Number: 2246552CE01

EU Quality Management System Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter I and III

Manufacturer:

Implandata Ophthalmic Products GmbH

Kokenstraße 5
30159 Hannover
Germany
SRN ID.: DE-MF-000005431

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EU- Regulation which apply to them:

0344

Supplement to certificate: 2188436CN

Additional certificate: 2246552TD01

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant requirements of EU Regulation 2017/745, including all subsequent amendments for the above mentioned conformity assessment. The manufacturer/ authorized representative is subject to periodic surveillance as required for the applicable conformity assessment in accordance to Regulation 2017/745.

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Principal Certification Manager

First Issued: **17 June 2021** Date: **24 June 2021** Expiry date**: 1 June 2026**

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 www.dekra-product-safety.com Company registration 09085396

Number: 2246552CE01

EU Quality Management System Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter I and III

This certificate covers the following device(s) / groups of device(s):

Class III
eyemate®-IO implant, 11.3mm, 11.7mm , 12.1mm (class III)
eyemate®-IO/KP, 11.3mm, 11.7mm , 12.1mm (class III)
eyemate®-SC implant (class III)
eyemate®-Reader Set, eyemate®-Key, eyemate®-Cable Antenna
Medicel ACCUJET Injector EM-IO 1.1 Silicone padding (class III)

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Conditions for or limitations to the validity of this certificate:

- N/A

Certificate History

Identification of the Common Specifications and Harmonized Standards complied with are documented within the technical documentation and audit assessments carried out. These are traceable through the DEKRA Certification B.V. Certification Notice. The Certification Notice also identifies the necessary information related to the quality management system of the manufacturer, including facilities.

Date of Issue certificate	Certification Notice Reference	Action
17 June 2021	2188436CN08	First issue
24 June 2021	2188436CN08.1	Revised

First Issued: **17 June 2021** Date: **24 June 2021** Expiry date**: 1 June 2026**

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