Declaration of conformity EU in according to annex IV EU Regulation 745/2017

The manufacturer

M2020 S.r.l.
Address: Via Turcoing, 10/11 – 59100 – Prato (PO)
PEC Address: m2020@pec.it

Declares under their own responsibility that the following medical devices, as described below are:

Name	Medical device named as MIIR
Device classification	Class I Single use Not sterile Type II R
Registration number	BD/RDM 1978002
Model	MIIR
Year of construction	2020

Devices are compliant with:

- Regulation (UE) 745/2017
- UNI EN 14683:2019 – Class I
- ISO 10993-1:2010

Date
21/07/20

Place
Prato

Signature

CE