

EC Certificate

The LGA InterCert GmbH – **Notified Body 1275** according to the EC Directive concerning medical devices – herewith confirms that the following company



MöllerMedical

Möller Medical GmbH
Wasserkuppenstraße 29 - 31
36043 Fulda
Germany

applies a

Full Quality Assurance System

for the design, manufacture and final inspection of the medical devices
as specified in the enclosure within the scopes:
fine-mechanical and electronic medical devices, assembly groups and single parts,
sterile disposables for medical technology and medical analyzing

according to

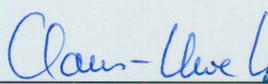
Directive 93/42/EEC, Annex II excluding (4).

In a certification audit carried out by LGA InterCert GmbH, evidence was furnished proving that the quality system of the company fulfills the requirement of the above-mentioned directive.
The surveillance through the LGA InterCert GmbH is effected according to Annex II (5).

This certificate is valid until June 30, 2016
Registration no.: **1911833-006-000**

Audit report no.: 1911833
of: May 27, 2011

Nuremberg, July 1, 2011


Claus-Uwe Hinrichs
Certification Office
Medical Devices



It is allowed for the company to sign the products specified in the enclosure with the adjoining sign.

CE 1275