

# EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

**No.****CE 556174**

Issued To:

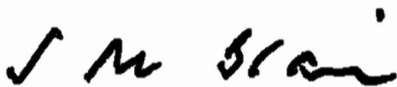
**CorneaGen Inc.  
101 North Chestnut Street, Suite 303  
Winston-Salem  
North Carolina  
27101  
USA**

In respect of:

**The manufacture of corneal endothelium delivery instruments**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):



Stewart Brain, Head of Compliance & Risk -  
Medical Devices

First Issued: **2010-04-21**Date: **2018-08-15**Expiry Date: **2020-04-20**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

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## List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

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**USA**

<b>Subcontractor:</b>	<b>Service(s) supplied</b>
Cathtek Inc 3825 Reidsville Road Winston-Salem North Carolina 27101 USA	<b>Manufacture</b>
Emergo Europe Prinsessegracht 20 2514 AP The Hague The Netherlands	<b>EU Representative</b>
Sterigenics 1148 Porter Avenue Haw River North Carolina 27258 USA	<b>Gamma Sterilization</b>

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# EC Certificate - Production Quality Assurance Certificate History

Certificate No: **CE 556174**  
 Date: **2018-08-15**  
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Date	Reference Number	Action
21 April 2010	7453974	First issue.
09 April 2015	8254909	Certificate renewal. Change of company name to Ocular Systems LLC and removal of Piedmont Triad Research Park from address. Change of subcontractor name from SteriPro Consulting to SteriPro Laboratories.
23 December 2016	8660044	Company name change from Ocular Systems LLC to SightLife Surgical Inc.
10 April 2017	8712730	Change of EU Representative subcontractor address.
Current	9629888	Company name change from Sightlife Surgical, Inc. to Corneagen Inc. Removal of subcontractor SteriPro Laboratories .

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