

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