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EU Declaration of Conformity for Medical Device

Document ID: 1067273 v02

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:	Universal table top, see Annex I_EN
Description:	The table tops are designed for the placement and positioning of the patient immediately prior to, during and after surgical interventions, as well as for examination and treatment.
Reference-No.:	See Annex I_EN
Basic UDI-DI (acc. to Part C of Annex VI)**:**	40467680118ABM
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

Directive 2011/65/EU on the restriction of the use of certain substances in electrical and electronic equipment

This declaration of conformity is valid from date of issue until 2025-05-25.

Rastatt, 2021-08-06

Holger Ullrich, Senior Director Regulatory Affairs, Center of Excellence EMEA
Signed on behalf of MAQUET GmbH

ANNEX I_EN

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

Product-No.	Product Description	Basic UDI-DI
1160.10D0	Universal table top	40467680118ABM
1160.30D0	Universal table top	40467680118ABM