

**AFTER-ACCREDITATION PROMOTION
INTERIM GUIDELINES FOR THE USE OF AND SAFETY MONITORING OF BEDAQUILINE
(SIRTURO™) FOR THE TREATMENT OF PULMONARY MULTIDRUG-RESISTANT
TUBERCULOSIS**

WB2329

FACULTY/CREDENTIALS:

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DATE: **October 25, 2013**

ORIGINATION/EXPIRATION DATES: **October 25, 2013 -
October 25, 2015**

LOCATION: **MMWR Journal**
<http://www.cdc.gov/mmwr>

HARDWARE/SOFTWARE: Computer-Internet;

MATERIALS: Access to computer

TARGET AUDIENCE: Physicians, Registered Nurses, Laboratorians, Medical Assistants, Physician Assistants, Public Health Professionals, Health care workers, health educators

PREREQUISITES: Knowledge of Pulmonary Multidrug-Resistant Tuberculosis

PROGRAM DESCRIPTION: The goal of this report is to provide guidance for the use of bedaquiline fumarate, a newly approved MDR TB drug for the treatment of Pulmonary Multidrug-Resistant Tuberculosis.

OBJECTIVES:

1. describe the basic mechanism of action of bedaquiline
2. describe the basic epidemiology of MDR TB
3. describe the way in which drug-resistant TB develops

4. describe the rationale and methods employed by CDC to develop new recommendations
5. describe the efficacy and safety data that led to the accelerated approval of bedaquiline by the FDA
6. describe the patients in whom bedaquiline may be considered
7. describe the dosing, administration, precautions and safety monitoring recommended for patients receiving bedaquiline

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ACCREDITATION STATEMENTS:

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Presentations will not include any discussion of the unlabeled use of a product or a product under investigational use with the exception of Kenneth Castro, Terence Chorba, Phillip LoBue and Sundari Mase's discussion of Bedaquiline. They will be discussing the approval of Bedaquiline, but its use for extrapulmonary and pediatric TB is off label.

CDC does not accept commercial support.

SUPPORT/FUNDING: CDC

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METHOD OF PARTICIPATION:

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