



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: XXXXXXXXXXX0XXX*

Accessories: *Including all options and accessories as defined in the User Manual*

Medical Device Class *Class IIb*
MDD 93/42/EEC:

Conformity Assessment Procedure *Annex II of MDD 93/42/EEC*

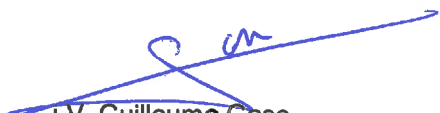
Scope of Application: *This Declaration of Conformity is valid for all products manufactured until 2021-11-29*


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-Dohm-Str. 8-10; 10589 Berlin, Germany