easyforce

EU Declaration of Conformity

Manufacturers Name:	Meloq AB
Manufacturers Address:	Drottning Kristinas Väg 53, 114 28 Stockholm, Sweden
SRN:	NA as EUDAMED is not ready yet.
Basic UDI-DI:	07350015485006
Name of the Device:	EasyForce
Product code:	2009006
Classification:	Class 1
Conformity assessment route:	Meloq AB uses the following procedures for the CE-labeling of their products according to the Regulation MDR: Class 1: EC conformity declaration according to annex VIII + annex IX.

This declaration of conformity is issued under the sole responsibility of Meloq AB. We hereby declare that he medical device specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices.

All supporting documentation is retained at the premises of the manufacturer.

Signature:

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Jerker Skogby
CEO

Place and date of issue:

Stockholm, 2021-07-01

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Document Name:
2009-1RE026-02 EasyForce Declaration of Conformity