



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>

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Measles Elimination Strategic Plan Part IV

Verifying elimination of Measles

The WHO Global Vaccine Action Plan for 2012–2020 has established the elimination of Measles in at least 5 WHO Regions by 2020 as a target. The five components of the strategy for elimination of Measles are:

1. Achieve and maintain high levels of population immunity by providing high vaccination coverage with two doses of Measles-containing vaccines.
2. Monitor disease using effective surveillance and evaluate programmatic efforts to ensure progress.
3. Develop and maintain outbreak preparedness, respond rapidly to outbreaks and manage cases.
4. Communicate and engage to build public confidence and demand for immunization.
5. Perform the research and development needed to support cost-effective operations and improve vaccination and diagnostic tools.

The achievement of Measles elimination should be verified for individual countries and areas and eventually for each of the WHO Regions, following a standardized process.

Conceptual framework for verifying elimination of Measles

A framework for considering the evidence to be assembled for monitoring progress towards and eventual elimination of Measles includes:

1. Explicit definitions

Endemic Measles virus transmis-

sion:

The existence of continuous transmission of indigenous or imported Measles virus that persists for ≥ 12 months in any defined geographical area

Declaring Measles elimination:

The absence of endemic Measles transmission in a defined geographical area (e.g. region or country) for ≥ 36 months in the presence of a well-performing surveillance system and maintaining high vaccination coverage to maintain high population level immunity

Measles eradication:

Worldwide interruption of Measles virus transmission in the presence of a sensitive surveillance system that has been verified to be performing well.....

Measles outbreak in an elimination setting:

a single laboratory-confirmed case is considered as an outbreak and intensive outbreak response activities need to be carried out

suspected case of Measles:

A patient in whom a health-care worker suspects Measles infection, or a patient with fever and maculopapular (non-vesicular) rash

Laboratory-confirmed Measles case:

suspected case of Measles that has been confirmed by a proficient laborato-

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ry (by serology and virology)

Epidemiologically linked confirmed Measles case:

A suspected case of Measles that has not been confirmed by a laboratory but was geographically and temporally related, with dates of rash onset occurring 7–21 days apart for Measles laboratory-confirmed case or, in the event of a chain of transmission, to another epidemiologically confirmed Measles case

Clinically compatible Measles case:

case with fever and maculopapular (non-vesicular) rash and one of cough, coryza or conjunctivitis, for which no adequate clinical specimen was taken and which has not been linked epidemiologically to a laboratory-confirmed case of Measles or another laboratory-confirmed communicable disease

Non-Measles discarded case:

suspected case that has been investigated and discarded as a non-Measles case using

- (a) laboratory testing in a proficient laboratory or
- (b) epidemiological linkage to a laboratory-confirmed outbreak of another communicable disease that is non- Measles

tion form has been completed and laboratory test results are available, suspected cases should be classified according to the algorithm below

3. Criteria for verifying elimination

Three essential criteria are considered during the time of declaring elimination.

- Documentation of the interruption of endemic Measles virus transmission for a period of at least 36 months from the last known endemic case;
- The presence of a high-quality surveillance system that is sensitive and specific enough to detect imported and import-related cases
- Genotyping evidence that supports the interruption of endemic transmission.

All 3 criteria are necessary for verification of elimination at the regional level to be supported by 5 lines of evidence already described to support these criteria.

Compiled By;
Dr. Saman Pathirana,
 Senior Registrar in community Medicine,
 Epidemiology Unit

2. Case classification system

Countries nearing elimination of Measles should investigate all suspected cases and obtain a clinical specimen for laboratory testing for blood samples for serological testing for Measles IgM and virus detection and genotype identification by throat and nasal swabs in VTM samples. . Once the case investiga-

