

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

Vol. 43 No. 36

27th – 02nd September 2016

Trachoma

Introduction

Trachoma is the leading infectious cause of blindness in the world which is a major public health concern in 42 countries. Trachoma is caused by the intracellular bacterium, *Chlamydia trachomatis*. Isolated infection is usually self limiting. Repeated infection leads to blindness. Trachoma is entirely a preventable cause of blindness.

Burden

Among the 42 endemic countries are countries in Africa, Central and South America, Asia, Australia and Middle East. Out of them, Africa is the most affected and the place where most intense control efforts are in place. Infection is particularly prevalent in dry, dusty regions and settings where hygienic practices are poor, overcrowded and poverty ridden. Just over 200 million people live in these endemic areas making them particularly vulnerable to get infected. In these endemic areas infection is common in childhood and infection usually subsides in adolescence. These children present with varying degrees of scarring.

As the leading preventable cause of blindness Trachoma is responsible for blindness in 1.9 million people. This accounts for 1.4% of the total number of people with blindness globally. In 2015 alone, in Africa, 176,000 underwent surgery for Trachoma and 54 million were treated with antibiotics.

Apart from the burden on health sector, Trachoma poses a significant impact on economy as well. Lost productivity due to blindness and visual impairment is estimated to be between 2.9 billion US\$ to 5.3 billion US\$.

Pathogenesis

Chlamydia trachomatis is a gram negative obligate intracellular pathogen. Infection with this pathogen causes inflammation of the conjunctiva characterized by lymphatic and monocyte infiltration with plasma cells and macrophages in follicles. Tissue damage is immunologically mediated. Repeated infection leads to sustained inflammation which ultimately ends up in scarring (trachomatous conjunctival scarring). Repeated infection is common in disease endemic communities.

In the process of scarring, the normal, loose vascular subepithelial stroma gets replaced by thick compact bands of type IV and type V collagen. Due to scarring conjunctival epithelium gets atrophied. It also leads to loss of goblet cells in the conjunctival epithelium.

Scarring leads to contraction and buckling of the tarsal plate of the upper eyelid producing entropian and trichiasis. In entropian the eyelid turns inward and in trichiasis eyelashes get introverted, due to which the eyelid rubs on cornea causing pain, discomfort and corneal damage. If left untreated this leads to formation of irreversible opacities and visual impairment.

Contents	Page
1. Leading Article – Trachoma 2. Summary of selected notifiable diseases reported -(20 th – 26 th August 2016) 3. Surveillance of vaccine preventable diseases & AFP -(20 th – 26 th August 2016)	1 3 4

Transmission

Incubation period of *Chlamydia trachomatis* infection is 5 to 12 days. It can spread directly or indirectly. The ocular discharges and nasopharyngeal secretions of the infected people contain the bacteria. These secretions can contaminate other individuals directly. On the other hand, secretions can contaminate other sites like clothes, towels etc. and indirectly spread the infection. Apart from that, particular types of flies, especially *Musca sorbens* aid in disease transmission, mostly in Middle East. Infected individuals are contagious as long as active lesions are present in conjunctiva and adnexsal mucosa.

Clinical features

There are two phases in the clinical course of Trachoma— active phase and scarring/ cicatricial phase. These two phases can co— exist. Most of the patients in the active phase are relatively asymptomatic. However, they can sometimes show symptoms similar to follicular conjunctivitis which include watery discharge, redness, photophobia and pain. Dry eyes is a feature of scarring phase as scarring destroys mucus and serous glands. Apart from that irritating foreign body sensation in the eyes and blepharospasms (involuntary tight closure of the eye lids) can be present. Trichiasis, entropian, corneal opacities are other unique features of the scarring phase.

WHO simplified Trachoma grading system

TF- Trachomatous Inflammation – Follicular

The presence of five or more follicles in the upper tarsal conjunctiva

TI - Trachomatous Inflammation - Intense

Pronounced inflammatory thickening of the tarsal conjunctiva that obscures more than half of the normal deep tarsal vessels

TS - Trachomatous Scarring

The presence of scarring in the tarsal conjunctiva

TT - Trachomatous Trichiasis

At least one eyelash rubs on the eyeball or evidence of recent removal of in turned eyelashes

CO - Corneal Opacity

Easily visible corneal opacity over the pupil

Diagnosis

History suggesting recent dwelling in an endemic area along with relevant clinical presentation aid in diagnosis. Laboratory investigations can be done for its confirmation. Polymerase Chain Reaction (PCR), Giemsa—stained smears for the detection of intracellular chlamydial elementary bodies in epithelial cells of conjunctival scrapings, immunofluorescence examina-

tion after methanol fixation of the smear, detection of chlamydial antigen by enzyme immunoassay or deoxyribonucleic acid by probe, isolation of the agent in special cell culture are some of the investigations that can be done. However, these methods are not readily available and feasible in resource poor settings, especially, in disease endemic areas.

