EC Declaration of Conformity

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

Manufacturer Carl Zeiss Suzhou Co., Ltd., Modern Industrial Square 3-B, No. 333 XingPu Road, Suzhou Industrial Park, 215126, Suzhou, China

We Carl Zeiss Suzhou Co., Ltd., 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:	*Surgical Microscope*
Medical Device Trade Name:	*EXTARO 300*
Models/Reference:	*6033*
Accessories:	*Video systems, HD camera, External camera adapter, Fluorescence Mode*
Medical Device Class: MDD 93/42/EEC	*Class I*
Conformity Assessment Procedure :	*Annex VII of MDD 93/42/EEC*
Scope of Application:	This Declaration of Conformity is valid for all products manufactured until 2020-03-14.
UMDNS code:	*12-539*
GMDN code:	*35191*
Standards Applied:	EN ISO 14971:2012 EN 60601-1:2006+A1:2013+A11:2011+A12:2014 EN 60601-1-2:2015 EN 60601-1-6:2010 EN 62366:2008 EN 62471:2008 EN ISO 13485:2012+AC:2012

We established and maintain a Quality Management System in accordance to EN ISO 13485:2012+AC:2012 which has been audited by DQS Medizinprodukte GmbH, August-Schanz-Straße 21, 60433 Frankfurt.

Any Modification to the product not authorized by Carl Zeiss Suzhou Co., Ltd. will invalidate this Declaration.

Lei Xie
General Manager

Xiaodong, Huang
Representative of Quality Management

Suzhou, 2017-03-15.