

## EU Declaration of Conformity

## for a medical device class I (according to the EU regulation 2017/745, annex IV)

The manufacturer:

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

declares under sole responsibility, that the class I medical device according to the classification rules of the EU regulation 2017/745, annex VIII

Item REF Description Brand Version 2670 Latex gloves Grip | powder-free Hygostar Packing unit: Carton Color: White Size: 10/XL Length: 24cm

complies and meets all the provisions of the conformity assessment procedure of the EU regulation 2017/745 (annex I).

in accordance with confirmed records, test results or certificates, complies/comply with the requirements of the relevant harmonization legislation:

EN 455 1-4

The achieved performance levels of this medical device correspond to:

Signed for and on behalf of Franz Mensch GmbH,

Buchloe, 31.05.2021

Amainda Kreuzma

Head of Quality Management

Updated 31.05.2021

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Franz Mensch is a ISO certified company DIN EN ISO 9001:2015