

EU Quality Management System Certificate

Regulation (EU) 2017/746, Annex IX Chapter I and III

IVDR 735494 R003

Manufacturer: BioFire Diagnostics, LLC

Address:

515 Colorow Drive
Salt Lake City
Utah
84108
USA

Single Registration Number: US-MF-000003311

EU Authorised Representative: bioMérieux SA

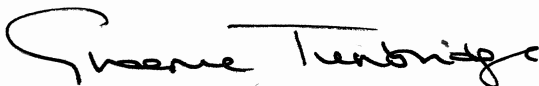
Address:

376 Chemin de l'Orme
69280 Marcy-l'Étoile
France

Scope: See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/746, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class D devices, and self-test, near-patient test and companion diagnostic devices an Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):



Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

First Issue Date: **2021-05-07**

Current Issue Date: **2026-08-03**

Starting Validity Date: **2026-08-03**

Expiry Date: **2031-05-06**

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.
This certificate was issued electronically and is bound by the conditions of the contract.

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Device Schedule: Class D, C and B devices

Class D Devices	Intended purpose
BIOFIRE® Respiratory Panel 2.1 <i>plus</i> (RP2.1 <i>plus</i>)	See IVDR 799673
BIOFIRE® FILMARRAY® Pneumonia Panel <i>plus</i> (BIOFIRE Pneumonia Panel <i>plus</i>)	See IVDR 799675
Class C Devices	Intended purpose
W010507 - Multiple Parameters - Infect. Immunology/NAT	In Vitro Diagnostic PCR devices intended for the detection and identification of infectious agents
IVP3011 - In vitro diagnostic devices which require knowledge regarding molecular biological testing including nucleic acid assays and next generation sequencing (NGS)	
Class B near-patient and professional use test devices	Intended purpose
BIOFIRE® SPOTFIRE® Respiratory/Sore Throat <i>plus</i> (R/ST <i>plus</i>) Panel	See IVDR 832356
BIOFIRE® SPOTFIRE® Respiratory/Sore Throat <i>plus</i> (R/ST <i>plus</i>) Panel Mini	See IVDR 832361
Class B Devices	Intended purpose
IVR 0503 – Devices intended to be used to detect presence of, or exposure to infectious agents	Nucleic acid devices intended to be used for the qualitative detection and identification of an infectious agent

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NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80
Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at Seventh and Eighth Floors, The Acre, 90 Long Acre, London, WC2E 9RA, UK.
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Certificate History

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)

Date	Reference number	Action
2021-05-07	3281374	First Issue
2023-01-24	3444335	Supplemented – Addition of device subcategory group IVR0503
2025-01-13	30336321	Amended – Change of EU Authorised Representative name and address to QbD RepS BV, Groenenborgerlaan 16, 2610 Wilrijk, Belgium
2026-03-03	30657526	Supplemented – Addition of two Class B near-patient and professional use test devices, BIOFIRE® SPOTFIRE® Respiratory/Sore Throat plus (R/STplus) Panel and BIOFIRE® SPOTFIRE® Respiratory/Sore Throat plus (R/STplus) Panel Mini Amended – Added updated description to EMDN scope for W010507
2026-04-20	30564062	Re-Issued – Certificate Renewal
2026-07-23	30753697	Amended – Change of EU Authorised Representative from QbD RepS BV to bioMérieux SA
Current	30727841	Supplement – Addition of devices BIOFIRE® Respiratory Panel 2.1 <i>plus</i> (RP2.1 <i>plus</i>) and BIOFIRE® FILMARRAY® Pneumonia Panel <i>plus</i> (BIOFIRE Pneumonia Panel <i>plus</i>)

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