EU Declaration of Conformity

For a single use medical device class I

The manufacturer:

Franz Mensch GmbH
Werner-von-Siemens-Str. 2
86807 Buchloe
Germany

SRN:	**DE-MF-000021137**

declares under its sole responsibility that the medical device of class I according to Annex VIII of the Regulation (EU) 2017/745

Item REF	**2700B**
Description	**Nitrile gloves Safe Premium | powder-free**
Brand	**Hygostar**
Version	**Colour: blue** **Size: 10/XL** **Length: 24cm** **PU: bag**
Basic – UDI	**40155440110GP**
Intended use	**For third-party protection (protection against germ transmission) in the hospital and care sector**
Applied standards:	**EN 455-1:2020** **EN 455-2:2015+A1:2011** **EN 455-3:2015** **EN 455-4:2009**

complies with all requirements of regulation EU 2017/745 and its annexes in accordance with the conformity assessment procedure set out in annexes II and III of regulation EU 2017/745.

Furthermore, the manufacture and release of the devices are carried out in accordance with the specifications defined in the associated technical documentation, applied standards and normative documents. The medical device bears the CE conformity marking.

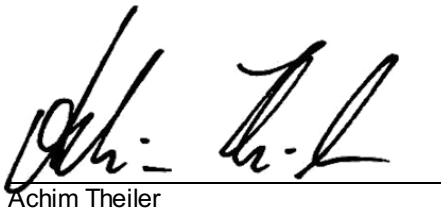
This declaration of conformity is valid until a new declaration of conformity is issued due to the modification of the medical device.

Signed for and on behalf of Franz Mensch GmbH,

Franz Mensch GmbH
Werner-von-Siemens-Str. 2
86807 Buchloe, Germany

phone: +49 8241 9633-0
fax: +49 8241 9633-100
internet: www.franz-mensch.de
e-mail: verkauf@franz-mensch.de

Buchloe, 08.06.2022



Achim Theiler
Management

Franz Mensch GmbH
Werner-von-Siemens-Str. 2
86807 Buchloe, Germany

phone: +49 8241 9633-0
fax: +49 8241 9633-100
internet: www.franz-mensch.de
e-mail: verkauf@franz-mensch.de