GETINGE

EU Declaration of Conformity for Medical Device

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Name and Address of the Manufacturer:	MAQUET GmbH Kehler Str. 31 D-76437 Rastatt
Single Registration Number:	DE-MF-000009119

On our sole responsibility, we hereby declare that the product(s)

Product- / Trade Name:	Pads, See Annex I_EN
Intended Purpose:	The pads are designed for the placement and positioning of the patient, immediately prior to, during and after surgical.
Reference-No.:	Refer to Annex I_EN of this document
Basic UDI-DI (acc. to Part C of Annex VI):	40467680122AB8
Classification (acc. to Annex VIII):	Class I

comply with the relevant provisions of the following Regulation(s) and Directive(s):

Regulation (EU) 2017/745 on Medical Devices

Conformity Assessment Procedure:	Acc. to Annex II and Annex III of Regulation (EU) 2017/745
Common Specifications used:	N/A

CB

This declaration of conformity is valid from date of issue until 19 January 2030.

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ANNEX I_EN

The products can be delivered in the following variants / with the following components:

Product-No.	Product- / Trade Name	Basic UDI-DI
1000.24A0	Padded roll for countertraction post	40467680122AB8
1000.5600	Pad	40467680122AB8
1000.5700	Head ring	40467680122AB8
1000.68C0	Pad	40467680122AB8
1000.6900	Plexus cushion	40467680122AB8
1000.77A0	Tunnel cushion	40467680122AB8
1001.94A0	Leg rest	40467680122AB8
1003.7400	Pad	40467680122AB8
1180.41A0	Pad for rectal positioning	40467680122AB8
1180.56A0	Transfer board pad	40467680122AB8
4000.05A0	Pad	40467680122AB8