

**EC DECLARATION OF CONFORMITY**

Medtrade Products Ltd of:

Electra House  
Crewe Business Park  
Crewe  
Cheshire  
CW1 6GL  
UK

Declares the class III medical device, with the GMDN code 46922 Chitosan Haemostatic Agent, described hereafter:

**Celox RAPID Z-Fold Gauze**

For Treatment of Life-Threatening Emergency Bleeding Trained Emergency Responders

In a sheet 7.6cm by 1.5m, with part number FG0XX39YY1

Where XX is a two-digit number representing a customer, and YY is a two-digit number representing the size, language and background colour of the pouch.

Are in conformity with the essential requirements and provisions of Council Directive 93/42/EEC (Medical Device Directive) amended by Directive 2007/47/EC.

And are subject to the procedure in Annex II of Council Directive 93/42/EEC, under the supervision of Notified Body Number 2797, British Standards Institute (BSI), Say Building, John M Keynesplein 9, 1066 EP, Amsterdam, The Netherlands.

Signed:   
Sue McLoughlin  
Regulatory Director

Date: 12<sup>th</sup> November 2019.

Issued in Crewe, Cheshire, U.K.

**Change History:**

1. Original declaration signed by J Ranfield 4<sup>th</sup> Dec 12
2. Re-issue for signature by current Head of Regulatory 16<sup>th</sup> Jan 17
3. Revised following approval of additional sterilisation facility; addition of GMDN code to DoC; and signature by I Walker 6<sup>th</sup> Feb 18.
4. Revised for change to BSI number and Netherlands address, and signature by S. McLoughlin.