



# DECLARATION OF CONFORMITY

Ultradent Products Inc. has evaluated the following product by using the Conformity Assessment Procedure of Annex VII of the Medical Device Directive 93/42/EEC, as amended by 2007/47/EEC:

**VALO<sup>®</sup>, VALO<sup>®</sup> Ortho, and Accessories  
(Lenses, Light Shield, Battery Pak, Barrier Sleeves, power supply, etc)**

and confirms 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 I medical device according to the Medical Device Directive 93/42/EEC, Section III Classification, 3.3 Rule 12.

**UMDNS Code:** 16386 Lights, Dental Activator  
**GMDN Code:** 35775 Light, Dental, Polymerization Activator

Ultradent Products GmbH  
Am Westhover Berg 30  
51149 Cologne  
Germany

Marie Hess  
Marie Hess  
Sr. Regulatory Affairs Associate

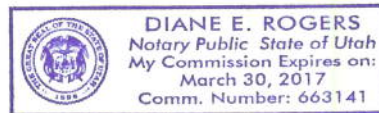
5 Mar 15  
Date

State of Utah  
County of Salt Lake

Subscribed and sworn to before me on this 5 day of March 2015

By Marie Hess

Diane E. Rogers  
Diane Rogers, Notary Public



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

EC Certificate 44 232 090234  
EN ISO 13485 44 221 090234