

# EU DECLARATION OF CONFORMITY

The Manufacturer  
**ANSELL HEALTHCARE EUROPE N.V.**  
**RIVERSIDE BUSINESS PARK, BLOCK J**  
**BOULEVARD INTERNATIONAL 55**  
**B-1070 BRUSSELS**  
**BELGIUM**

declares under his sole responsibility, that the PPE described hereafter:

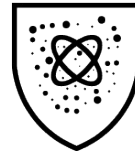
## **AlphaTec® 1500 PLUS FR Model 111** **PPE to be used against category III risks**



**TYPE 5**



**TYPE 6**



**EN 1073-2**



**EN 1149-5**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards , EN ISO 13982-1:2004 + A1:2010, EN 13034:2005 + A1:2009, EN 1073-2:2002, EN 1149-5:2008 (with exception of puncture resistance for which class 1 was achieved.) and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 0598/PPE/24/2525, issued by the Notified Body:

**SGS FIMKO OY (0598)**  
**TAKOMOTIE 8,**  
**FI-00380 HELSINKI,**  
**FINLAND**

and is subject to the procedure set out in Annex VIII (Module D) of the Regulation under the supervision of the Notified Body:

**SGS FIMKO OY (0598)**  
**TAKOMOTIE 8,**  
**FI-00380 HELSINKI,**  
**FINLAND**

**Ulf Nystrom**  
**Sr Manager, Regulatory Affairs PPE Products**

**Place: Malmö**  
**Date: 2024/09/06**