

Declaration of Conformity of Medical Device

Producer: Danish Care Technology ApS
Energivej 3
DK-4180 Sorø
Denmark

This EU declaration of conformity is issued under the sole responsibility of the manufacturer

Product: Epi-Care 3000
Type no: 6070
Classification: Class I

The Classification is made pertaining to MDR Article 51(1) according to Annex VIII, rule 13.

The Epi-Care 3000 is in conformity with Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices (MDR).

Sorø, June 9, 2021

Jens Jørgen Eriksen
Director / CEO


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Signature