



EUMEDICAL
EUROPA S.R.L.S

EC Declaration of Conformity

Manufacturers Name: Eumedical Europa S.r.l.s

Manufacturers Address: Via Piastrarelle, 9 00075, Lanuvio, Roma

Authorized Representative Name: Luigi Rosario Franco Inderst

Name of the Device: **QUANOCENS PRONTOUSO 750 ml, QUANOCENS PRONTOUSO 1000 ml QUANOCENS PRONTOUSO 5 Litri QUANOCENS CONCENTRATO 500 ml QUANOCENS MONODOSE box 30 da 10 ml**

Classification: Class I CND D99

Conformity assessment route: Eumedical Europa Sr.l.s. uses the following procedures for the CE-labeling of their products according to Rule 5 of MDD 93/42/CE and Rule 16 and art 2 of MDR 2017/745.
The product is produce in a manufacturing plant provided of the ISO 9001 and ISO 13485 certification

This declaration of conformity is issued under the sole responsibility of Eumedical Europa Sr.l.s. We hereby declare that he medical device(s) specified above meet the provision of the MDD 93/42/CE and, following, the Regulation (EU) MDR 2017/745 for medical devices.
All supporting documentation is retained at the premises of the manufacturer.

Signed for and on behalf of Eumedical Europa srls

Pf Dr Inderst Luigi Rosario Franco

Place and date of issue: Lanuvio

Research and development affair

Lanuvio 09.06.2022

NAME

FUNCTION

.....

EUMEDICAL EUROPA S.R.L.S.
VIA PIASTRARELLE, 9 00075 LANUVIO (RM)
P.IVA 14228071008
REA : RM1505537

.....