EC Declaration of Conformity

in accordance with EC Directive 93/42/EEC on Medical Devices

Manufacturer Carl Zeiss Meditec AG, Goeschwitzer Strasse 51-52, 07745 Jena, Germany

We Carl Zeiss Meditec AG herewith declare with sole responsibility that the following Medical Device meets the Requirements of the European Directive 93/42/EEC. The device is provided with CE Marking with the number of the Notified Body.

Product identification:	***Autorefractor / Keratometer***
Medical Device Trade Name:	***VISUREF 150***
Models/Reference:	***VISUREF 150***
Accessories:	***n/a***
Medical Device Class:	
MDD 93/42/EEC	***Class IIa***
Conformity Assessment Procedure :	***Annex II of MDD 93/42/EEC***
Scope of Application:	This Declaration of Conformity is valid for all products manufactured until 2021-11-29.
UMDNS code:	***12-811, 13-313***
GMDN code:	***12811, 36386***

Digitally signed by Cucerenco Olesea
Date: 2021.07.16 13:00:19 EEST
Reason: MoldSign Signature
Location: Moldova

Notified Body: DQS Medizinprodukte GmbH, August-Schanz-Straße 21, 60433 Frankfurt - notified under 0297.

Any Modification to the product not authorized by Carl Zeiss Meditec AG will invalidate this Declaration.

i. V. Dr. Christian Münster
Head of CoCe
Ophthalmologic Diagnostics and Therapy Jena

i.V. Peter Schruttka-Rechtenstamm
Senior Director Regulatory Affairs

Jena, 2019-03-11