

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention Coordinating Center for Infectious Diseases, Mail Stop G-<mark>25</mark> Atlanta, Georgia 30333

OMB Form NO. <u>0920-0600</u> Exp. Date <u>05/31/2013</u>

Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* and Nontuberculous Mycobacteria Drug Susceptibility Testing

<u>WARNING</u>: The panel provided in this survey consists of viable cultures of Mycobacterium tuberculosis complex, some of which are drug resistant. The cultures in the panel should be considered hazardous and capable of transmitting infection. Testing should only be done if the recommended safety procedures are followed as described in the *Centers for Disease Control and Prevention's* Biosafety in Microbiological and Biomedical Laboratories, 2007, 5th Edition.

This manual can be accessed at http://www.cdc.gov/od/ohs/biosfty/bmbl5/BMBL 5th Edition.pdf. Biosafety Level 3 practices should be used when testing MTBC cultures.

Check the contents of your package. It should contain:

- (1) Cover letter
- (2) Results Worksheet for recording testing results with instructions.
- (3) Shipping container with five (5) cultures labeled "TB Cultures." The culture tubes are labeled with individual identification codes.

If the contents of your package are not complete, or if additional cultures are required, please call Lois Diem at 404-639-2862 immediately.

INSTRUCTIONS FOR ENTERING RESULTS

Results must be entered in the on-line data entry system only no later than **June 6, 2011.** You will need your TPEP number and password. If you have forgotten or misplaced your password please contact Suzette Brown at 404 498-2283 or 888-465-6062.

- 1. After testing your samples, enter your results at the CDC Tuberculosis Drug Susceptibility Website: http://wwwn.cdc.gov/mpep/mtbds/login.aspx
- 3. If you can not use the on-line data entry system, please complete the Results Worksheet and contact the project coordinator at (888) 465-6062 or 404-498-2283.
- 4. For multiple choice questions beginning on page 4 of the Results Worksheet, **fully blacken the circle** to the left of the appropriate answer. **Please do not use check marks** (☐) **or cross marks** (X) **within the circles**.

MTBC Results Worksheet

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CDC DRUG SUSCEPTIBILITY TESTING PROGRAM FOR MYCOBACTERIUM TUBERCULOSIS RESULTS FORM

Report your results Online (password required) at:
http://wwwn.cdc.gov/mpep/mtbds/login.aspx
TPEP number: (you will need this to enter your results online)
DEADLINE for submission June 6, 2011

Please note: Treat these cultures in the same manner that you routinely treat MTBC isolates. Please test each MTBC isolate against first line drugs and any second line drug tested in your laboratory. This will provide you with an opportunity to evaluate your performance for testing second-line drugs.

<u>WARNING</u>: The panel provided in this survey consists of viable cultures of Mycobacterium tuberculosis complex, some of which are drug resistant. The cultures in the panel should be considered hazardous and capable of transmitting infection. Testing should only be done if the recommended safety procedures are followed as described in the *Centers for Disease Control and Prevention*'s Biosafety in Microbiological and Biomedical Laboratories, 2007, 5th Edition.

This manual can be accessed at http://www.cdc.gov/od/ohs/biosfty/bmbl5/BMBL 5th Edition.pdf.

Biosafety Level 3 practices should be used when testing MTBC cultures.

If you do not have the capacity to enter your results online or if you need assistance contact Suzette Brown at:

- telephone (888) 465-6062 or (404) 498-2283
- email <u>MTBNTMDST@CDC.GOV</u>

Perso	n(s) Com	pleting Form:	
1.	Name:		
2.	Title:		
		MTBC Worksheet	

3.	Please indicate the primary classification of your laboratory. (Please blacken only one circle.)
ı	Hospital [e.g., city, county, district, community, state, regional, military, Veterans Administration, Federal government (other than military), privately-owned, university, HMO/PPO-owned and operated, religious-associated]
1	Health Department [e.g., city, county, state, regional, district, national reference laboratory]
1	Independent (non-hospital-based) [e.g., commercial, commercial manufacturer of reagents, HMO satellite clinic, reference laboratory (non-government affiliated)]
ı	Other [e.g., university-associated research, Federal government research (nonmilitary), privately-funded research]
4.	In the last calendar year (January 1 - December 31), how many <i>Mycobacterium tuberculosis</i> isolates (excluding quality control isolates) did your laboratory test for drug susceptibilities? (Please write the number of <i>Mycobacterium tuberculosis</i> isolates your laboratory tested for susceptibility in the boxes below.) <i>Mycobacterium tuberculosis</i> isolates:
cas the app	e following questions pertain to the receiving and testing of the culture panel. In most les, blacken the circle corresponding to your response in the circle provided to the left of answer. Some questions may require more than one response; please blacken all that oly. In some cases, you will be asked to fill in the boxes to the right of the answer with appropriate comment or number.
5.	On what date was the culture panel received in your laboratory? Month Day Year

