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.

Product identification:	***Intraocular lens***
Medical Device Trade Name:	***AT LISA tri 839MP***
Models/Reference:	*Serialnumber identification: XXXXXXXX0XXX*
Accessories:	*Including all options and accessories as defined in the User Manual*
Medical Device Class MDD 93/42/EEC:	***Class IIb***
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 classification:	***16-071***
GMDN Code	***35658***

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. Guillaume Gasc
Vice President Competence Center &
Order fulfillment consumables

i.V. Christina Jakob
Director Clinical & Global Regulatory Affairs
Implants / OVDs / Sterile Products

Berlin, 2016-11-22

Carl Zeiss Meditec AG, Berlin Site, Max-Dohrn-Str. 8-10; 10589 Berlin, Germany