

Certificate

Full Quality Assurance



No. CE 553140

Issued to:

NeoVista, Inc
47865 Fremont Blvd
Fremont
California
94538
USA

In respect of:

The design, development and manufacture of the NeoVista VIDION® Anti-NeoVascular Therapy (ANV®) System including non-sterile Reusable Delivery Module and sterile Disposable Delivery Assembly and Applicator

on the basis of our examination under the requirements of Council Directive 93/42/EEC, Annex II, Section 3.2.

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

A handwritten signature in black ink, appearing to be 'D. Ford', written over a horizontal line.

David Ford, Director, Healthcare and Testing Services

First Issued: **29 Jul 2009**

Date: **29 Jul 2009**

Expiration Date: **28 Jul 2014**

Page: 1 of 1

Conditions of Approval

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive. 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.

raising standards worldwide™



List of Significant Subcontractors

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

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Subcontractor	Service(s) supplied
AR-MED Ltd Runnymede Malthouse Egham United Kingdom TW20 9BD	EU Representative
Hantel Technologies, Inc. 721 Sandoval Way Hayward California USA 94544	Design Manufacture
nuclitec GmbH Gieselweg 1 38110 Braunschweig Germany	Design Manufacture
Centurion Sterilization Services A Division Of Tri-State Hospital Supply 301 Catrell Drive, Howell Michigan 48843 USA	Sterilization

History of Quality Assurance Certificate

Certificate No: CE 553140
Issue Date: 29 July 2009
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Date	Reference Number	Action
29 July 2009	7432438	First issue