

# WEEKLY EPIDEMIOLOGICAL REPORT

# A publication of the Epidemiology Unit Ministry of Healthcare and Nutrition

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# 20<sup>th</sup> - 26<sup>th</sup> February 2010

# Guidelines on Immunization against tetanus

### (A) Introduction

Tetanus is a fatal infectious disease caused by toxigenic strains of Clostredium tetani. Tetanus and neo- natal tetanus is still a major public health problem in a considerable number of developing countries. However, both tetanus and neo- natal tetanus have reached elimination levels in Sri Lanka as a result of the successful immunization programme.

Protection against tetanus could be achieved either through active immunization (by giving tetanus toxoid containing vaccine) or by passive immunization (by giving tetanus specific immunoglobulin). Neonatal tetanus is prevented by immunizing the pregnant mothers with tetanus toxoid vaccine.

Tetanus vaccines are based on tetanus toxoid which is a chemically inactivated tetanus toxin which could induce production of antibodies against tetanus toxin. Tetanus toxoid is available as mono valent tetanus toxoid (TT), combined with diphtheria, pertussis, hepatitis B and Hib vaccines (pentavalent vaccine), combined with diphtheria and pertussis vaccines (DPT), combined with diphtheria (DT) or as low dose diphtheria Toxoid (aTd).

The goal of Tetanus immunization in the EPI programme is to eliminate tetanus and neonatal tetanus from Sri Lanka. With the emergence of new knowledge on immunological response of patients after receiving tetanus toxoid, the following recommendations have been made regarding immunization with tetanus containing vaccines.

### (B) Recommended schedule (pre exposure vaccination)

### Immunization from infancy up to adolescence

Dose	Age (on completion of)	Type of vaccine	Minimum duration to be kept between different doses of tetanus containing vaccine when tetanus toxoid containing vaccines have not been re- ceived according to the scheduled age							
1st	2 months		6-8 weeks between							
2nd	4 months	tavalent	$2^{nd}$ and the $3^{rd}$ dose							
3rd	6 months									
4th	18 months	DPT	12 months between 3 <sup>rd</sup> and 4 <sup>th</sup> doses							
5th	5 years	DT	3 years between 4 <sup>th</sup> and 5 <sup>th</sup> doses							
6th	12 years ( in grade 7, in school medical inspection)	aTd	years between 5 <sup>th</sup> and 6 <sup>th</sup> doses							

### 2) Immunization during pregnancy

The number of doses required and timing of boosters during pregnancy will depend on the past immunization history of the pregnant mother with tetanus containing vaccines.

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Immunization of pregnant mothers who have not received tetanus containing vaccines in infancy and childhood as per the EPI schedule

Pregnant mothers who have not received tetanus containing vaccine according to the EPI schedule during infancy and childhood should be immunized according to the following schedule.

1 <sup>st</sup> dose	=	During 1 <sup>st</sup> pregnancy after 12 weeks of gestation
2 <sup>nd</sup> dose	=	During 1 <sup>st</sup> pregnancy (6-8 weeks after the first dose, two weeks before the delivery)
3 <sup>rd</sup> dose	=	During 2 <sup>nd</sup> pregnancy
4 <sup>th</sup> dose	=	During 3 <sup>rd</sup> pregnancy
5 <sup>th</sup> dose	=	During 4 <sup>th</sup> pregnancy
Not indicated		Pregnant mothers who have received 5 doses of tetanus toxoid during previous pregnancies do not

Immunization of pregnant mothers who have documented evidence of receipt of six doses of tetanus containing vaccines as per the national EPI schedule (3 doses of DPT/ penta in infancy + DPT at 18 months + DT at 5 years + aTd at 12 years)

One booster dose (TT)	=	If the gap between the 6 <sup>th</sup> dose of tetanus containing vaccine or any subsequent tetanus contain- ing vaccine and the current pregnancy is more than 10 years
		If the gap between the 6 <sup>th</sup> dose of tetanus containing vaccine and the current pregnancy is less than 10 years
Not indicated		If the gap between the subsequent tetanus containing vaccine dose after the 6 <sup>th</sup> dose and the
	=	current pregnancy is less than 10 years

#### © Post exposure vaccination

Decision on post exposure vaccination should be taken after considering the nature of the lesion and the previous history of the immunization with tetanus containing vaccine.

