a third dose should be given 4-6 weeks later. Even in countries where measles is not usually severe, vitamin A should be given in all cases of severe measles

Compiled By;

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Table 1: Water Quality Surveillance Number of microbiological water samples July 2018

District	MOH areas	No: Expected *	No: Received
Colombo	15	90	78
Gampaha	15	90	NR
Kalutara	12	72	NR
Kalutara NIHS	2	12	NR
Kandy	23	138	70
Matale	13	78	31
Nuwara Eliya	13	78	NR
Galle	20	120	79
Matara	17	102	67
Hambantota	12	72	76
Jaffna	12	72	133
Kilinochchi	4	24	33
Manner	5	30	22
Vavuniya	4	24	56
Mullatvu	5	30	NR
Batticaloa	14	84	95
Ampara	7	42	26
Trincomalee	11	66	44
Kurunegala	29	174	114
Puttalam	13	78	42
Anuradhapura	19	114	58
Polonnaruwa	7	42	27
Badulla	16	96	126
Moneragala	11	66	102
Rathnapura	18	108	103
Kegalle	11	66	13
Kalmunai	13	78	0
* N.T. C. 1	. 1 (6 / 1) (6)	r (M 4)	-

* No of samples expected (6 / MOH area / Month)

NR = Return not received

Table 1: Selected notifiable diseases reported by Medical Officers of Health 11th - 17th August 2018(33rd Week)

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Source: Weekly Returns of Communicable Diseases (WRCD).

-Table Transitions of Communicable Diseases (WRCD).

-Table Transitions received on or before 17th August, 2018 Total number of reporting units 353 Number of reporting units data provided for the current week: 351 C**-Completeness

A = Cases reported during the current week. B = Cumulative cases for the year.

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Table 2: Vaccine-Preventable Diseases & AFP

11th - 17th August 2018(33rd Week)

Disease	No. of Case		y Province	Э						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
	W	С	S	N	Е	NW	NC	U	Sab	week in 2018	week in 2017	2018	2017	cases to date in 2018 & 2017
AFP*	00	00	00	01	00	00	00	00	00	01	00	39	45	- 13.3 %
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0%
Mumps	01	01	02	00	00	01	00	00	01	06	03	230	216	6.4 %
Measles	00	00	00	00	00	00	03	00	00	03	08	84	159	- 47.1 %
Rubella	00	00	00	00	00	00	00	00	00	00	00	04	05	- 20 %
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	01	0%
Tetanus	00	00	00	00	00	00	00	00	00	00	00	15	11	36.3 %
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %
Japanese Encephalitis	00	00	00	00	00	01	00	00	00	01	00	21	21	0 %
Whooping Cough	00	00	00	00	00	00	00	00	00	00	00	35	11	218.1 %
Tuberculosis	108	32	15	26	27	21	01	03	06	229	247	5404	5418	- 0.25 %

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

Influenza Surveil	lance in Sentinel	Hospitals - ILI & SARI					
M. d	Human				Animal		
Month	No Total	No Positive	Infl A Infl B Pooled samples Serum Samples Positive	Positives			
August	65	13	07	06	1439	609	0
Source: Medical	Research Institut	e & Veterinary Research Institute					

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ON STATE SERVICE

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