EU Declaration of Conformity

for In Vitro Diagnostic Medical Devices
according to Annex IV of Regulation (EU) 2017/746

Manufacturer: BioMaxima S.A., Vetterów 5, 20-277 Lublin, Poland
SRN: Not Available

Product Name: Mueller Hinton 2 LAB-AGAR™
Basic UDI-DI: Not Available
Classification (IVDR, Annex VIII): Class A, Rule 5

We herewith under our sole responsibility declare that the above mentioned products meet the provisions of the Regulation (EU) 2017/746
The manufacturer is exclusively responsible for the declaration of conformity.

Place and date of issue:
Lublin, 27.05.2022

Signed on behalf of BioMaxima S.A.:

Henryk Lewczuk
VicePresident

Patrycja Paniak -Sankowska
Proxy