

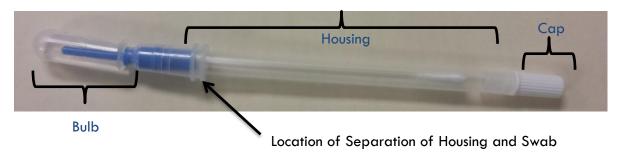
# T-COR 8<sup>TM</sup> BW C2T<sup>TM</sup> Multiplex 2 Burkholderia, SEB, Bot, IC Instructions

TC-8021-064

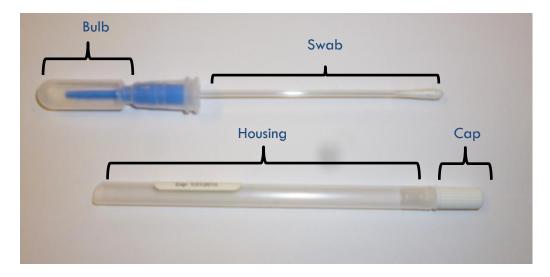
# **Instrument Setup**

- 1. Review all information in the Quick Start Guide, Hardware Manual and Software Manual prior to operating the T-COR 8<sup>TM</sup>.
- 2. Plug in the Magnetic Mixer into the T-COR 8 via the USB ports before powering on the T-COR 8.
- 3. Power on the T-COR 8 as directed in the Quick Start Guide.
  - Available usernames are "user" and "admin" (no password required). Refer to the Software Manual for adding new users.
  - It is recommended that a password be added to the "admin" user account. See the Software Manual for details on changing a password.

## **Sample Processing Device**

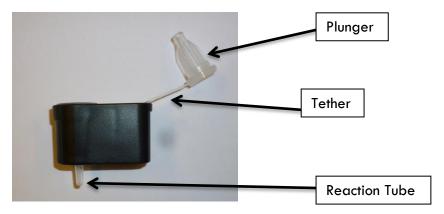


#### **Dissembled Sample Processing Device**





## **C2T Cartridge**



### **Sample & Reaction Preparation**

- 1. If using the Sample Processing Device:
  - a. Carefully remove the swab from the Sample Processing Device by pulling housing away from the bulb. Ensure that the swab does not touch any surfaces.
  - b. Swab the desired area or sample.
    - i. Use a sweeping motion to ensure the area was thoroughly swabbed.
    - ii. Use one swab per area or sample.
  - c. Return the swab into the housing. (Do not remove the cap.)
  - d. Snap the bulb at the top of the device. (Do not remove the cap.)
  - e. Squeeze the bulb to move the fluid into the housing. (Do not remove the cap.)
  - f. Open each assay pouch containing the C2T cartridge and place each cartridge on the C2T rack.
  - g. Un-screw the cap on the bottom of the Sample Processing Device.
  - h. Squeezing the housing and/or bulb, dispense 5 drops from the Sample Processing Device into each C2T cartridge. Dispensing more than 5 drops will not affect the performance of the C2T cartridge.
  - i. Re-screw the cap on the bottom of the Sample Processing Device and retain until the end of the test.
- 2. If NOT using the Sample Processing Device:
  - a. Open each assay pouch containing the C2T cartridge and place each cartridge on the C2T rack.
  - b. Use a micropipette to add 100µL of sample into the C2T cartridge.
  - c. Carefully pick up the open cartridge from the C2T rack. Hold cartridge upright and carefully but firmly tap the reaction tube of the open cartridge on a hard surface a few times. This will remove bubbles from the cartridge and will improve load. Place open cartridge back on the C2T rack.
- 3. Closing the C2T cartridge:

**NOTE:** This action moves the sample onto the reaction tube of the cartridge.

- a. The cartridge <u>must</u> be on the C2T rack to ensure proper closure of the cartridge.
- b. Align the plunger with the corresponding cartridge cavity opening.
- c. Firmly press down on the plunger until the plunger is flush with the cartridge.
- d. Confirm that the cartridge is fully closed by visually inspecting the plunger/cartridge at eye level. If the plunger is elevated from the cartridge, push on the plunger again to firmly close the cartridge.



#### **Experimental & Run Setup**

- 1. For complete instructions, refer to the Software Manual, Guided Workflow Section.
- 2. After signing in to the T-COR 8, go to the Menu button and select "New Run."
- 3. Select the "Barcode" button, located under Guided Setup. The barcode scanner will illuminate on the right hand side of the T-COR 8.



**WARNING:** Laser Radiation: Class 2 Laser Product. Do not stare into barcode beam. Never direct a laser at other people, highly reflective surfaces, or any other object not intended for designated use.

- 4. Hold the barcode on the cartridge approximately 3 inches away from scanner, centering the crosshair to the middle of the barcode.
- 5. Follow the instructions on the screen to properly mix and load the sample.
  - a. Remove the cartridge from the rack and place it in the mixer and press [Mix] to start mixing.
  - b. When Mixing is completed, press [Next].
  - c. Open the well indicated. Press [Yes] to confirm visual of the blue light in the reaction well.
  - d. Remove the cartridge from the mixer.
  - e. Visually confirm that the reaction tube meets the following requirements:
    - 1. Contains mix that is blue in color and NOT clear.
      - The blue color indicates that the magnetic mixer worked properly and the mix in the reaction tube mixed successfully.
    - 2. Contains volume that fills more than 50% of the reaction tube.
      - Cartridges that fail to meet this requirement should be discarded.
    - 3. No air bubbles are present at the bottom or side of the tube.
  - f. If the cartridge meets the above requirements, proceed to place cartridge into the well indicated, and press [Next].
  - g. If desired, name the sample by pressing in the gray box marked Sample or by pressing on the Barcode button. Press [Next] to continue with the setup.
  - h. Press [Add well] to load the next cartridge.
  - i. The barcode scanner will illuminate on the right hand side of the instrument. Hold the barcode located on the cartridge approximately 3 inches away from the scanner, centering the crosshair to the middle of the barcode.
  - j. Repeat steps "a" through "i" for the remaining samples.
  - k. After all the cartridges have been loaded, press [Start Run].
  - 1. If desired, enter a Run Name. Press [Start Run] to begin the run.
- 6. When a run is complete, analyze samples (follow Steps in Run Analysis Section).
- 7. After analysis is complete, dispose of the used T-COR 8 cartridges properly.

#### Run Analysis

1. Use the table button, to determine the results of the run. The instrument will identify the run as either:



- a. Not Detected
- b. Invalid (the sample should be re-tested)
- c. Detected (each target detected will be listed)
  - i. To confirm results of a detected sample, re-test the sample.
- 2. Due to space constraints, some assays are abbreviated in the table/protocol. See table below for assays and abbreviations.

Assay Name	Abbreviations Used
Burkholderia	Burk
S. aureus B	SEB
C. botulinum	Bot

3. Additional analysis can be performed, refer to the T-COR 8 Software Manual for guidance.

### **IMPORTANT:**

- 1. When analyzing the run, please note the thermal cycling protocol has 10 cycles hidden from view. Therefore, a sample with a Ct of 25 is actually a Ct of 35.
- 2. SEB can be found in the environment (such as skin, surfaces and other reagents). Care should be used to interpret low level positive samples as these may be environmental.

#### **Technical Assistance**

If technical assistance is needed, please contact Tetracore at 240-268-5400.