EU – DECLARATION OF CONFORMITY

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

Hereby the manufacturer

anmed
Am Gewerbegebiet 5
09474 Crottendorf
Federal Republic of Germany

SRN: DE-MF-000006213

declares under sole responsibility that the named product

ANABOX® VIDA4

(WEPA, CINIBOX, SUPAIRBOX AND OTHER PRIVATE LABELS)

MEDICATION DISPENSER FOR 1 WEEK

BASIC UDI-DI: 4260041530085.VIDA4DH

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 MEDICAL DEVICE RISK CLASS I

INTENDED PURPOSE:

Medication dispensers to alleviate or compensate for injuries or disabilities

CROTTENDORF, 2024/09/01
place, date

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