EU Declaration of Conformity

In accordance with the Regulation (EU) 2017/745 on medical devices (MDR 2017/745)

Manufacturer's Name:	ILLUCO Corporation Ltd.
Manufacturer's Address:	102-304 SK Ventium, #166 Gosan-ro Gunpo-si, Gyeonggi-do 15850 Republic of Korea
Manufacturer's SRN (Single Registration Number):	KR-MF-000012082
Authorized Representative Name (if applicable):	AeMi World e.K.
Authorized Representative SRN (Single Registration Number):	DE-AR-000007704
Authorized Representative Address (if applicable):	Bugenhagenstr.8, 10551 Berlin, Germany Tel: 0049 (0) 172 3199 811 e-mail: dwis22@gmx.de
Basic UDI-DI:	88001906IDS1100BK
Description of the Device(s):	Dermatoscope IDS-Series Class I, non-sterile and non-measuring medical devices
Name of Device(s):	IDS-1100, IDS-1100C, IDS-1000, IDS1000 PLUS, IDS-3100
Risk Class:	Medical Device Class I according to the Annex Ⅷ
Notified Body:	Not required for medical device class I
Conformity assessment route:	The procedures according to the Regulation (EU) 2017/745 on medical devices: Class I: EU conformity declaration according to Annex I, Annex II, Annex III, and Annex XIV.
References to the relevant harmonized standards or specifications in relation to this conformity:	ISO 13485 Medical devices – Quality management system – Requirements for regulatory purposes EN 60601-1-2 Medical electrical equipment – Part 1-2 General requirements for basic safety and essential performance -Collateral Standard: Electromagnetic disturbances - Requirements and tests

This declaration of conformity is issued under the sole responsibility of ILLUCO Corporation Ltd. We hereby declare that the medical device(s) specified above meet the provision of the **Regulation (EU) 2017/745 on medical devices (MDR 2017/745).**

Signature: ILLUCO CO., LTD ip Hong PRESIDENT / ILPYO. HONG

……

Name: Ilpyo Hong

Function: CEO

Place and date (dd.mm.yyyy) of issue:

Gunpo, Republic of Korea, 01.01.2023

…….