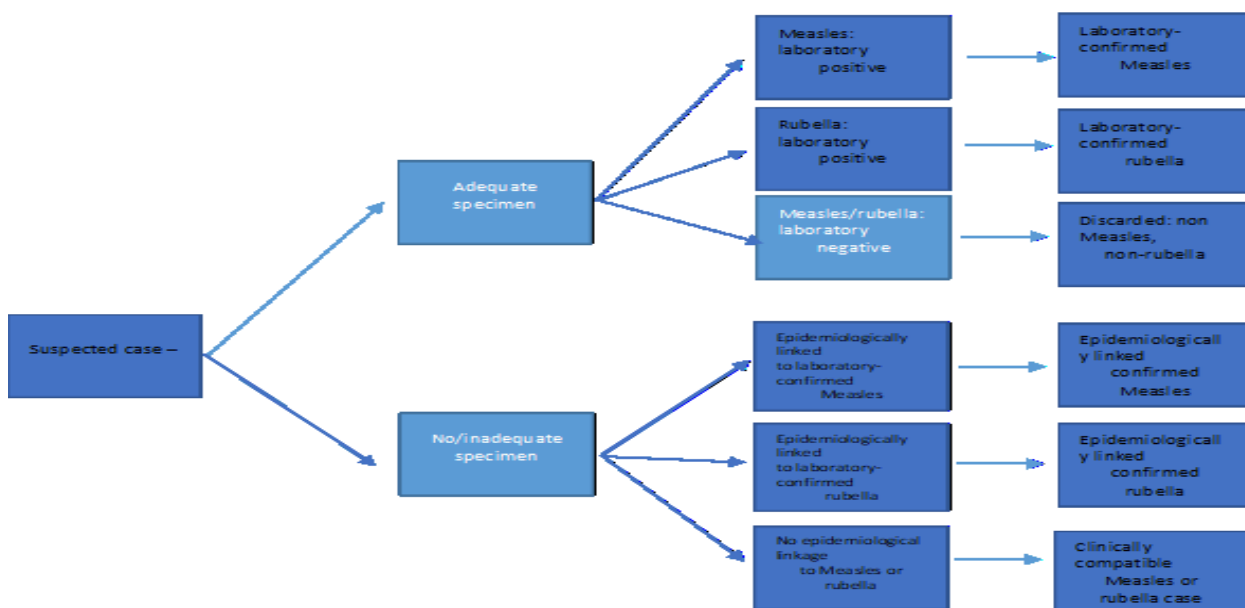


Table 1: Selected notifiable diseases reported by Medical Officers of Health 25th - 31st August 2018(35th Week)

