ANABOX®

anmed GmbH

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® 1 x 7

ALSO THE BRAND NAMES CINIBOX UND SUPAIRBOX

MEDICATION DISPENSER FOR 1 WEEK

BASIS – UDI: PP01228ABX7INONE76

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 225116 119 / HRB 18717 Chemnitz
Mail: info@anmed.de
www.anabox.de

CROTTENDORF, 2021/07/01

place, date

SEBASTIAN RICHTER

operating manager / proxy