

For the Class IIb medical device

modula

a Central Reverse Osmosis System

with serial numbers 20 MODULA 0001 to 20 MODULA 9999,

we,

DWA GmbH & Co. KG Großer Sand 8 76698 Ubstadt-Weiher Germany,

declare in our function as manufacturer and under our sole responsibility conformity with the requirements of the following directive:

93/42/EEC

Council directive issued 14 June 1993 concerning medical devices last amended by directive 2007/47/EC.

Conformity of the named product with the Essential Requirements of Annex I of the directive is demonstrated in technical documentation.

All applicable harmonized standards are met.

Compliance with the requirements is demonstrated with the certificates of conformity according to Medical Devices Directive 93/42/EEC Annex II (exclusive 4), Reg. No. 44 232 150580.

Notified Body

EC reference no. 0044 TÜV NORD CERT GmbH Langemarckstraße 20 45141 Essen Germany

Ubstadt-Weiher, 2020-02-07

B. Gegenhuber, General Manager