

## EC DECLARATION OF CONFORMITY

according to the Directive 2017/746/EC 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](mailto:export@biolab.hu)

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

Product classification: **devices mentioned in Annex VIII of the Directive 2017/746/EC**

***Hereby we declare***

*under our sole responsibility that the above mentioned devices meet the applicable provisions of the Directive 2017/746/EC on „In Vitro Diagnostic Medical Devices”.*

All the supporting documents, as required by Annex II of the 2017/746/EC Directive, 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, 02. 06. 2021.**

Signature:

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

László Ferenci  
Managing Director

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Product identification: **Plastic disposables**

Product classification: **devices mentioned in Annex VIII of the Directive 2017/746/EC**

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Product identification: **Antimicrobial susceptibility discs**

Product classification: **devices mentioned in Annex VIII of the Directive 2017/746/EC**

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