Campfield Road
Shoeburyness
SS3 9BX

EU- Declaration of Conformity

issued under the sole responsibility of the manufacturer listed below:

Manufacturer's Name	**Cantel (UK) Ltd**
Manufacturer's address/ SRN	**Campfield Road, Shoeburyness, SS3 9BX. UK**
EU Representative Address/ SRN	**Cantel Medical (Italy) S.r.l. a socio unico Via Laurentina, 169 00071 Pomezia (RM) – Italia**
Quality System Cert No.	**LRQ00001040/B**
Notified Body No./Name	**LRQA**
CE Certificate No.	**N/A**

As the manufacturer listed above, we declare that the devices listed:

Product(s):	**FLEXISLIDE™ Standard Single-Use Patient Transfer Device FLEXISLIDE™ Extra Wide Single-Use Patient Transfer Device FLEXISLIDE™ Mini Size/Extra Wide Single-Use Patient Transfer Device FLEXISLIDE™ Mini Size Single-Use Patient Transfer Device**
REF:	**MW1029, MW1061, 105093,105089**
Basic UDI-DI:	**506018978TDF-00029M2**
Intended Purpose:	**FLEXISLIDE™ Lateral Patient Transfer Device is a single use lateral transfer sheet designed to slide a patient between two surfaces in a pre / peri-operative or ER environment.**
Applied standards:	**EN ISO 13485:2016, EN ISO 14971:2019, EN ISO 20417:2021, EN ISO 15223-1:2016, EN 62366-1:2015+A1:2020, EN ISO 10993-1:2020**

Meets all conformity requirements of the safety and performance requirements of the Medical Devices Regulation (MDR, Regulation (EU) 2017/745 as amended).

Product classification according to the requirements described in Annex VIII *of the Medical Devices Regulation, the medical device is assigned to risk class I (conformity assessment per* Annex IX*).*

Name: **Richard Manford**

Signature: DocuSigned by: 8E1EFA113D314CA...

Title: **Director Regulatory Affairs, EMEA**

Place of Issue: **Shoeburyness, UK**

Date of Issue: **23 May 2021**