

# 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<sup>®</sup> 1500 PLUS Model 111P**

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 1149-5:2018, EN ISO 13688:2013, (with exceptions for puncture resistance being class 1 and resistance to ignition not tested) 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/23/3096, 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: 2024/10/17