

CE	EC – Declara	nitätserklärung tion of Conformity ration de conformité	Date Day / Month / Year 10.05.2021
Tetcon Global BV Ruysdaelbaan 3G 5642 JJ Eindhoven The Netherlands			
declares, solely responsible, that the medical devices and its accessory			
S-CAPEPLUS Evacuation Mats Model: Basic Model (Art. Nr. 5N88888886), Premium Model (Art. Nr. 5NOUT88886), Youth Model (Art. Nr. 5N888888831), Inflatable Model (Art. Nr. 5NINFL6021), Bariatric Model (Art. Nr. 5N88BAR866)			
meet all the provisions of the regulation MDR 2017/745/EU of the European Parliament and of the council of 5 April 2017 on medical devices (MDR), which apply to him.			
The product is classified in Class 1 according to Regulation MDR 2017/745/EU.			
Applied standards: NEN-EN-ISO 13485:2016 EN NEN-EN-ISO 14971:2019 EN			
Corresponding technical documentation and quality manual is available for examination.			
Eindhoven, 15.04.2021 Place, Date		5642 +31	

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