EUROPEAN JOURNAL OF PHARMACEUTICAL AND MEDICAL RESEARCH

www.ejpmr.com

Review Article ISSN 2394-3211 EJPMR

IMPLEMENTATION OF DOTS AS PER REVISED NATIONAL TUBERCULOSIS CONTROL PROGRAMME (RNTCP) IN DADRI REGION OF UTTAR PRADESH

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Article Received on 20/12/2018

Article Revised on 10/01/2019

Article Accepted on 31/01/2019

ABSTRACT

DOT (Directly Observed Treatment) against tuberculosis under the guidance of the revised national tuberculosis control programme (RNTCP) has already covered 450 million population of the country and has succeeded in achieving an overall cure rate of 80% for new smear-positive tuberculosis (TB) cases. For more than a decade, designated microscopy health centers and medical hospitals have been providing diagnostic services, treatment, referral for treatment, recording and reporting data, carrying out advocacy for RNTCP and conducting operational research relevant to RNTCP. Health centers are also contributing to diagnosis and treatment of human immunodeficiency virus (HIV)-TB co-infection and development of laboratory infrastructure for early diagnosis of multidrug-resistant and/or extensively drug-resistant TB (M/XDR-TB) and DOTS-Plus sites for treatment of MDR-TB cases. It tries to evoke a uniform consciousness among treating doctors both at government and private levels to follow a uniform diagnostic algorithm and treatment protocols which is very much readily available under RNTCP-DOTS throughout our nation. The objective of the study is to overview the guidelines put by RNTCP in various health centers of Dadri region of Uttar Pradesh for the evaluation and treatment of suspected TB cases.

KEYWORDS: DOT, Tuberculosis, Dadri, RNTCP.

INTRODUCTION

Tuberculosis (TB) is a major public health problem in India.^[1] India accounts for one-fifth of the global TB incident cases. Each year nearly 2 million people in India develop TB, of which around 0.87 million are infectious cases. It is estimated that annually around 330,000 Indians die due to TB.^[2] Since 1993, the Government of India has been implementing the WHOrecommended DOTS strategy via RNTCP. The revised strategy was pilot-tested in 1993 and launched as a national programme in 1997. By March 2006, the programme was implemented nation-wide in 633 districts, covering 1114 million (100%) population. Phase II of the RNTCP started from October 2005, which is a step towards achieving the TB-related targets of the Millennium Development Goals. Since 2006, RNTCP is implementing the WHO recommended "Stop TB Strategy", which in addition to DOTS, addresses all the newer issues and challenges in TB control.^[3,4]

The objectives of RNTCP are

- To achieve and maintain at least 85% cure rate amongst New Smear Positive (NSP) pulmonary TB cases.
- To achieve and maintain at least 70% detection of such cases.

RNTCP program provides, free of cost, quality antitubercular drugs across the country through the numerous Primary Health Centres and the growing number of private-sector DOTS-providers. Tuberculosis is an infectious disease caused by T.B bacteria (Mycobacterium tuberculosis and Mycobacterium Bovis) that primarily affect the lungs but it can also affect organs in the central nervous system, lymphatic system, and circulatory system among others. Tuberculosis mostly affects young adults, in their most productive years. However, all age groups are at risk. Over 95% of cases and deaths are in developing countries. People who are co-infected with HIV and TB are 21 to 34 times more likely to become sick with TB (see TB and HIV section). Risk of active TB is also greater in persons suffering from other conditions that impair the immune system.^[5,6] About half a million children (0-14 years) fell ill with TB, and 64 000 children died from the disease in 2011. Tobacco use greatly increases the risk of TB disease and death. More than 20% of TB cases worldwide are attributable to smoking.^[7]

MATERIALS AND METHODS Study area and population

The study was conducted in Dadri a medium sized town of district Greater Noida, Uttar Pradesh, India. The RNTCP has been operational in Dadri region and a survey was carried out by direct interview with patients and doctors of few private and government hospitals. 414 Patients of both sex were interviewed randomly from a age group of 10 to more than 70 years old. Doctors from 10 private hospitals and 2 government hospitals were also randomly interviewed about same.