#### Immunization of persons who have been immunized with tetanus containing vaccine during infancy and childhood

If a person has documented evidence of receipt of six doses of tetanus containing vaccine (with 4 doses of DPT/ penta, DT, aTd) he/she does not need to be immunized with tetanus toxoid up to 10 years after the 6<sup>th</sup> dose of tetanus containing vaccine. If a patient presents 10 years after the 6<sup>th</sup> dose of tetanus containing vaccine a booster dose should be given. If a patient presents with a severely contaminated wound but less than 10 years from the 6<sup>th</sup> dose of tetanus containing vaccine a booster dose of tetanus containing vaccine a booster dose of tetanus containing vaccine a booster dose of tetanus toxoid can be given with the discretion at the treating physician.

#### Immunization of persons who have not been immunized with tetanus containing vaccine according to the EPI schedule

Any person who has not been immunized with tetanus containing vaccine during infancy and adolescence according to the national EPI schedule, he / she should be given a dose of tetanus toxoid  $(1^{st} dose)$  if there is a risk of developing tetanus. Second dose of tetanus toxoid should be given 4 weeks after the  $1^{st}$  dose and the third dose should be given 6 months after the second dose. A booster dose  $(4^{th})$  of tetanus could be given after 5 years. Fifth dose given 10 years after the  $4^{th}$  dose will produce a long lasting, probably a life long immunity.

#### (D) Development of immunity after immunization with tetanus containing vaccines

If there is documentary evidence to show that any infant, child or an adult have been immunized as per EPI schedule, it is not necessary to immunize them in between the routine doses again whenever they present with trauma as they are protected against tetanus infection in between the doses.

#### (E) Storing and administering tetanus containing vaccines

Tetanus containing vaccines should be stored and transported between + 2C and+8 C and should not be exposed to freezing as freezing can reduce the potency of these vaccines. Opened tetanus containing vaccines should be stored between + 2C and+8 C and could be reused in subsequent immunization sessions within 4 weeks of opening of the vaccine vials. Tetanus containing vaccines should be given intra muscularly.



# Table 1: Vaccine-preventable Diseases & AFP

# 13th - 19th February - 2010(07th Week)

20<sup>th</sup> - 26<sup>th</sup> February 2010

Disease			1	No. of Cas	ses by P	rovince		Number of cases during current	Number of cases during same	Total number of cases to date in	Total num- ber of cases to date in 2009	Difference between the number of cases to date			
	W	С	S	N	E	NW	NC	U	Sab	week in 2010	week in 2009	2010		in 20010 & 2009	
Acute Flaccid Paralysis	02	01	03	00	00	00	00	00	01	07	00	16	08	+100%	
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	-	
Measles	00	00	00	00	00	00	00	00	01	01	00	21	15	+ 40.0 %	
Tetanus	00	00	00	00	00	00	00	00	00	00	01	04	05	- 20.0 %	
Whooping Cough	00	00	00	00	00	00	00	00	00	00	01	03	12	- 75.0 %	
Tuberculosis	25	14	41	16	11	12	00	13	09	141	177	1331	1148	+ 16.0 %	

# Table 2: Newly Introduced Notifiable Disease

### 13th - 19th February - 2010(07th Week)

Disease			l	No. of Ca	ases by	Provinc	е	Number of	Number of	Total	Total num-	Difference			
	W	С	S	N	E	NW	NC	U	Sab	cases during current week in 2010	cases during same week in 2009	cases to date in 2010	ber of cases to date in 2009	number of cases to date in 2010 & 2009	
Chickenpox	17	05	13	06	01	05	15	06	08	76	161	496	1081	- 54.1 %	
Meningitis	03 KT=1 CB=1 GM=1	00	02 GL=1 MT=1	00	00	07 KR=5 PU=2	01 AP=1	00	07 KG=2 RP=5	20	17	268	142	+ 88.7 %	
Mumps	01	03	05	01	01	01	02	00	04	18	34	120	265	- 54.7 %	
Leishmaniasis	00	00	02 MT=2	00	00	00	06 AP=6	00	00	08	14	54	50	+ 08.0 %	

#### Key to Table 1 & 2

W: Western, C: Central, S: Southern, N: North, E: East, NC: North Central, NW: North Western, U: Uva, Sab: Sabaragamuwa. DPDHS 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:

Provinces:

Weekly Return of Communicable Diseases: Diphtheria, Measles, Tetanus, Whooping Cough, Chickenpox, Meningitis, Mumps.

