

DECLARATION OF CONFORMITY

Ultradent Products, Inc. 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:

Opalescence Quick PF 45%

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 Classification 2.2, Rule 6

UMDNS Code: 17619, Restorative Materials, Dental, Other

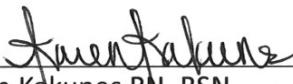
GMDN Code: 38785, Dental bleaching agent

EC Representative:

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Am Westhoven Berg 30
51149 Cologne
Germany

Notified Body:

TÜV Nord Cert GmbH
Unternehmensgruppe TÜV Nord
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ID No. 0044


Karen Kakunes RN, BSN

Regulatory Affairs Management


Date

State of Utah
County of Salt Lake

Subscribed and sworn to before me on this 2 day of December 2020

By Karen Kakunes

Amy Henderson - Nielson
Notary Public



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

EC Certificate 44 232 090234 valid through 26 May 2024