



DECLARATION OF CONFORMITY

The company Curasept S.p.A., with registered office in Via G. Parini 19, I-21047 Saronno (VA)

Declares

that the Medical Device described below:

CURASEPT ADS DNA MOUTHWASH

risk class III based on rule 13 and 17 of Annex IX of Directive 93/42/EEC (absorbed into Italian law by Legislative Decree No. 46 of 24 February 1997 and subsequent amendments)

is manufactured according to the quality system which meets the requirements set forth in Annex II of the aforementioned rule, as per certificate No. QCT-0141-19 Addendum n° 01-20 issued on 10.10.2019 by the I.S.S. (Istituto Superiore Sanità), in Rome, Italy, expiration 26.05.2024,

conforms to the essential requirements and the provisions of the Directive 93/42/EEC according to Annex II of the above Directives as reported in the Certificate No. EPG-0249-19 Addendum n° 01-20 issued on 10.10.2019 by I.S.S., Rome, Italy, expiration 26.05.2024 (Notified Body No. 0373).

Saronno, 15/07/2020

Stefano GIOVANNARDI
General Director

Curasept S.p.A.

Via G. Parini 19, 21047 Saronno (VA), Italy, Tel. 02 9622 799, Fax 02 9670 9243
info@curaseptspa.it, www.curaseptspa.it
Cap. soc. € 3.000.000,00 i.v., R.E.A. VA 284921, C.F. P.IVA 13268170159