Findings

Recommended treatment

Standardized treatment regimens are one of the pillars of the DOTS strategy Isoniazid, Rifampicin, Pyrazinamide,

Table 1: Recommended	Treatment f	or TB. ^[11,12]
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Ethambutol, and Streptomycin are the primary antitubercular drugs used. Most DOTS regimens have thrice-weekly schedules and typically last for 6 to 8 months, with an initial intensive phase and a continuation phase.^[8]

Based on the nature/severity of the disease and the patient's exposure to previous anti-tubercular treatments, RNTCP classifies tuberculosis patients into two treatment categories.

Table 1: Recommended Treatment for TD.	
New	Previously treated
New sputum smear-positive,	Sputum smear-positive relapse,
New sputum smear-negative,	Sputum smear-positive failure,
New extrapulmonary tuberculosis,	Sputum smear-positive treatment after default,
Others	others#
$2\mathbf{H}_3\mathbf{R}_3\mathbf{Z}_3\mathbf{E}_3 + 4\mathbf{H}_3\mathbf{R}_3$	$2\mathbf{H}_{3}\mathbf{R}_{3}\mathbf{Z}_{3}\mathbf{E}_{3}\mathbf{S}_{3} + I\mathbf{H}_{3}\mathbf{R}_{3}\mathbf{Z}_{3}\mathbf{E}_{3} + 5\mathbf{H}_{3}\mathbf{R}_{3}\mathbf{E}_{3}$
2 months Intensive phase + 4 months continuation phase	3 months Intensive phase + 5 months continuation phase
Four drugs at Thrice-weekly Schedule for 2 months	Five drugs at Thrice-weekly Schedule for initial 2 months
Intensive phase & Two drugs at Thrice-Weekly Schedule	followed by Four drugs for next 1 month Intensive phase.
for remaining 4 months continuation phase.	Three drugs at Thrice-weekly Schedule for remaining 5
	months continuation phase.

H: Isoniazid (600 mg), R: Rifampicin (450 mg),

Z: Pyrazinamide (1500 mg), E: Ethambutol (1200 mg),

S: Streptomycin (750 mg).

- Patients who weigh 60kg or more receive additional 1. Rifampicin 150mg.
- Patients who are more than 50 years old receive 2. Streptomycin 500mg. Patients who weigh less than 30kg receive drugs as per Pediatric weight band boxes according to body weight.

1.1. Symptoms and diagnosis methods

Common symptoms of active lung TB are cough with sputum and blood at times, chest pains, weakness, weight

loss, fever and night sweats. Many countries still rely on a long-used method called sputum smear microscopy to diagnose TB. Trained laboratory technicians look at sputum samples under a microscope to see if TB bacteria are present. With three such tests, diagnosis can be made within a day, but this test does not detect numerous cases of less infectious forms of TB.

Diagnosing MDR-TB (see Multidrug-resistant TB section below) and HIV-associated TB can be more complex. A new two-hour test that has proven highly effective in diagnosing TB and the presence of drug resistance is now being rolled-out in many countries.^[9,10]

Table 2: (Categories	Of Cases	And Treatment	Regimens Unde	r Rntcp. ^[13-14]
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Category	Characteristic of TB Cases	Intensive phase	Continuation phase
Category-I	New sputum smear- positive seriously ill, sputum	2 (HP7E)3	A (LID)2
Red box	smear- negative, seriously ill, extra-pulmonary	2 (IIRZE)	4 (IIK)3
Category-II	Palance foilure treatment ofter default	2 (SHRZE)3 followed by	5 (UDE)2
Blue box	Relapse failure treatment after default	1 (HRZE)3	J (HKE)5
Category-III	Sputum smear-negative not seriously ill, extra-	2 (1107)2	4(IID)2
Green box	pulmonary	2 (HRZ)5	4(HK)5

H: Isoniazid (600 mg), R: Rifampicin (450 mg), Z: Pyrazinamide (1500 mg), E: Ethambutol (1200 mg), S: Streptomycin (750 mg)

Out of 12 hospitals, 486 patients presented to these hospitals during the study period. About all these was free availability of drugs in only 2 primary hospitals. Sufficient information education & communication is provided only in the 2 primary hospitals. Being, tertiary care centre , many patients were referred to primary DOTS centres as per RNTCP guidelines as such only 414 patients with TB had gone treatment at primary

health centres during this study period. Most percent of patients preferred government hospitals because of their financial constraints. The treatment outcome is tabulated in the table 3.

S. No.	Evaluation Parameter	No of Patients	Number %
1	Registered Culture case	414	100
2	MDR-TB	57	14
3	Cured	289	70
4	Failured	9	2
5	Default	21	5
6	Died	17	4
7	Continuous treatment on DOTS	21	5

 Table 3: Treatment outcome in 414 patients at DOTS center.

