



EU DECLARATION OF CONFORMITY

We,

IDCP BV
Manuscriptstraat 12-14
1321 NN Almere
The Netherlands
SRN: NL-MF-000000755

hereby declare under our sole responsibility that the CE-marked products to which this declaration relates,

Dino-Lite CapillaryScope Basic UDI-DI: 87202991629MEDLN97

CapillaryScope 500 (type number MEDL4N5),
CapillaryScope Pro 200 (type number MEDL4N Pro),
CapillaryScope Pro 200 Wireless (type number MEDLW4N Pro),
CapillaryScope Pro 500 (type number MEDL4N5 Pro)
and
CapillaryScope Pro 500 Wireless (type number MEDLW4N5 Pro)

having the intended purpose: The CapillaryScope is intended to make images of microcapillary vessels,

and have been classified as Class I, according to Annex VIII, Rule number 10, and the related DinoCapture software is classified as Class I, because the software drives the CapillaryScope or influences the use of CapillaryScope and consequently shall fall within the same class as the CapillaryScope,

and are in conformity with the General Safety and Performance Requirements of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices,

and are in conformity with the standards EN-ISO 20417:2021 (cor. 2022-04) and EN-ISO 15223-1:2021, EN ISO 15223-1:2021/Amd 1:2025,

and are in conformity with the requirements of directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment.

Signature:

Almere, The Netherlands
Date: Nov 5, 2025
Name: Jan Boers
Function: Managing Director