EU Declaration of Conformity

TO REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL

Manufacturer:	Guilin Refine Medical Instrument Co., Ltd. Address: NO.8-3, Information Industrial Park, High-Tech Zone, Qixing District, Guilin, Guangxi, 541004, P.R.China SRN: CN-MF-000012204
European Representative:	MedNet EC-REP C IIb GmbH Address: Borkstraße 10, 48163 Münster, Germany SRN: DE-AR-000011194
Product Name:	Curing Light
Model:	MaxCure 3, MaxCure 5, **MaxCure7**, MaxCure 9, SWAN, **SWAN+**, A-Cure, A-Cure Plus, MaxCureG, MaxCureH.
GMDN Code:	35775
Basic UDI-DI:	697156045RCLS8
Device Photograph:	Please refer to section 1.7 Device Photograph in RF-RCL-T004 Device Description and Specification.
Intended Use:	Curing Light is intended for use by trained dental professionals for the purpose of light curing dental resins and composites.
Risk Class:	Class I
Classification Rule:	Rule 13 in Chapter III of Annex VIII of the Regulation (EU) 2017/745
Conformity Assessment Route	Annex II & III of the Regulation (EU) 2017/745
We herewith declare that the above-mentioned product(s) meet the Regulation (EU) 2017/745 of THE EUROPEAN PARLIAMENT AND OF THE COUNCIL and the transposition into national law. All supporting documentation is retained at the premises of the manufacturer. We, the manufacturer, are exclusively responsible for the DoC.	
Applied Standards:	See RF-RCL-T003 Applied Standard List
Place, Date of Issue:	Guilin, Guangxi, 2023-9-11
Signature:	Name: Judong Chen Function: Management representative