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TÜV SÜD Product Service GmbH · Ridlerstr. 65 · 80339 Munich · Germany

B. Braun Medical AG
Seesatz 17
6204 SEMPACH
SWITZERLAND

Via Email: Michael.Gluschke@bbraun.com

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
CBW 61585	PS-MHS-AU-3/ Dr. Gertrud Schönmann	+49 89 50084-978 Gertrud.Schoenmann@tuvsud.com		2024-05-23 Dokument2	1 of 2

TÜV SÜD Product Service GmbH
Receipt of formal application

Reference: 713337991 | 2156110

To whom it may concern,

Confirmation of the status of a formal application in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 (in the following referenced as MDR) as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.

With this letter TÜV SÜD Product Service GmbH, designated under MDR and identified by the number 0123 on NANDO, confirms that we have received **a formal application** in accordance with Section 4.3, first subparagraph of Annex VII of MDR with the above stated manufacturer with the following SRN Number:

SRN Number: CH-MF-000017781

The devices covered by the formal application mentioned above are identified in the Table below.

Please note that this letter only confirms the status of the formal application.

To benefit from the additional transitional provisions in the framework of Regulation EU 2023/607, TÜV SÜD Product Service GmbH and the manufacturer need to sign a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR latest until 26 September 2024.

In case of inquiries please contact medical_devices@tuvsud.com.

On behalf of the Notified Body TÜV SÜD Product Service GmbH,

Registered Office: Munich
Trade Register Munich HRB 85 742
UniCredit Bank AG · BIC HYVEDEMMXXX
IBAN DE13 7002 0270 0048 8522 11
VAT ID No. DE129484267
Information pursuant to § 2 [1] DL-InfoV
(Germany) at tuvsud.com/imprint

Supervisory Board:
Holger Lindner (Chairman)
Board of Management:
Walter Reithmaier (CEO)
Patrick van Welij

TÜV SÜD Product Service GmbH tuvsud.com/ps
Nicht-Aktive Medizinprodukte / Non-Active Medical Devices
Hotline: +49 89 50084-747
Ridlerstr. 65
80339 Munich
Germany





2024-05-23

TÜV SÜD Product Service GmbH
Medical and Health Services

Gertrud Schönmann

Gertrud Schönmann (23. Mai 2024 12:05 GMT+2)

Dr. Gertrud Schönmann (PH)
Conformity Assessment Responsible (CARE)

Devices covered by the formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR

Device name or Basic UDI-DI (under MDR application)
Device 1 Uro-Tainer PHMB Basic UDI-DI 40392390000009262W