MTBC Worksheet

6.	What was the condition of the cultures in the panel when they arrived? (Please blacken only one circle.)
	○ Satisfactory
	○ Broken
	Other (please explain):
7.	What method(s) was used in your laboratory to perform drug susceptibility testing on the MTBC isolates in this shipment ? (Please blacken all that apply.)
	Agar Proportion (Middlebrook 7H10)
	Agar Proportion (Middlebrook 7H11)
	Genotype MTBDRplus (<i>Hain</i> Lifescience)
	Genotype MTBDRsl (<i>Hain</i> Lifescience)
	 Lowenstein Jensen (LJ) proportion method
	○ MGIT System
	Radiometric (BACTEC 460)
	○ VersaTREK Myco
	XPERT MTB/RIF (Cepheid)
	Laboratory Developed Test (LDT) (please specify):
	Other (please specify):
8.	If your laboratory uses more than one method for testing routine samples for first-line drugs for MTBC susceptibility, please indicate the initial method that is used. (Please blacken only one circle.)
	Agar Proportion (Middlebrook 7H10)
	Agar Proportion (Middlebrook 7H11)
	Genotype MTBDRplus (Hain Lifescience)
	Genotype MTBDRsl (<i>Hain</i> Lifescience)
	Lowenstein Jensen (LJ) proportion method
	○ MGIT System
	Radiometric (BACTEC 460)
	○ VersaTREK Myco
	XPERT MTB/RIF (Cepheid)
	Laboratory Developed Test (LDT) (please specify):
	Other (please specify):

MTBC Woksheet

9.	If you use Middlebrook 7H10 or 7H11 media as either an initial or secondary method of MTBC drug susceptibility testing, your media is: (Please blacken all that apply.)
	ourchased "commercially-prepared" containing anti-tuberculosis drugs
	 prepared in-house with disks containing anti-tuberculosis drugs
	 prepared in-house by reconstituting and adding anti-tuberculosis drugs
	 Not Applicable – We do not use Middlebrook media
10a.	In your opinion, is there a need for offering performance evaluation of NTM strains? Yes No
10b.	If yes – For your laboratory, would it be more advantageous to offer evaluation of:
	Rapidly growing NTMSlowly growing NTM

Continue to the next page.

MTBC Worksheet

11. For each antimicrobial concentration tested: Select the antimicrobial, test method, the concentration of the antimicrobial and a result (R=Resistant, S=Susceptible, O=Other). If the isolates in the panel were tested using more than one concentration of an antimicrobial, record those results on lines that correspond to the antimicrobial you are testing (**Example 1**). If you need more lines than are provided for that antimicrobial, please record results in the blank lines provided at the bottom of the result page. Do not cross out an existing antimicrobial and write another drug name over it (example 2).

If you are testing an antimicrobial not listed on the result page, record the entire drug name (no abbreviations), a concentration and a result in the blank lines provided at the bottom of the result page. Please make sure that each result is recorded on a provided line and not written in the margins outside the form. Make a copy of the result page if you do not have enough room on the provided page to record all results.

Other responses related to susceptibility results such as Borderline, Contaminated, No Growth, etc. can be abbreviated and recorded to the right of the "O" selection in the result columns (examples 1 and 3).

1. Following are examples of **CORRECTLY** reported *M. tb* results.

Isoniazid	$A \bullet C \odot$		0		1	$\mathbb{R} lue{\mathbb{O}}$	● S O	$\mathbb{R} \bullet \mathbb{O}$
Isoniazid	● B C O		0		2	$\mathbb{R} \bullet \mathbb{O}$	● \$0	$\mathbb{R} \bullet \mathbb{O}$
Isoniazid			1	•	0	$\mathbb{R} \bullet \mathbb{O}$	● \$ ®	®S ● NG

2. Following are examples of **INCORRECTLY** reported *M. tb* results.

Isoniazid	ABCO	1	2	-	•	-	0		RØ	0	3 (S)(O)
Isoniazid	● B C ●							$\mathbb{R} lacktriangle$		•	

MTBC Worksheet **Please provide the Test Method, the Concentration, and the Test Results for each line reported.

