# **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® Meningitis/Encephalitis (ME) Panel (RFIT-ASY-0118, RFIT-ASY-0119)

meets the provisions of the European Directive 98/79/EC for *in vitro* Diagnostic Medical Devices. The device is classified as an *In Vitro* Diagnostic (IVD) Device under Annex II list B. BioFire Diagnostics' quality system is registered to ISO 13485:2016 and 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

#### EN 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 IV 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).

Salt Lake City, UT, USA 11-20-18

(Place and date of issue)

Randy Rasmussen
Chief Executive Officer



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