CE	**EG – Konformitätserklärung** **EC – Declaration of Conformity** **CE - La déclaration 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. 5N88888831), 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

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Signed by Tetcon management and representative: Bob Hezemans