Title: MIC82000-BioFire Torch Type: Laboratory Services Program SOP

Issuing Authority: Director of Health Services

Next Review Date:

Policy Number: Date Approved:

PROGRAM Standard Operating Procedure – Laboratory Services				
Title: MIC82000 -	Policy Number:			
BioFire Torch				
Program Name: Laboratory Services				
Applicable Domain: Lab, DI and Pharmacy Services				
Additional Domain(s):				
Effective Date:	Next Review Date:			
Issuing Authority:	Date Approved:			
Director of Health Services				
Accreditation Canada Applicable Standard: N/A				

GUIDING PRINCIPLE:

The BioFire Torch is an automated in vitro diagnostic (IVD) device intended to detect multiple nucleic acid targets contained in clinical specimens using nested multiplex PCR (nmPCR) in a closed system.

PURPOSE/RATIONALE:

This standard operating procedure describes the BioFire Torch and its components.

SCOPE/APPLICABILITY:

This procedure applies to Medical Laboratory Technologists processing specimens using the BioFire Torch.

SAMPLE INFORMATION:

Refer to assay specific BioFire procedures for sample information

REAGENTS and/or MEDIA:

Refer to assay specific BioFire procedures for reagent information

SUPPLIES:

Refer to assay specific BioFire procedures for supply information

EOUIPMENT

BioFire Torch

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Policy Number: Date Approved: Page 1 of 5 Title: MIC82000-BioFire Torch

Type: Laboratory Services Program SOP

Issuing Authority: Director of Health Services

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ENVIRONMENTAL CONTROLS:

Operating temperature: 15°C to 30°C

Relative humidity: 20 to 80% relative humidity, non-condensing

SPECIAL SAFETY PRECAUTIONS:

Containment Level 2 facilities, equipment, and operational practices for work involving infectious or potentially infectious materials or cultures:

- Ensure that appropriate hand hygiene practices be used
- Lab gown must be worn when performing activities with potential pathogens.
- Gloves must be worn when direct skin contact with infected materials is unavoidable
- Eye protection must be used when there is a known or potential risk of exposure of splashes
- All procedures that may produce aerosols or involve high concentrations or large volumes should be conducted in a biological safety cabinet (BSC)
- The use of needles, syringes and other sharp objects should be strictly limited

All patient specimens are assumed to be potentially infectious. Routine Practices must be followed. Since viable micro-organisms are used, all cultures must be handled with appropriate precautions. All equipment in contact with cultures should be decontaminated by appropriate methods.

QUALITY CONTROL:

• Refer to specific BioFire procedures for quality control information

PROCEDURE INSTRUCTIONS:

Step	Action					
System Components						
1	 BioFire Torch Instrument: The BioFire Torch is composed of one to twelve BioFire Torch Modules connected to a BioFire Torch System Base running BioFire Torch Software The major components of the BioFire Torch includes: BioFire Torch System Base – Includes barcode scanner, touch screen and USB ports. The BioFire Torch System Base houses up to two BioFire Torch Modules. BioFire Torch Module – Interacts with the reagent pouch to purify and amplify targeted nucleic acid sequences using nmPCR, includes pouch slot and LED status light. Each BioFire Torch Module can be randomly and independently accessed to run a reagent pouch. BioFire Torch Duplex – Module enclosure; houses up to two BioFire Torch Modules. Up to five BioFire Torch Duplex Module enclosures, each capable of housing up to two additional BioFire Torch Modules, may be added on top of the BioFire Torch System Base. 					

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Policy Number: Page 2 of 5

Title: MIC82000-BioFire Torch

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Next Review Date:

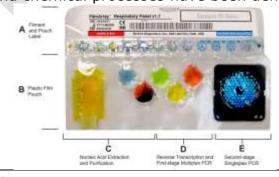
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Policy Number: Date Approved:



BioFire Pouch:

- Each BioFire Pouch is a self-contained, closed-system disposable packet that houses all the chemistry required to isolate, amplify and detect nucleic acid from a sample
- The reservoirs in the rigid plastic component or fitment of the pouch (A) contain freeze-dried reagents
- The flexible plastic film portion of the pouch (B) is divided into discrete segments (blisters) which, via interactions with actuators and sensors in the BioFire Torch Module, are where the chemical processes are performed:
 - (C) Extraction and purification of nucleic acids from a clinical sample using mechanical lysis (bead beating) and magnetic bead technology
 - (D) First-stage multiplex PCR (including reverse transcription of target RNAs when appropriate)
 - > (E) Second stage singleplex PCR and melting analysis within a multiwell array
- Each pouch contains at least one internal process control. Control material is lysed and the nucleic acids of the control material are extracted along with that of the organisms contained in the sample. When the internal control is positive, proper operation of the BioFire Torch Module and chemical processes have been demonstrated



