Umbra Medical Products

Declaration of Conformity

PRODUCT IDENTIFICATION	
Product name	**Model/number**
InMEDFLATOR™ Inflation Device	12107 ADD Y KIT # 33330– CLICK 13108 ADD Y KIT # 33336 – CLICK

MANUFACTURER		
Name of company	**Address**	**Representative**
Umbra Medical Products	8930 Roan Lane East Inverness, Florida USA 34450-1807	Larry Junker

AUTHORIZED REPRESENTATIVE		
Name of company	**Address**	**Telephone/email**
Emergo Europe	Molenstraat 15 2513 BH The Hague, Netherlands	+31.70.345.8570 - phone +31.70.346.7299 - fax Europe@emergogroup.com

REGISTRATION INFORMATION	
Notified Body and ID #	**CE certificate number**
Intertek CE 0473	1167-02 CE, 1167-02 DE

CONFORMITY ASSESSMENT		
Device classification	**Route to compliance**	**Standards applied**
Class I	Annex II of MDD 93/42/EEC Council Directive, Section 4	

Umbra Medical Products, Inc. declares that the above mentioned products meet the provision of the Council Directive 93/42/EEC for Medical Devices and Directive 93/42/EEC as transposed in the national laws of the Member States.

COMPANY REPRESENTATIVE: Larry G. Junker

TITLE: President **SIGNATURE:** *Larry G. Junker*

DATE: 23/05/2017

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