Special Surveillance: Acute Flaccid Paralysis. Leishmaniasis is notifiable only after the General Circular No: 02/102/2008 issued on 23 September 2008.

10th South East Asia Regional Scientific Meeting of the International Epidemiological Association 23rd - 26th May 2010

### Colombo, Sri Lanka

Theme

## "Epidemiological Methods in Evidence Based Healthcare"

### Visit http://www.episea2010.com

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# Table 4: Selected notifiable diseases reported by Medical Officers of Health

13th - 19th February - 2010(07th Week)

DPDHS Division	Denç ver /	gue Fe- ' DHF*	Dyse	entery	Enc	ephali tis	En Fo	teric ever	F Poi:	ood soning	Lepto	ospiros is	Ty F	phus ever	Vi Hep	ral atitis	Hu Ral	man pies	Returns Re- ceived
	А	В	А	В	А	В	А	В	А	В	Α	В	А	В	А	В	А	В	%
Colombo	136	954	2	23	1	3	1	12	0	5	16	61	0	2	2	12	0	1	85
Gampaha	86	1004	1	6	1	6	0	5	1	1	17	56	0	0	1	17	0	1	67
Kalutara	22	207	4	30	0	3	0	4	0	6	1	28	0	0	1	7	0	0	92
Kandy	32	319	0	57	0	0	0	2	0	1	0	9	1	27	2	15	0	2	87
Matale	28	202	102	128	0	0	0	6	33	36	3	20	0	0	2	10	0	0	83
Nuwara	4	38	1	13	0	0	2	22	0	0	0	4	1	15	4	10	0	0	100
Galle	19	94	1	28	01	1	0	0	0	4	0	2	0	1	2	4	0	1	95
Hambant	40	138	0	7	0	2	1	1	0	0	3	17	1	28	1	2	0	0	100
Matara	15	71	1	19	0	1	0	1	0	34	7	23	3	41	0	5	0	0	94
Jaffna	83	1557	1	28	0	1	18	166	0	4	0	0	8	73	1	11	0	0	67
Kili-	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Mannar	5	42	2	11	0	0	1	17	0	0	0	0	0	0	1	8	0	0	60
Vavuniya	22	414	0	10	0	1	3	19	0	0	0	0	0	0	1	4	0	0	75
Mullaitivu	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Batticaloa	57	488	3	18	1	1	0	5	0	9	0	1	0	1	0	0	0	0	92
Ampara	0	20	2	17	0	0	0	2	2	6	0	12	0	0	0	5	0	0	71
Trincomal	12	430	7	38	0	2	0	2	0	1	0	6	0	4	0	5	1	1	60
Kurunega	15	291	3	48	0	3	1	7	0	0	15	68	1	12	3	15	0	0	80
Puttalam	23	360	0	17	0	2	0	21	0	0	1	7	0	0	0	0	0	0	89
Anuradha	35	482	0	14	0	0	0	2	0	0	2	7	3	9	0	10	0	3	68
Polonnar	11	62	3	16	1	1	0	0	1	2	1	23	0	0	2	13	0	0	100
Badulla	13	125	7	36	0	0	5	19	0	6	2	14	2	12	1	12	0	0	87
Monaraga	9	73	3	43	0	0	0	14	0	1	1	10	1	9	0	1	0	0	73
Ratnapur	32	177	6	51	0	3	0	4	0	8	9	51	0	19	2	29	0	3	67
Kegalle	33	176	0	11	0	4	0	12	0	2	4	34	1	4	3	22	0	0	100
Kalmunai	38	277	4	29	0	0	1	3	0	0	0	0	0	0	0	5	0	0	77
SRI LANKA	770	8001	153	698	05	34	33	346	37	126	82	453	22	257	29	222	01	10	80

Source: Weekly Returns of Communicable Diseases WRCD).

\*Dengue Fever / DHF refers to Dengue Fever / Dengue Haemorrhagic Fever. \*\*Timely refers to returns received on or before 19th February, 2010 Total number of reporting units =311. Number of reporting units data provided for the current week: 255 A = Cases reported during the current week. B = Cumulative cases for the year.

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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@sltnet.lk.

# **ON STATE SERVICE**

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