Patients were also interviewed regarding the side effects of the treatment. This report is presented in the figure 1. Fatigue was found to be the most common side effect among TB patients as per the survey report. Vomiting and gastrointestinal upset was also compliant in a significant percentage. About 94% patients were satisfied with the therapy.



Figure 1. Side effect report of TB patients

DISCUSSIONS

With the data procured from the various hospitals, we came to the inference that primary health centres of Dadri Area of west U.P. were following RNTCP guidelines. In addition to this, we observed that there was rise in MDR-TB case & awareness by the trained staff. So, a powerful TB control programme such as RNTCP is required to combat resurgent of TB due to HIV, multi drug therapy. Majority of patients were symptomatically improved within two months of treatment, this improved patient satisfaction. No cost treatment in primary health centres also improved patient's satisfaction.

The findings of the study showed that most % of patients know that TB is a curable disease, although many of them were not aware about RNTCP, by government of India. Majority of patients were also aware about duration of treatment and method of treatment. Knowledge about the side effects and consequences of incomplete treatment was not among all patients. This study emphasises for continuing education programme for TB patients, to teach them more about DOTS and consequences of incomplete treatment.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

ACKNOWLEDGEMENT

The authors would like to thank Vishveshwarya Group of institutions, Greater Noida, Uttar Pradesh and health centres of Dadri region for helping us to complete this study.

REFERENCES

1. World Health Organization. Global tuberculosis report, 2014. Available: http:// apps.who.int/iris/bitstream

/10665/137094/1/9789241564809_eng.pdf?ua=1.

- 2. National Tuberculosis Institute; Tuberculosis in a rural population of south India: A five year epidemiological study; Bull. World Health Organization, 1974; 51: 473.
- 3. Sangoian IL and Apresian EA. Health education and knowledge regarding tuberculosis among healthy person and patients with tuberculosis. Problemy Tuberkkuleza, 1990; (10): 20-22.
- 4. Regional Meeting of National TB Programme Managers, WHO/SEARO, New Delhi, India. Geneva: WHO; Nov 2-5, World Health Organization. TB/HIV in the South-East Asia Region Status Report, 2009; 2–3.
- 5. Emerging Infectious Diseases, 2013 Oct; 19(10). www.cdc.gov/eid. http://www.cdc.hbi.ir.
- 6. India's Progress Toward Achieving the Millennium Development Goals. Nath A Indian J Community Med, 2011 Apr; 36(2): 85-92.
- Human resources development for TB control: report of a consultation held on 27 and 28 August, 2003. Geneva: World Health Organization/Rockefeller Foundation, 2004. (WHO/HTM/TB/2004.340).
- Effectiveness of the direct observation component of DOTS for tuberculosis: a randomised controlled trial in Pakistan. Walley JD, Khan MA, Newell JN, Khan MH Lancet, 2001 Mar 3; 357(9257): 664-9.
- Diagnostic delays in access to tuberculosis care in counties with or without the National Tuberculosis Control Programme in rural China. Xu B, Jiang QW, Xiu Y, Diwan VK Int J Tuberc Lung Dis, 2005 Jul; 9(7): 784-90.
- 10. History of tuberculosis and drug resistance. Marais BJ, Zumla A, N Engl J Med, 2013 Jan 3; 368(1): 88.
- 11. Dhingra VK, Rajpal S, Aggarwal N, Aggarwal JK. Treatment of tuberculous pleural effusion patients

and their satisfaction with DOTS: 1¹/₂ year follow up. Ind J Tuberc, 2004; 51: 209–12.

- 12. Unit 5, part 1: The DOTS strategy for controlling TB. [Last accessed 2018 Dec 22].
- Pandit S, Dey A, Chaudhuri AD, Saha M, Sengupta A, Kundu S, et al. Five-years experiences of the Revised National Tuberculosis Control Programme in northern part of Kolkata, India. Lung India, 2009; 26: 109–13.
- 14. Chhetri AK, Saha A, Verma SC, Palaian S, Mishra P, Shankar PR. Study of adverse drug reactions caused by first line anti-tubercular drugs used in directly observed treatment, short course (DOTS) therapy in Western Nepal, Pokhara. J Pak Med Assoc, 2008; 58: 531–6.