

EU - DECLARATION OF CONFORMITY

according to Regulation (EU) 2017/745 article 19 read in conjunction with annex IV

Hereby the manufacturer

anmed GmbH

Gerichtsstr. 140
09474 Crottendorf
Federal Republic of Germany

SRN: DE-MF-000006213

declares under sole responsibility that the named product

ANABOX® DAILY BOX

ALSO THE BRAND NAMES CINIBOX UND SUPAIRBOX

MEDICATION DISPENSER FOR 1 DAY

BASIS - UDI: PP01228ABXDAILY83

complies with the relevant provisions of Regulation (EU) 2017/745 of April 5,
2017 on medical devices.

A conformity assessment procedure was carried out in accordance with article 52 (7) of Regulation
(EU) 2017/745.

In accordance with annex VIII rule one of Regulation (EU) 2017/745 the product mentioned above is
classified as

RISK CLASS I MEDICAL DEVICE I

anmed GmbH

Gerichtsstraße 140

D-09474 Crottendorf

Tel.: +49(0) 37344 763800

DE 243016119 / HRB 18717 Chemnitz

Mail: info@anmed.de

www.anabox.de

SEBASTIAN RICHTER

operating manager / proxy

CROTTENDORF, 2021/07/01

place, date