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CE DECLARATION OF CONFORMITY

according to EU_2017/746 Regulation on „In Vitro Diagnostic Medical Devices”

Manufacturer: **BIOLAB Inc.**
Address: **Öv u. 43., H-1141 Budapest**
Phone: **+36 1 221 9614**
Fax: **+36 1 364 2006**
E-mail: export@biolab.hu

Product identification: **Microbiological culture media and supplements**

Product classification: **devices mentioned in Annex VIII of the EU_2017/746 Regulation**

Hereby we declare

under our sole responsibility that the above mentioned devices meet the applicable provisions of the on
EU_2017/746 Regulation on „In Vitro Diagnostic Medical Devices”.

All the supporting documents, as required by Annex II of the EU_2017/746 Regulation in order to prove
conformity to the Essential Requirements as listed in Annex I, are retained under premises of the Manufacturer.

Applicable standard: **ISO 9001**

Place and date: **Budapest, January 2022**

Signature:

 **BIOLAB Zrt.**
1141 Budapest, Öv utca 43.
Adószám: 11838658-2-42

László Ferenci
Managing Director

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according to the EU_2017/746 Regulation on „In Vitro Diagnostic Medical Devices”

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Product identification: **Plastic disposables**

Product classification: **devices mentioned in Annex VIII of the EU_2017/746 Regulation**

Hereby we declare

under our sole responsibility that the above mentioned devices meet the applicable provisions of the EU_2017/746 Regulation on „In Vitro Diagnostic Medical Devices”.

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Applicable standard: **ISO 9001**

Place and date: **Budapest, January, 2022**

Signature:



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Product identification: **Antimicrobial susceptibility discs**

Product classification: **devices mentioned in Annex VIII of the EU_2017/746 Regulation**

Hereby we declare

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Applicable standard: **ISO 9001**

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