Certificate of CE-Registration

This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

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USA

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

UMDNS Code	Description	Classification	Registration Number
10573	Eye Cannula	IIa	DE/CA09/0760/A15/007-01
12440	Markers	I	DE/CA09/0760/A15/004-01
15621	Microsurgical Instrument	I	DE/CA09/0760/A15/003-01
16223	Positioning Aids	I	DE/CA09/0760/A15/006-01
17543	Inserters	I	DE/CA09/0760/A15/002-01
17544	Inserters, Intraocular Lens, non-sterile, non-measuring	I	DE/CA09/0760/A15/001-01
17596	Cataract Extraction Units, Phacoemulsification	I	DE/CA09/0760/A15/005-01

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 23 November 2018

Werner Sander
President