

EU Quality Management System Certificate

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

IVDR 780538 R000

Manufacturer: Microbix Biosystems Inc.

Address:

265 Watline Avenue
Mississauga
Ontario
L4Z 1P3
Canada

Single Registration Number: CA-MF-000028149

EU Authorised Representative: OBELIS S.A.

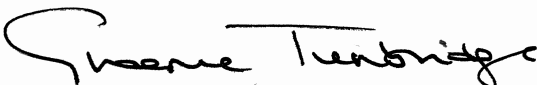
Address:

Bd Général Wahis, 53
B-1030 Brussels
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 Global Regulatory & Quality

First Issue Date: **2024-09-26**

Current Issue Date: **2024-09-26**

Starting Validity Date: **2024-09-26**

Expiry Date: **2029-09-25**

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Device Schedule: Class D, C and B devices

Class B devices	Intended purpose
IVR 0701 – Devices which are controls without a quantitative assigned value.	In vitro diagnostic unassayed controls intended to be used for monitoring performance of assays for the detection of infectious agents.



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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
Current	3789066	Issued



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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.