LABOMED

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CE EU DELCLARATION OF COMFORMITY CE

Labo America

Declare under sole responsibility that

Our medical devices: Slit lamps

Models: SL 20i / SL 25i
Model Nos.: 8126700, 8126701 / 8127700, 8127701
Models:SLx 40, SLx 45
Model Nos.: 8126400,
8126401 / 8127400, 8127401
Models: eVO 300 / eVO 400 / eVO 350 / eVO 450
Model Nos.: 8126200, 8126201 / 8127200, 8127201 / 8126300, 8126303 / 8127300, 8127303
Basic UDI-DI 81017855SLITLAMPAT

Are AC powered devices intended for use in eye examination of the anterior eye segment, from the cornea epithelium to the posterior capsule and used to aid in the diagnosis of diseases or trauma which affects the structural properties of the anterior eye segment.
Have been subjected to conformity assessment procedure complying with Article 52 (7) and technical documentation accordance to Annexure II and III of council regulation (EU) 2017/745.

These Devices are Class 1 (without a measuring function) per Rule 13 annex VIII of the regulation MDR 2017/745

Labo America designs and manufactures all products under Quality Systems registered to meet the requirements of ISO 9001, ISO 13485, 21CFR 820 and MDR 2017/745. In case of alteration of the device, not agreed upon by us, this declaration will lose its validity.

Applied harmonized standards, national standards and other normative documents:

EN/ISO: 15223-1 EN
ISO: 10993-1
EN /IEC 60601-1-1
EN /IEC 60601-1-2
EN/ISO: 14971
EN/ISO: 10936
EN 62366

EN 61010-1
EN/IEC 61326-1
EN/IEC 61000-3-3
EN/IEC 61000-3-2
CISPR11/EN 5501

AUTHORIZED SIGNATORY

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EC REP
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