



EU DECLARATION OF CONFORMITY

HEINE Optotechnik GmbH & Co. KG
Dornierstr. 6, 82205 Gilching, Germany
www.heine.com

Single Registration Number: DE-MF-000006269

Medical device

Product family: Retinometer
Product group: LAMBDA 100

We hereby declare, under our sole responsibility, the conformity of the following product in accordance with the Regulation (EU) 2017/745 of the European Parliament and of the Council of 05 April 2017 concerning medical devices.

Product name	LAMBDA 100
Basic UDI-DI	4053755_RM_01_8Y
GMDN	35148
UMDNS	14-382
EMDN	Q0299
Classification	Class I according annex VIII




HEINE Optotechnik GmbH & Co. KG hereby declares that the product covered by this declaration is in conformity with this Regulation and, where applicable, with other relevant EU legislation providing for the issuing of an EU declaration of conformity.

References to any common specifications: None

Conformity assessment procedure chosen: Technical documentation according Annexes II and III

This declaration of conformity is valid until a revised declaration of conformity is issued.

Gilching, 12 July 2024
(Place and date of issue)


Thomas Sauerer / PRRC
(Name/function and signature)

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