

Adult Infectious Diseases Clinic – Tuberculosis Treatment – Follow-up Form Intensive Phase – (5)

Patient Initials _____ IDC Number _____ TB number _____
 Contacts: Tel No 1 _____ (_____, _____) Tel No 2 _____ (_____, _____)
 Type of TB: PTB smear positive PTB smear negative EPTB, site: _____
 Category of treatment: Category 1 Category 2 Date of TB Treatment Initiation ____/____/____

Expected date of TB follow-up visit	2 weeks (____/____/____)		End of Intensive Phase ¹ (____/____/____)		End of Extended Intensive Phase ¹ (____/____/____)	
Visit date	____/____/____		____/____/____		____/____/____	
Name of medical officer:						
Treatment Response	Absent	Present	Absent	Present	Absent	Present
Symptoms						
Cough	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Excessive night sweats	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Weight Loss	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Anorexia	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Other:	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Drug Side Effects						
Joint Pains ²	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Peripheral neuropathy ³	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Skin rash ^{4,5}	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Deafness ^{4,6}	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Dizziness ^{4,6}	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Jaundice ^{4,7}	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Vomiting ^{4,8}	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Difficulty with vision ^{4,9}	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Other:	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Signs						
Pleural effusion	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____	<input type="checkbox"/>	<input type="checkbox"/> _____
Temperature (°C)	<input type="checkbox"/> afebrile <input type="checkbox"/> _____		<input type="checkbox"/> afebrile <input type="checkbox"/> _____		<input type="checkbox"/> afebrile <input type="checkbox"/> _____	
Weight (kg)						
Drug Adherence (%)¹⁰						
Treatment Regimen¹¹						
Follow-up investigations¹²						
Date of Investigation			____/____/____		____/____/____	
Sputum Smear			<input type="checkbox"/> ZN <input type="checkbox"/> Auramine <input type="checkbox"/> Neg <input type="checkbox"/> 1+ <input type="checkbox"/> 2+ <input type="checkbox"/> 3+		<input type="checkbox"/> ZN <input type="checkbox"/> Auramine <input type="checkbox"/> Neg <input type="checkbox"/> 1+ <input type="checkbox"/> 2+ <input type="checkbox"/> 3+	
X-ray			Improvement? <input type="checkbox"/> Yes <input type="checkbox"/> No Other:			

Labs ¹³	Baseline results	If baseline abnormalities			
	Date:	Date:	Date:	Date:	Date:
RFTs ¹⁴	Creat: BUN:	Creat: BUN:	Creat: BUN:	Creat: BUN:	Creat: BUN:
LFTs ⁷	AST: ALP: ALT: GGT: Other:	AST: ALP: ALT: GGT: Other:	AST: ALP: ALT: GGT: Other:	AST: ALP: ALT: GGT: Other:	AST: ALP: ALT: GGT: Other:

Please turn over for comments and legend

Does this patient need an extended intensive phase¹⁵ ? No Yes
 Can this patient start continuation phase¹⁵ ? Yes, continue on TB Continuation Phase Form
 No, restart treatment according to guidelines
 No, other treatment outcome: _____

Name _____ Signature _____ Date ____/____/____

Adult Infectious Diseases Clinic – Tuberculosis Treatment – Follow-up Form Intensive Phase – (5)

Notes (please add date):

Legend:

1. Intensive phase = 2 months (category 1 regimen), 3 months (category 2 regimen). Extended intensive phase = extension of 1 month if sputum is smear positive at end of regular intensive phase. See note 15.
2. Frequent with PZA, treat with aspirin/NSAID
3. Usually due to INH, increase dose of pyridoxine to 100mg
4. Major side effect according to WHO and NTLF; consult with TB-coordinator
5. Mild rash: treat with antihistamines; petechiae: check for thrombocytopenia due to RIF, stop RIF; generalized erythematous rash with fever and/or mucous membrane involvement (Stevens- Johnson Syndrome): stop all drugs immediately. If severe TB: start 3 new TB drugs, when rash has subsided reintroduce drugs one by one.
6. Ototoxicity due to streptomycin. Stop SM.
7. Moderate=AST 5-10x upper limit of normal, severe= >10x. Stop TB drugs immediately if AST >5x upper limit of normal without symptoms, or >3x with symptoms, or with significant rise in bilirubin or ALP. Check for hepatitis A/B/C. See SOP for guidelines on management and follow-up.
8. No absorption of drugs. Check for cause and treat.
9. Retrobulbar optic neuritis due to EMB, first sign is colour blindness. Immediately stop EMB.
10. Assess adherence to TB drugs using TB treatment card and pill counts. Record in percentages.
11. Record the current regimen (f.e. HERZ, EH), plus possible changes due to toxicity or hepatic/renal insufficiency.
12. The investigation that originally led to the diagnosis of TB is to be used for the assessment of treatment response. Repeat the chest X-ray in PTB smear negative cases with initial radiographic abnormalities at the end of intensive phase and at the end of treatment. Repeat the abdominal ultrasound in abdominal EPTB only at the end of treatment. If the diagnosis was made on the basis of other investigations, assessment of the outcome on clinical grounds may suffice. Discuss with TB coordinator.
13. Order RFTs and LFTs in all patients at TB treatment initiation. Check these baseline results at the first follow-up visit. If abnormal, follow-up during the rest of the treatment is indicated. See SOP for guidelines on management and follow-up.
14. If new renal insufficiency with creatinine clearance <30ml/min: adjust dosage of ethambutol, pyrazinamide and streptomycin. Discuss with TB coordinator.
15. First sputum evaluation at the end of intensive phase (category 1: 7 weeks, category 2: 11 weeks). If negative: start continuation phase. If positive and negative at treatment initiation: send for culture & sensitivity testing, consider case as treatment failure and restart treatment with category 2 regimen (if on category 1) or discuss with TB coordinator (if on category 2). If positive and positive at treatment initiation: counsel for adherence, extend intensive phase with 1 month, and redo sputum examination afterwards. If positive after extended intensive phase: send for culture & sensitivity, consider case as treatment failure and restart treatment with category 2 regimen (if on category 1) or discuss with TB coordinator (if on category 2).
If the patient is transferred out, defaults, dies or stops treatment for any other reason during intensive phase, please tick the appropriate box and fill out this treatment outcome.