

DECLARATION OF CONFORMITY



Ultradent Products, Inc., 505 W. Ultradent Drive (10200 S.), South Jordan, UT, USA 84095 has evaluated the following product by using the Conformity Assessment Procedure of Annex II of the Medical Device Directive 93/42/EEC, as amended by 2007/47/EEC:

UltraCal XS

and confirms in sole responsibility that the product is compliant with the Essential Requirements of Annex I of the Medical Device Directive 93/42/EEC. Technical documentation is located in the Regulatory Affairs Department.

This product system is classified as Class IIa medical device according to the Medical Device Directive 93/42/EEC, Annex IX, Section III, Rule 8

UMDNS Code: 16182 Cavity Liner

GMDN Code: 47232 Calcium hydroxide dental suspension

EMDN Code: Q01010201 Root Canal Filling Dental Pastes

EC Representative:

Ultradent Products GmbH
Am Westhover Berg 30
51149 Cologne
Germany

Notified Body:

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ID No. 0044


Adam Black, RAC

Senior Manager – Regulatory Affairs

22 Feb 2024
Date

State of Utah
County of Salt Lake

Subscribed and sworn to before me on this 22 day of February 2024

By Adam Black


Notary Public



This document is in force as long as the following EC certificates are valid:

EC Certificate 44 232 090234 with valid extension through 31 December 2028