BIO-RAD	GLOBAL FORM	04.01.GLB.FRM.00289

EU DECLARATION OF CONFORMITY

Division/Group: RAQA	Revision: 1

ID-Diluent 2

Id-n°: 05761

REF 009260, 009280, 009290

BUDI-DI : 361052A002678D

DiaMed GmbH
Pra Rond 23, 1785 Cressier, Switzerland
SRN : CH-MF-000020826

EC REP

FRANCE, Bio-Rad, 3 boulevard Raymond Poincaré
92430 Marnes-la-Coquette
SRN : FR-AR-000006264.

We, as the manufacturer of the device(s) take sole responsibility for and hereby declare that the above mentioned product(s) meet(s) the provisions of the following Regulation(s) / Directive(s):

■ Regulation EU 2017/746 on *in vitro* Diagnostic medical devices

Risk CLASS:

☒ A ☐ B ☐ C ☐ D

CONFORMITY ROUTE

☒ ANNEX I & II+III

Digitally signed by Levința Alexandru
Date: 2025.07.21 12:13:43 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

Common Specification (CS): Not Applicable

EU DECLARATION OF CONFORMITY

Division/Group: RAQA	Revision: 1

Date of the first issuance of the EU Declaration of Conformity: 04-05-2022

Place, Date:	Cressier, 04.05.2022
Signed by:	Jérémy Poropane
Function:	Associate Director, Regulatory Affairs, Bio-Rad CDG
Signature:	DocuSigned by: Jeremy Poropane Signer Name: Jeremy Poropane Signing Reason: I am the author of this document Signing Time: 04-May-2022 | 10:22:23 AM PDT F9BD0CB5E5B347DE9FEC587797500EC0