EC Certificate Full Quality Assurance System: Certificate US15/842192

The management system of

Katena Products, Inc.

23175 224th Pl. SE, Suite C,
Maple Valley, WA, 98038, United States

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Sterile ophthalmic lenses, forceps, and lens supports for diagnostic use in the examination of the eye and to aid visualization during surgery.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 13 April 2016 until 5 June 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 28 June 2016
Issue 3. Certified since 5 June 2015

Certification is based on reports numbered WW/MW 604442

Authorised by

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