EC Certificate Production Quality Assurance System: Certificate
US13/82939

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 V
Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions.

For the following products

Annex V Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions:

Sterile transient ophthalmic lenses.

This certificate is valid from 13 April 2016 until 6 August 2018
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 28 June 2016
Issue 3. Certified since 6 August 2013

Certification is based on reports numbered WW/MW 604442

Authorised by

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