

24th– 30th November 2018 Vol. 45 No. 48 Influenza-related deaths investigation – 2018, Southern Province Part IV Therefore, the aetiological agent for the first epi-

sode of illness was not clear for most of the cases.

Number of days from the onset of symptoms to virology sampling was as follows.

Table 14: Time gap between the onset of symptoms to virology sampling (Range, Mean, Median)

No. of days from onset of symptoms to virology sampling	Days
Range	1 day – 28 days
Mean duration	10.5 days
Median duration	8.5 days

The range of days from onset of symptoms to taking samples for virology testing was 1 day to 28 days. Mean duration was 10.5 days. There was a significant delay in taking samples for virology tests from the onset of symptoms. There was a higher possibility of missing the primary source of viral infection as influenza viruses usually gives positive results within the first 5 days of the illness.

The number of days from the admission to the hospital to taking samples for virology testing was as follows.

Table 15: Time gap between the date of admission and virology sampling (Range, Mean, Median) among deceased children

No. of days from admission to virology sampling	Days
Range	0 – 24 days
Mean	7.8 days
Median	4 days

The range of days for taking samples for virology testing from admission to the hospital was 0-24 days. The mean duration was 7.8 days from admission to the hospital. There is a significant delay observed in taking samples for virology. Evidence from virological findings is inadequate to conclude on the initial viral aetiology due to late sampling.

Some of the samples for virology testing has been sent after starting Oseltamivir. Following table depicts the time duration between virology sampling and Oseltamivir treatment.

Table 16: Time gap between virology sampling and starting Oseltamivir treatment

Con	tents	Page
1. L	Leading Article – Influenza-related deaths investigation – 2018, Southern Province Part IV	1
2. S	bummary of selected notifiable diseases reported (17 th – 24 th November 2018)	3
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Virology sample collection is	No.	%
Before starting Oseltamivir	2	10
Same day	1	5
1 day after Oseltamivir	2	10
2 days after Oseltamivir	3	15
3 days after Oseltamivir	3	15
4 days after Oseltamivir	0	
5 days after Oseltamivir	2	10
6 days after Oseltamivir	0	
> 7days after Oseltamivir	5	25
Virology not done	2	10
Total	20	100

Nearly 50% of the cases, virology had been done more than 3 days after the commencement of the treatment with Oseltamivir. There was a high possibility that index samples giving negative results and missing the opportunity to identify the primary source of infection.

Results of the virology sampling are demonstrated in Table 18.

Table 17: Distribution of cases according to virology findings

Virology results	No	%
influenza A	01	5
influenza A + Adeno	02	10
Influenza B+ Adeno	01	5
Influenza A+ Adeno +RSV	02	10
Adeno	03	15
RSV + Adeno	01	5
RSV	03	15
Sample not done	02	10
Sample negative for virus / inconclusive	05	25
Total	20	100

Out of all deceased children, there were **only 6 (30%)** patients who were positive for **Influenza virus infection.** Multiple viruses were isolated from some of the patients. Another 6 patients were positive for RSV. Samples were not sent for virology in 2 patients.

There were 9 positives for Adenoviral infection among these

24th- 30th November 2018

children. The timing of sending samples from the onset of ill-

ness and time gap between starting of Oseltamivir were ana-

lysed among Adeno virus-positive patients as follows.

 Table 18: Distribution of adenovirus positives according to the time gap between virology sampling and onset / starting Oseltamivir

	No. of days from on- set of symptoms to sample collection	No. of days from the start of Oseltamivir to sample collection
Range	5 -28 days	1 -22 days
Mean	13.7 days	7.5 days
Median	8	5

Mean number of days from onset of illness to sample collection was 13.7 days and the number of days from starting Oseltamivir treatment and sample collection was 7.5 days. In summary, adeno positive samples collected average 13 days after onset of symptoms. Reported Adenoviral infections may be a possible secondary viral infection.

Antibiotic treatments:

Several categories of broad-spectrum antibiotics were given to these patients throughout their hospital stay.

Table 19: Distribution of cases according to the number of antibiotics given

	Categories of broad- spectrum antibiotics given to the children	Categories of Anti- biotics
1	Range	2-15
2	Mean	8.5
3	Median	7

All the children were treated with multiple broad-spectrum antibiotics during the illness.

The average number of broad-spectrum antibiotics received by a child was 8.

Compiled by,

Dr. Chintha Jayasinghe Consultant Epidemiologist Epidemiology Unit Ministry of Health

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24th- 30th November 2018

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Table 2: Vaccine-Preventable Diseases & AFP

24th- 30th November 2018

17th-23rd Nove 2018 (47th Week)

Disease	No. of	Cases b	y Province	9					Number of cases during current	Number of cases during same	Total num- ber of cases to	Total num- ber of cases to date in	Difference between the number of	
	W	С	S	N	E	NW	NC	U	Sab	week in 2018	week in 2017	2018	2017	2018 & 2017
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Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %
Mumps	01	01	02	01	01	01	00	01	00	08	06	331	282	17.3 %
Measles	00	00	01	00	01	00	00	00	00	02	03	112	188	- 40.4 %
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	00	00	00	00	00	00	00	19	16	18.7 %
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	25	25	0 %
Whooping Cough	01	00	00	00	00	00	00	00	00	01	00	47	19	147.3 %
Tuberculosis	31	14	49	16	11	71	24	05	48	269	179	7996	7660	4.3 %

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



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ON STATE SERVICE

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