

# EC Certificate

## FULL QUALITY ASSURANCE SYSTEM

### Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

**Certificate Number**  
41314041-01

**Initial Certification Date**  
December 13, 2001

**Certificate Valid from**  
December 14, 2016

**Certificate Expiry Date**  
December 13, 2021

*The certification is subject to the organization maintaining their system in compliance with the regulations stated in this certificate, allowing regular assessments and following the contracted requirements of the Notified Body.*

*Intertek Semko AB is a Notified Body according to Directive 93/42/EEC on medical devices, with identification number 0413.*

*Intertek Semko AB  
Box 1103, SE-164 22 Kista,  
Sweden  
Telephone +46 8 750 00 00  
medtechsweden@intertek.com*



Ackred. nr 1003  
ISO/IEC 17021

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the Swedish national legislation LVFS 2003:11 to which the undersigned is subjected, transposing Annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive, and the result entitles the organization to use the CE 0413 marking on those products listed below.

#### Organization:

## Consol AB

Krokslättsgatan 7, SE- 431 67 Mölndal, Sweden  
Box 2094, SE-431 02 Mölndal, Sweden

#### Product Category:

- Contact lens care products
- Comfort solutions

For further identification of the products covered, see the MDD product list/product schedule.

December 12, 2016

Signed date



Mats Premfros, Certification Authority MDD  
Intertek Semko AB, Kista, Sweden