

## 欧盟符合性声明

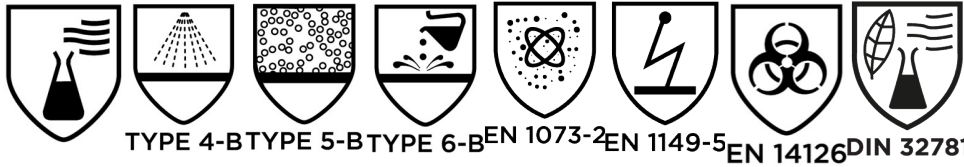
制造商

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

声明以下所述的个人防护设备由其全权负责：

**AlphaTec® 2000 Ts PLUS Model 111**

用于防护category III风险的PPE



is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards , EN 14605:2005 + A1:2009, EN ISO 13982-1:2004 + A1:2010, EN 13034:2005 + A1:2009, EN 1073-2:2002, EN 1149-5:2018, EN 14126:2003, EN ISO 13688:2013, DIN 32781:2010 (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/23/4169, issued by the Notified Body:

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

并须遵守该规例附件VI (模块D) 所载的程序：

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

Ulf Nystrom  
Sr Manager, Regulatory Affairs PPE Products

地点： Malmö  
日期： 2024/01/17