11. Use the blank lines provided at the end of the form for other drugs or additional concentrations. Please provide the complete drug name when filling in additional spaces.

A=Agar Proportion (7H10) B= Agar Proportion (7H10),

C= BACTEC
D= VERSA
E= MGIT

F= L-J Proportion M= Molecular Method

O=Other

Culture Identification Codes (Fill in ONE letter for each culture)

R=Resistant S=Susceptible B=Borderline C=Contaminated N=No Growth

Antimicrobial	Test Method	Con µg/ı		A	В	С	D	Е	
Isoniazid	ABCDEFMO			RSBCNO	RSBCN0	RSBCNO	BSBCNO	RSBCNO	
Isoniazid	ABCDEFMO			RSBCNO	RSBCN0	RSBCNO	8SBCN0	RSBCNO	
Isoniazid	ABCDEFMO		-	RSBCNO	RSBCN0	RSBCN0	RSBCN0	RSBCNO	
Isoniazid	ABCDEFM0			RSBCNO	RSBCNO	RSBCN0	RSBCN0	RSBCNO	
Rifampin	ABCDEFMO		11.1	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Rifampin	ABCDEFMO		11.1	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Rifampin	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Pyrazinamide	ABCDEFMO			RSBCNO	RSBCN0	RSBCN0	RSBCN0	RSBCN0	
Pyrazinamide	ABCDEFMO		1 .	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Pyrazinamide	ABCDEFMO		 	RSBCNO	RSBCNO	RSBCNO	RSBCN0	RSBCNO	
Ethambutol	ABCDEFMO			RSBCNO	RSBCN0	RSBCN0	RSBCN0	RSBCNO	
Ethambutol	ABCDEFMO		1 .	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Ethambutol	ABCDEFMO		-	RSBCNO	RSBCN0	RSBCN0	RSBCN0	RSBCN0	
Streptomycin	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	RSBCN0	RSBCNO	
Streptomycin	ABCDEFMO		 	RSBCN0	RSBCN0	RSBCN0	RSBCNO	RSBCNO	
Streptomycin	ABCDEFMO			RSBCN0	RSBCN0	RSBCN0	RSBCN0	RSBCNO	
Ethionamide	ABCDEFMO		-	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Ethionamide	ABCDEFMO			RSBCN0	RSBCN0	RSBCNO	8SBCN0	@\$BCNO	
Kanamycin	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	RSBCN0	RSBCNO	
Kanamycin	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	RSBCN0	RSBCNO	
Capreomycin	ABCDEFM0		1 .	RSBCNO	RSBCNO	RSBCNO	8SBCN0	RSBCNO	
Capreomycin	ABCDEFM0		11.1	RSBCNO	RSBCNO	RSBCNO	8SBCN0	RSBCNO	
Cycloserine	ABCDEFM0		1 -	RSBCNO	RSBCNO	RSBCNO	8SBCN0	RSBCNO	
Cycloserine	ABCDEFMO		11.1	RSBCNO	RSBCN0	RSBCNO	RSBCN0	RSBCNO	
p-Aminosalicylic acid	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
o-Aminosalicylic acid	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
Amikacin	ABCDEFMO		1 -	RSBCNO	RSBCN0	RSBCNO	8SBCN0	RSBCN0	
Amikacin	ABCDEFMO		1 -	RSBCNO	RSBCN0	RSBCNO	8SBCN0	RSBCN0	
Ofloxacin	ABCDEFMO		1 -	RSBCNO	RSBCNO	RSBCNO	8SBCN0	RSBCNO	
Ofloxacin	ABCDEFMO		1 -	RSBCNO	RSBCN0	RSBCNO	8SBCN0	RSBCN0	
Ciprofloxacin	ABCDEFMO			RSBCNO	RSBCNO	RSBCNO	8SBCN0	RSBCNO	
Ciprofloxacin	ABCDEFMO		1 -	RSBCNO	RSBCNO	RSBCNO	8SBCN0	RSBCN0	
1	ABCDEFMO		11.	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	
	ABCDEFM0			RSBCNO	RSBCNO	RSBCN0	RSBCN0	RSBCNO	
	ABCDEFMO		1 _	RSBCNO	RSBCNO	RSBCNO	RSBCNO	RSBCNO	

Note: Please provide the complete drug name when filling in additional spaces.

Public reporting burden of this collection of information is estimated to average 6 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB Control Number. Send comments regarding this burden estimate or any other aspect of this collection of information,

including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia, Attn: PRA 0920-0600