

# CROSPON

Crospon Ltd., Galway Business Park, Dangan, Galway, Ireland.  
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## Declaration of Conformity

We, Crospon Ltd., of Galway Business Park, Dangan, Galway Ireland declare under our sole responsibility that the following products:

EF-100	EndoFLIP System
EF-400	EndoFLIP System Cart
EF-500	EndoFLIP Power Supply
LK-100	EndoFLIP System Localisation Kit (UK/IRE)
LK-101	EndoFLIP System Localisation Kit (EU)
LK-102	EndoFLIP System Localisation Kit (Scandanavia)
LK-103	EndoFLIP System Localisation Kit (US)
LK-104	EndoFLIP System Localisation Kit (South America)
LK-105	EndoFLIP System Localisation Kit (Australia)
LK-106	EndoFLIP System Localisation Kit (Japan)
EF-325	Catheter – 25mm (Box of 5)
EF-335	Catheter – 35mm (Box of 5)
EF-350	Catheter – 50mm (Box of 5)
EF-325N	Catheter, Nasal – 25mm (Box of 5)

to which this declaration relates, and which bear the CE Marking, are in conformity with the applicable requirements of EC Directive 93/42/EEC as amended by 2007/47/EC concerning medical devices.

This declaration is supported by EC Quality System Approval Registration No. MD19.4354 and Quality System Approval Certificate No. 252.784 both issued by the National Standards Authority of Ireland, Notified Body No. 0050 in accordance with Annex II of the above directive.



Signature of Authorised Person

CEO

Title

JOHN O'DEA

Print Name

8th Mar 2010

Date