



# NSAI

## Quality System Approval Certificate

Medical Devices Directive 93/42/EEC

*The National Standards Authority of Ireland as a duly designated  
Notified Body, (identification number 0050), for the purposes of the European Communities  
(Medical Devices) Regulations (S.I. No. 252 of 1994)*

*APPROVES THE QUALITY SYSTEM APPLIED BY*

### Lumitex Medical Devices, Inc.

**8443 Dow Circle  
Strongsville  
Ohio 44136  
USA**

*to the Product Family*

### Surgical instrument fibreoptic light (LightMat)

**GMDN Code: 48036**

*on the basis of examination under the requirements of Directive 93/42/EEC on Medical Devices, Annex  
II, excluding (4)*

*The use of the NSAI Notified Body identification number 0050 in conjunction with CE Marking of  
Conformance for this product family is hereby authorised.*

<b>Registration Number:</b>	<b>252.914</b>
<b>Original Approval:</b>	<b>14 November 2013</b>
<b>Last Amended on:</b>	<b>25 September 2018</b>
<b>Remains valid until:</b>	<b>13 November 2021</b>

**Signed:**

Approved by:  
Geraldine Larkin  
Chief Executive Officer, NSAI

Approved by:  
Susan Murphy  
European Medical Device Operations Manager

**This certificate remains valid on condition that the Approved Quality System is maintained in an adequate and efficacious manner.**  
Details of the operational locations included within the scope of this approval can be obtained from NSAI

In the case of a Class III device, this certificate must be supported by a valid design examination certificate  
**National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland.**