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Version: A

EU Declaration of Conformity

Manufacturer:	**Guilin Kevin Peter Technology Co., Ltd.** North side of Linsu Rd, Lingui Town, Lingui Dist., Guilin, Guangxi 541100, P. R. China
SRN of Manufacturer:	CN-MF-000022934
European Representative:	**MedNet EC-REP C IIb GmbH** Borkstrasse 10, 48163 Muenster, Germany
SRN of European Representative:	DE-AR-000011194
Product Name:	Surgical Headlight
Models:	i Light Wireless
Classification:	I / Rule 1 & Rule 10 (MDR, Annex VIII)
EMDN Code:	L0399 GENERAL SURGERY INSTRUMENTS, REUSABLE - OTHER
Basic UDI-DI:	69753843318Z5
Conformity Assessment Route:	Annex II, Annex III of (EU)2017/745, MDR

We herewith declare that the above-mentioned product(s) meet the Regulation (EU) 2017/745. All supporting documentations are retained under the premises of the manufacturer. We are exclusively responsible for the Declaration of Conformity. General applicable regulations: Medical Device Regulation, (EU)2017/745. Every article we sell is covered by a technical file. Technical files are provided to the European Authorized representative and kept up to date by us.

Name: Wei kunsong

Position: PRRC

Location: Guilin, China

Date: 2026/03/10

Signature: