

DECLARATION OF CONFORMITY

VALO CORDLESS

Ultradent Products Inc., 505 West Ultradent Drive (10200 South), South Jordan, UT, 84095, has evaluated this product by using the Conformity Assessment Procedure of Annex IX of the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on Medical Devices, and confirms that this product is compliant with the General Safety and Performance Requirements of Annex I of the Regulation (EU) 2017/745. Technical documentation is located in the Regulatory Affairs Department.

Identified Legislations:	<i>Regulation (EU) 2017/745 – European Parliament and of the Council of 5 April 2017 on medical devices</i>
EC Representative:	Ultradent Products GmbH Am Westhover Berg 30 51149 Cologne Germany
Device Classification:	This product system is classified as a Class I medical device according to Regulation (EU) 2017/745, Annex VIII, Chapter III, Section 6.5, Rule 13.
Intended Purpose:	Source of illumination for curing photo-activated dental restorative materials and adhesives.
UMDNS Code:	16386 – Light, Dental Activator
GMDN Code:	35775 – Light, Dental, polymerization activator
CND Code:	Z12119003 – Cold Light Dental Equipment
Basic UDI-DI:	0883205-TF086BF
Eudamed SRN:	DE-AR-000006666, DE-IM-000010855, US-MF-000013697
Common Specifications:	N/A
Model Numbers:	5941-
Images:	

This Declaration of Conformity is issued under the sole responsibility of the manufacturer and is in force as long as the following certificate is valid:



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DIN EN ISO 13485:2021 Reg.-No. 44 221 090234 valid until 05 March 2029

Adam Black

Director - Regulatory Affairs & Clinical Operations

23 Feb 2026

Date

State of Utah, County of Salt Lake

Subscribed and sworn to before me on this

23 day of February 2026

By Adam Black

Notary Public

