



# 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*

### Crospon Ltd

**Galway Business Park  
Dangan  
Co. Galway  
Ireland**

*to the Product Family*

### **Esophageal Balloon Catheter and Inflator (EndoFLIP® Functional Lumen Imaging Probe)**

*on the basis of examination under the requirements of Annex II, Section 3.2 of Directive 93/42/EEC.  
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>252.784</b>
<b>Original Approval:</b>	<b>09 January 2009</b>
<b>Last Amended on:</b>	<b>09 January 2009</b>
<b>Remains valid until:</b>	<b>08 January 2012</b>

**Signed:**

Patrick Breen – Director of NSAI Client Services

**This certificate remains valid on condition that the Approved Quality System is maintained in an adequate and efficacious manner .  
Details of the current product range and operational locations included within the scope of this approval can be obtained from NSAI  
National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland.**