

# WEEKLY EPIDEMIOLOGICAL REPORT

# A publication of the Epidemiology Unit Ministry of Health

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### Vol. 42 No. 31

### 25<sup>th</sup> – 31<sup>st</sup> July 2015

Revised guidelines on Introduction of injectable Inactivated Polio Vaccine (IPV) to the National Immunization Programme

This is the first in a series of two article on introduction of IPV to the National Immunization Programme.

Injectable Inactivated Polio Vaccine (IPV) was introduced into the National Expanded Programme on Immunization (EPI) from1<sup>st</sup> July 2015, in line with the recommended Global Poliomyelitis eradication endgame strategies. Despite the declaration of the certification of South-East Asia as polio-free in March 2014, the risk persists until the disease is eradicated globally.

The requirement of shifting over from trivalent Oral Polio Vaccine (tOPV) to bivalent Oral Polio Vaccine (bOPV) (with Polio Virus type 2 withdrawal plans) during 2016 is universally identified. Wild Polio Virus type 2 (WPV type 2) has been eliminated globally since 1999 but some countries are still experiencing symptomatic cases due to Vaccine Derived Polio Virus type 2 and Vaccine Associated Paralytic Polio type 2 (VDPV2 & VAPP type 2). In fact, ensuring the maintenance of immunity to polio virus type 2 (PV), after withdrawal of PV type 2 from tOPV, introduction of at least one dose of IPV in to the National EPI schedule as an additional dose before shifting over from tOPV to bOPV, is a requirement.

Introduction of IPV will supplement the immunity to polio virus and is recommended in addition to OPV but does not replace OPV. Inactivated Polio Vaccine has been proven to be an extremely safe and effective vaccine but does not produce adequate gut immunity which OPV would provide.

### Introduction of injectable Inactivated Polio Vaccine (IPV) into the National EPI schedule

One dose of IPV should be given to infantsfrom 1<sup>st</sup> of July 2015 <u>on completion of 4 months of age , together with the 2<sup>nd</sup> dose of Pentavalent vaccine and OPV vaccine.</u>

#### Dose, route and site of administration

IPV is liquid suspension, which does not require reconstitution. A single dose of <u>0.5 ml of IPV</u> <u>should be administered by intramuscular route</u> (IM) into the right thigh of the baby.

Sequence of vaccination of infants at 4 months of age (at the clinic)

Step 1: Give OPV first

Step 2: Give IPV to Right thigh

Step 3: Give Pentavalent vaccine to Left thigh

#### Contraindications

Should not vaccinate if:

- Known or documented allergy to vaccine components, including: Streptomycin, Neomycin, Polymyxin B
- History of an allergic reaction following a previous IPV injection

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- Thrombocytopenia (insufficient blood platelets, which play an important role in coagulation)
- Other bleeding disorders
- Anyone with a fever over 38.5°C (101°F)

But IPV can be administered on schedule to immune deficient infants (such as HIV) or infants born prematurely (on completion of 2 months)

Vaccination is better postponed if ;

The recipient is under temporary treatment that suppresses the immune response in which the treatment could reduce immune response to the vaccine.

#### Side effects

Side effects are rare, the most common side effects of the vaccine are redness, swelling and pain at the injection site, fever and discomfort.

Allergic reactions are extremely rare.

#### Storage

- IPV in 5 dose (multi-dose) vials
- IPV should be stored in the upper compartment in the lce lined Refrigerator and middle compartment in the domestic refrigerator.
- IPV should be transported and stored at +2<sup>o</sup> C to +8<sup>o</sup> C temperature. (should not be exposed to heat)
- The vaccine should be kept in the clinic in a container with cool water or inside the form pad of the vaccine carrier to maintain the cold chain (+2<sup>o</sup> C to +8<sup>o</sup>C) and should be protected from direct sunlight.
- IPV should not be stored in the freezer compartment since it is freeze sensitive. (unlike OPV which can be frozen)

It is important to ensure that the vaccine is not frozen. If vaccines are frozen( as indicated by "X" in the freeze tag or any electronic monitor), potency will be lost and will not provide adequate protection against the disease. The "shake test" is ineffective in determining whether IPV has been frozen since it does not contain aluminum as an adjuvant. If there is any suspicion that IPV was frozen, the vial must be discarded.

