

EC Declaration of Conformity

We,

Vertex-Dental B.V.
Centurionbaan 190
3769 AV Soesterberg
The Netherlands

Hereby declare under our sole responsibility that the CE marked product to which this declaration relates,

Dental LT Clear

Productcodes are available in the annex.

has been classified as Class IIa, according to Annex IX, rule 5, and is in conformity with the essential requirements and provisions of the Council Directive 93/42/EEC concerning medical devices as amended by Directive 2007/47/EC.

and is conformity with EN ISO 1641:2009; EN ISO 10993-1:2009/AC2010; EN ISO 10993-3:2009, EN ISO 10993-5:2009 and EN 908:2008.


and is subject to the procedure set out in Annex II (excluding section 4) of the Council Directive 93/42/EEC as amended by Directive 2007/47/EC.

This declaration is made on base of the EC Certificate Full Quality Assurance System N° BE12/223574916 Issue 9, valid until 17-12-2020 delivered by:

Notified Body n° 0120
SGS United Kingdom Ltd.
202B Worle Parkway
Weston-super-Mare, BS22 6WA
United Kingdom

This Declaration of conformity covers batches produced as of batches:
XK234N01

Soesterberg, June 08 2017


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Mrs. N.C. Peterse – van der Koppel
General Manager Research & Development

Annex I: Product references Dental LT Clear

Description	Size	Productcode
Formlabs Dental LT Clear / Clear	1000 ml	NPFLOCCLE01000