CERTIFICATE OF EUROPEAN UNION AUTHORISED REPRESENTATIVE

CE

This is to certify that Dechoker LLC.

has duly submitted the following relevant product types with the UK Competent Authority (MHRA Ref No. CA015553) through its Appointed Representative in accordance with *Article 14* of the Council Directive 93/42/EEC (revised by 2007/47/EC) concerning medical devices

(The "Medical Devices Directive") (UK Medical Devices Regulation 1994: Regulation 14)

********** Scope of Supply **********

Antichoking Products (IDCH01/IDCH02) Class I non-sterile, single use (MHRA Code Z999)

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In accordance by self-declaration with *Article 11* and *Annex VII* for Class I devices may apply the CE Mark

********** Appointment **********

We certify that M. Devices Group was appointed as the Authorised Representative on 18 May 2016

Signature
Authorised Representative

Date 1 July 2016

Certificate No. MDG-1034-AR Valid to 17 May 2017

Healthcare Education Centre,
Portland Street, Southport, PR8
1HU, England