

EU Quality Management System Certificate

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

IVDR 735494 R000

Manufacturer: BioFire Diagnostics, LLC

Address:

515 Colorow Drive
Salt Lake City
Utah
84108
USA

Single Registration Number: US-MF-000003311

EU Authorised Representative: Qarad EC-REP BV

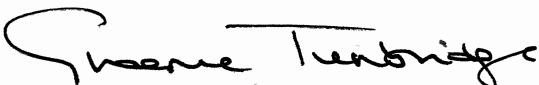
Address:

Pas 257
2440 Geel
Belgium

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 Medical Devices

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

Current Issue Date: **2023-01-24**

Starting Validity Date: **2023-01-24**

Expiry Date: **2026-05-06**

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Device Schedule: Class D, C and B devices

Class C Devices

W010507 - Multiple Parameters - Infect. Immunology

IVP3011 - In vitro diagnostic devices which require knowledge regarding molecular biological testing including nucleic acid assays and next generation sequencing (NGS)

Intended purpose

In Vitro Diagnostic PCR devices intended for the detection and identification of infectious agents

Class B Devices

IVR 0503 – devices intended to be used to detect presence of, or exposure to infectious agents

Intended purpose

Nucleic acid devices intended to be used for the qualitative detection and identification of an infectious agent

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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
Current	3444335	Supplemented – Addition of device subcategory group IVR0503



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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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