Mechanical Lysis:

 The first step in processing a sample is to break the outer membrane of the target cells contained in the sample using a device called a beadbeater

• A sensor detects the speed and operation of the bead-beater motor and aborts the run if the bead-beater is not working properly

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Policy Number:

Date Approved:

Page 3 of 5

3

2

Title: MIC82000-BioFire Torch

Type: Laboratory Services Program SOP

Issuing Authority: Director of Health Services

Next Review Date:

Policy Number:

Date Approved:

4	 Nucleic Acid Extraction: Following bead-beating, the nucleic acids contained in the sample are purified by magnetic bead technology 				
	A retractable magnet is used to capture or release the magnetic beads during washes				
	Thermal Control:				
	 The purified nucleic acids are mixed with PCR reagents, which amplify all of the targets identified by the pouch as well as the control material A Peltier device drives the thermocycling of the reverse transcription 				
5	and/or PCR reactions				
	A second Peltier device controls thermocycling for second-stage PCR and DNA melting				
	The thermocycling conditions are controlled by the run protocol associated with each specific reagent pouch and sample type				
	Optics and Imaging:				
	 To identify targets from positive PCR reactions, DNA melting curve 				
	analysis is performed				
	The fluorescence emitted by the dye is imaged by a camera				
	DNA melting curves are captured by slowly increasing the temperature				
6	of the PCR array and capturing the fluorescent signal				
	These images are processed automatically by the System Base and the				
	data is analyzed to determine if the control reactions passed and which targets were detected in the sample				
	 Proper operation and calibration of BioFire Torch Module optics is 				
	monitored by the BioFire Torch Module self-tests and internal controls				
	BioFire Torch Software:				
	The BioFire Torch Software controls the function of each BioFire Torch				
	Module				
	 The software also collects, stores and analyzes data generated by the BioFire Torch Module 				
	 Results of analyses are presented in a test report The BioFire Torch Software automatically starts when the BioFire Torch 				
	is powered on – no login is required				
	The BioFire Torch Toolbar always displays at the top of the screen and				
	consists of three options: Dashboard – Displays the status of each BioFire Torch Module within				
7	the BioFire Torch and guides the operator through the process of				
	operating the modules. The dashboard also displays the status of all				
	runs that are in progress, along with the sample ID, pouch type and				
	time remaining until completion				
	➤ Browse Runs – Allows operators to search for runs and perform				
	operations on individual runs or on group of runs. The runs are presented as a table that lists the date of the run, sample ID and				
	other information about the run				
	 Settings – Allows users to perform administrative type tasks, such as 				
	print options, managing operators, pouch modules, instrument				
	modules, archive, view system logs and switch to admin mode				

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Policy Number: Date Approved: Page 4 of 5

Title: MIC82000-BioFire Torch

Issuing Authority: Director of Health Services

Next Review Date:

Type: Laboratory Services Program SOP
Policy Number:
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BioFire Torch Module Status:

- Solid white light → Module is initializing
- Solid yellow light → Warning, i.e., not connected
- Solid blue light → Module idle and available to run a pouch
- Blinking blue light → Module waiting for pouch insertion
- Fast blinking green light → Pouch inserted but run not started yet
- Solid green light → Run in progress
- Slow blinking green light → Run complete, remove pouch
- Solid red light → Error. Operator intervention required

INTERPRETATION OF RESULTS:

Refer to assay specific BioFire procedures for interpretation of results

REPORTING INSTRUCTIONS:

Refer to assay specific BioFire procedures for reporting of results

REFERENCES:

8

1. BioFire Torch Operator's Manual, HTFA-PRT-0001-05, March 2020

APPROVAL:	
Date	

REVISION HISTORY:

REVISION	DATE	Description of Change	REQUESTED BY
1.0	10 Aug 20	Initial Release	L. Steven
2.0	17 Oct 22	Procedure reviewed	L. Steven

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Policy Number: Date Approved: Page 5 of 5