Risk factors

Children are more at risk of developing Trachoma. Apart from that, women are affected up to four times as often as men mainly because of their close contact with infected children. Environmental risk factors include poor hygiene, crowded households, exposure to dry wind, dust and fine sand, water shortage and inadequate latrine and sanitary facilities.

Treatment

"SAFE" (S- Surgical treatment, A- Antibiotic therapy, F- Facial cleanliness, E- Environmental improvement) is the strategy developed by WHO, indicating the key components of the treatment of Trachoma.

Eyelid surgery is done in patients with trichiasis and entropian. Surgical management reduces the rate of progression of corneal scarring. It also improves visual acuity for some degree by restoring visual surface and reducing ocular secretions and blepharospasms. Oral Azithromycin and Tetracycline are the recommended antibiotic therapy.

Research evidence suggest that facial cleanliness in children reduces both the risk and severity of active Trachoma.

Environmental improvement include improved water supplies, improved household sanitation especially safe disposal of human feces.

All theses practices need to be complimented with improvement of knowledge, attitudes and practices in the community which are necessary to identify and control Trachoma.

Sources

World Health Organization official web site

Center for Disease Control and Prevention official website

Control of Communicable diseases Manual, 20th edition

Compiled by Dr. S.A.I.K. Sudasinghe of the Epidemiology Unit

Ω

⋖

Ω

⋖ \sim

Ω

⋖

Ω

⋖

Ω /

 \forall

Ω

⋖

Ω

⋖ \sim

Ω

⋖

Ω

⋖

Δ

⋖

Ω

Meningitis

Chickenpox

Human Rabies

Viral Hepatitis

Typhus Fever

Leptospirosis

Food Poisoning

Enteric Fever

Encephaliti

Dysentery

Dengue Fever

⋖

Colombo

Gampaha

/

_ $^{\circ}$ _

_

 ∞

9/

Kalutara

=

Matale Kandy

 ∞

/

ω

Galle

Hambantota

 ∞

NuwaraEliya

 \sim

_

Matara

 ∞

Jaffna

 \sim

Kilinochchi

/

WER Sri Lanka - VOI. 43 NO. 36 2711- U2114 September 2016															2016														
Table 1: Selected notifiable diseases reported by Medical Officers of Health 20th - 26th Aug 2016 (35th Week)															Neek)														
WRCD	*.	88	73	100	100	100	100	85	92	100	92	100	100	100	100	98	7.1	83	67	79	89	86	82	91	89	100	85	91	
M	<u>*</u>	75	33	64	87	77	77	20	83	100	92	75	09	75	80	11	14	67	76	69	42	57	71	82	50	73	54	89	
Leishmani- asis	В	0	7	0	8	17	0	က	247	156	1	0	0	9	2	_	Ω	2	99	33	174	66	3	33	1	2	0	842	
Leish asis	⋖	0	0	0	0	0	0	0	13	4	0	0	0	0	0	0	0	0	m	0	4	_	0	0	0	0	0	25	

 ∞

_

 \sim

=

Mannar

 Ampara

_

_

Puttalam

 ∞

Anuradhapura Polonnaruwa

Kurunegala

 \sim

Frincomalee

 \sim

/

Batticaloa Mullaitivu Vavuniya

ω

Badulla

-

_ _ $\overline{}$

_

Ratnapura

ω

Monaragala

-

SRILANKA

 $^{\circ}$

Kalmune

Kegalle

· T=Timeliness refers to returns received on or before 26* August, 2016 Total number of reporting units 339 Number of reporting units data provided for the current week: 313 C**-Completeness A = Cases reported during the current week. B = Cumulative cases for the year. Source: Weekly Returns of Communicable Diseases (WRCD),

Page

Table 2: Vaccine-Preventable Diseases & AFP

20th - 26th Aug 2016 (35th Week)

Disease				No. of Ca	ses by F	Province	9		Number of cases during current	Number of cases during same	Total number of cases to	Total num- ber of cases to date in	Difference between the number of cases to date		
	W	С	S	N	Е	NW	NC	U	Sab	week in 2016	week in 2015	date in 2016	2015	in 2016 & 2015	
AFP*	00	00	00	00	00	01	00	00	00	01	02	50	51	-2.1%	
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0%	
Mumps	00	01	01	00	00	00	01	01	00	04	05	279	260	+7.3%	
Measles	02	00	00	01	00	00	00	01	00	04	67	316	2009	-84.2%	
Rubella	00	00	00	00	00	00	00	00	00	00	00	07	08	-12.5%	
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	00	0%	
Tetanus	01	00	00	00	00	00	00	00	00	01	00	08	14	-43.1%	
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	13	07	+85.7%	
Whooping Cough	01	00	00	01	00	00	00	00	01	03	00	49	59	-17.1%	
Tuberculosis	92	08	18	24	15	04	16	08	04	189	213	6452	6763	-4.5%	

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