

## EU Declaration of Conformity

### for Personal Protective Equipment (PPE) of Category 3

The manufacturer:

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

declares under sole responsibility, that the following product

<b>Item REF</b>	<b>27841</b>
<b>Description</b>	<b>Coveralls Type 5B+6B   Microporous</b>
<b>Brand</b>	<b>Hygostar</b>
<b>Version</b>	<b>Colour: white</b> <b>Size: L</b>

complies and meets all the provisions of Regulation (EU) 2016/425 on Personal Protective Equipment (PPE).

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

**EN ISO 13688:2013**  
**EN ISO 13982-1:2004+A1:2010**  
**EN 13034:2005+A1:2009**  
**EN 14126:2003+AC:2004**  
**EN 1073-2:2002**  
**EN 1149-5:2018**

The achieved performance levels of this medical device correspond to:

*EN ISO 13982-1: Typ 5B*  
*EN 13034: Typ 6B*  
*EN 1073-2: Klasse 1*

The notified body

*ANCCP Certification Agency S.r.l. Via dello Struggino 6 57121 Livorno Italien*

*Number 0302*

performed the EU type-examination (Module B) and issued the EU type-examination certificate *PPE-1607-20617-Cert*

The PPE is subject to the conformity assessment procedure:

Module C2 - conformity to type based on internal production control plus supervised product checks at random intervals

under surveillance of the notified body  
*ANCCP Certification Agency S.r.l. Via dello Struggino 6 57121 Livorno Italien*

*Number 0302*

Signed for and on behalf of der Franz Mensch GmbH,

Buchloe, 24.08.2022



Achim Theiler  
General manager