# **EC Declaration of Conformity**

Manufacturer/
Supplier Information:

BioFire Diagnostics, LLC

515 Colorow Drive

Salt Lake City, Utah 84108, USA

Phone: 1-801-736-6354 Regulatory@BioFireDX.com http://www.BioFireDX.com

We, BioFire Diagnositics, LLC, declare under our sole responsibility, that the product

FilmArray® Blood Culture Identification (BCID) Panel (RFIT-ASY-0126, RFIT-ASY-0127)

meets the provisions of the European Directive 98/79/EC for *In vitro* Diagnostic Medical Devices. The device is classified as a General In Vitro Diagnostic (IVD) Device.

BioFire Diagnostics' quality system is registered to ISO 13485:2016/EN ISO 13485:2016.

The following relevant standards have been met:

## ISO 13485:2016/EN ISO 13485:2016

Medical devices - Quality Management System - Requirements for regulatory purposes

#### EN ISO 14971:2012

Medical devices - Application of risk management to medical devices

### EN 13641:2002

Elimination or reduction of risk of infection related to in vitro diagnostic reagents

#### FN 62366:2008

Medical devices-Application of usability engineering to medical devices

#### EN 13612:2002

Performance evaluation of in vitro diagnostic medical devices

## EN ISO 23640:2015

In vitro diagnostic medical devices - Evaluation of stability of in vitro diagnostic reagents

## EN ISO 15223-1:2016

Medical Devices – Symbols to be used with medical device labels, labeling and information to be supplied – Part 1: General requirements

## EN ISO 18113-1:2011

In vitro diagnostic medical devices – Information supplied by the manufacturer (labeling) – Part 1: Terms, definition and general requirements

## EN ISO 18113-2:2011

In vitro diagnostic medical devices – Information supplied by the manufacturer (labeling) – Part 2: In vitro diagnostic reagents for professional use

Technical documentation demonstrating compliance as described in Annex III of the European Directive 98/79/EC is kept by the manufacturer and can be made available by the authorized representative in Europe - QARAD byba, Cipalstraat 3, B-2440 Geel, Belgium.

The notified body for this product is BSI (Notified body #0086; Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes, MK5 8PP, United Kingdom). Keynes, MK5 8PP, United Kingdom).

Salt Lake City, UT, USA

(Place of issue) 7-31-20

(Date of issue)

Randy Rasmussen
Chief Executive Officer