IPV 5-dose vial can be used under Multi-dose Vial Policy. (MDVP)

Opened IPV 5-dose vial can be used up to 28 days after opening if following criteria are fully met.

The expiry date of the vaccine has not passed. The vaccine vial has been, and will continue to be stored at  $+2^{0}-8^{0}$ C and the Vaccine Vial Monitor (VVM) has not passed its discard point.

Further information on MDVP is available in circular no:01-06/2015 dated 11/02/2005.

#### Injection safety:

Should explain adequately the safety of 2 injections ( IPV and pentavalent vaccines) at a single visit to parent/s of the child vaccinated at 4 months

AD syringes provided in the National EPI programme should be used in vaccine administration and used AD syringes should be discarded into safety boxes provided.

AD syringes and safety boxes will be provided for the National EPI programme by the Regional Medical Supplies Division in coordination with the Epidemiology Unit. Regional Directors of Health Services, Regional Epidemiologists, Medical Officers of Health and Heads of Medical Institutions are responsible for ensuring adequate supply, availability and use of injection safety items at all Immunization clinics in their respective areas.

Appropriate and safe disposal of sharps should be ensured in all instances .

#### Accountability of the IPV

IPV vials are presented as 5-dose vials and measures should be taken to minimize wastage. Significant wastage should be clearly documented and should be reported to both Epidemiology Unit and RDHS office. Open 5-dose multi-dose vial can be used under Multi-dose Vial Policy.

### Compiled by Dr H.H.W.S.B Herath of the Epidemiology Unit

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Table	1:	Sel	ecte	ed n	otif	iabl	e di	sea	ses	rep	ort	ed b	y M	edi	cal	Offi	cers	s of	Hea	lth	1	8 <sup>th</sup>	-24	th J	uly	201	5 (3	0 <sup>th</sup>	Week)
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RDHS Division		Colombo	Gampaha	Kalutara	Kandy	Matale	NuwaraEliya	Galle	Hambantota	Matara	Jaffna	Kilinochchi	Mannar	Vavuniya	Mullaitivu	Batticaloa	Ampara	Trincomalee	Kurunegala	Puttalam	Anuradhapura	Polonnaruwa	Badulla	Monaragala	Ratnapura	Kegalle	Kalmunei	SRILANKA	Source: Weekly Ret A = Cases reported du
Page	3																												

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### Table 2: Vaccine-Preventable Diseases & AFP

# 25<sup>th</sup> July 31<sup>st</sup> 2015

18 <sup>th</sup> - 24 <sup>th</sup>	Jul	2015	(30 <sup>th</sup> V	Veek
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Disease			Ν	lo. of Cas	es by P	rovince		Number of cases during current	Number of cases during same	Total number of cases to	Total num- ber of cases to	Difference between the number of			
	w	С	S	N	Е	NW	NC	U	Sab	week in 2015	week in 2014	date in 2015	date in 2014	cases to date in 2014& 2015	
AFP*	01	01	00	00	00	00	00	00	00	02	01	-13.4%			
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0%	
Mumps	02	01	00	00	00	00	00	03	00	06	15	234	444	-47.2%	
Measles	46	02	07	01	03	07	03	04	05	78	39 1588 229		2297	-31.1%	
Rubella	01	00	00	00	00	00	00	00	00	01	00	07	13	-46.1%	
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	04	-100%	
Tetanus	00	00	00	00	00	00	00	00	00	00	00	11	09	+22.2%	
Neonatal Teta- nus	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	07	19	-63.1%	
Whooping Cough	00	00	01	01	01	00	00	00	00	03	00	54	33	+63.6%	
Tuberculosis	47	07	13	12	07	06	08	00	31	131	210	5449	5731	-5.1%	

#### 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

AFP and all clinically confirmed Vaccine Preventable Diseases except Tuberculosis and Mumps should be investigated by the MOH

### **Dengue Prevention and Control Health Messages**

Look for plants such as bamboo, bohemia, rampe and banana in your surroundings and maintain them

### 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**

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