RDHS Division	Dengue Fever		Dysentery		Encephalitis		Enteric Fever		Food Poisoning		Leptospirosis		Typhus Fever		Viral Hepatitis		Human Rabies		Chickenpox		Meningitis		Leishmaniasis		WRCD	
	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	A	B	T*	C**
Colombo	110	7189	2	61	0	7	0	33	0	29	3	143	1	11	0	4	0	0	12	495	0	34	0	2	63	100
paha	95	3943	0	53	0	8	0	17	1	16	7	163	0	4	0	11	0	0	19	536	1	34	2	34	66	100
Kalutara	64	2366	0	64	1	4	0	10	0	53	11	414	1	6	0	11	0	0	11	454	0	73	0	9	53	100
Kandy	54	2667	6	85	0	5	0	3	5	16	5	58	2	80	0	16	0	0	7	253	0	29	1	22	60	100
Matale	9	754	1	16	0	1	0	4	0	31	2	72	0	2	0	6	0	0	3	32	0	11	1	84	61	100
NuwaraEliya	5	154	3	46	0	3	0	9	0	47	2	30	8	110	0	21	0	0	0	180	0	27	0	0	29	100
Galle	13	759	0	39	0	10	0	4	0	9	6	294	4	43	0	2	0	1	4	236	1	45	0	5	22	100
Hambantota	17	692	0	12	0	4	0	2	0	4	4	51	5	54	0	2	0	1	3	194	0	6	5	525	72	100
Matarra	36	796	2	31	0	6	0	5	0	22	7	175	3	37	1	12	0	0	9	214	1	11	14	309	55	100
Jaffna	18	2284	1	116	0	4	0	37	0	212	0	10	1	253	0	1	0	2	6	214	0	9	0	3	37	93
Kilinochchi	2	240	0	23	0	1	1	16	0	2	0	3	0	15	0	0	0	1	0	29	0	2	0	1	51	100
Mannar	0	186	0	17	0	0	0	3	0	2	0	1	0	0	0	0	0	0	0	27	0	4	0	3	38	100
Vavuniya	8	465	0	15	0	4	0	35	0	12	0	31	0	7	0	0	0	1	0	38	0	5	0	8	59	100
Mullaitivu	4	86	0	5	0	0	0	9	1	11	0	8	0	5	0	0	0	1	0	6	0	1	0	2	23	100
Batticaloa	16	4202	5	123	0	5	0	4	0	24	1	39	0	1	0	2	0	3	8	123	0	17	0	0	66	100
Ampara	2	193	0	49	0	3	0	2	0	5	0	33	0	0	0	5	0	1	18	212	1	20	0	2	66	100
Trincomalee	4	906	0	36	0	2	0	4	0	13	3	45	2	21	0	1	0	0	2	159	2	9	0	18	27	100
Kurunegala	28	1888	0	101	0	13	0	13	0	3	0	108	3	20	0	17	1	2	3	383	0	73	7	254	63	100
Puttalam	3	1372	0	32	0	6	0	4	0	4	1	32	0	11	0	2	0	0	5	106	0	61	0	2	63	100
Anuradhapura	6	704	2	40	0	7	0	3	0	38	2	106	0	17	0	9	0	1	3	315	1	31	14	308	44	95
Polonnaruwa	1	241	1	27	0	2	0	0	0	12	2	93	0	0	0	4	0	1	13	193	1	16	6	166	58	88
Badulla	12	407	4	93	1	6	0	7	1	12	0	121	5	61	2	30	0	0	8	353	0	87	0	7	46	100
Monaragala	10	704	0	56	0	2	0	1	0	2	6	224	2	105	1	21	0	0	9	132	9	96	5	34	67	100
Ratnapura	30	1768	4	131	0	36	1	20	0	5	13	490	0	22	2	18	0	2	2	222	1	91	11	167	47	100
Kegalle	26	1106	0	47	0	7	0	6	2	78	13	179	1	57	0	11	0	0	8	263	1	37	0	10	66	100
Kalmune	14	1515	2	31	0	3	0	2	0	31	0	5	0	1	0	1	0	0	0	146	1	10	0	1	51	100
SRILANKA	587	37587	33	1349	2	149	2	253	10	693	88	2928	38	943	6	207	1	17	153	5515	20	839	66	1976	53	99

Source: Weekly Returns of Communicable Diseases (WRCD).

*T=Timeliness refers to returns received on or before 31st 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.

Table 2: Vaccine-Preventable Diseases & AFP

25th – 31st August 2018(35th Week)

Disease	No. of Cases by Province									Number of cases during current week in 2018	Number of cases during same week in 2017	Total number of cases to date in 2018	Total number of cases to date in 2017	Difference between the number of cases to date in 2018 & 2017
	W	C	S	N	E	NW	NC	U	Sab					
AFP*	00	01	00	00	00	01	00	00	00	02	00	43	47	- 8.5 %
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0%
Mumps	01	00	00	00	03	01	00	01	00	06	07	243	226	7.5 %
Measles	00	00	00	00	01	00	00	00	00	01	04	87	169	- 48.5 %
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	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	01	00	00	00	00	01	00	23	21	9.5 %
Whooping Cough	00	00	00	00	00	00	00	00	00	00	01	36	12	200 %
Tuberculosis	154	23	34	08	25	05	21	08	20	298	120	5768	5647	2.1 %

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

Look for plants such as bamboo, bohemia, rampe and banana in your surroundings and maintain them free of water collection.

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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@slt.net.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