symmetry surgical Bovie

To whom it may concern

Ogni Dispositivo Medico, di Protezione e IVD deve essere registrato presso MHRA prima dell'immissione sul mercato in UK. Fabbricanti con sede fuori UK devono nominare un loro Rappresentante avente sede in UK (UKRP).

All medical devices, IVDs, PPE must be registered with the MHRA before they can be placed on the market in Great Britain. Non-UK manufacturer are required to appoint a single UK Responsible Person (UKRP) within the UK.

La sottoscritta società/The undersigned company

Bovie Medical Corporation

con sede in/based in

5115 Ulmerton Road, Clearwater Florida 33760

Dichiara/Declares

Di aver nominato la seguente società come UKRP affinchè operi in UK per suo nome e conto per i prodotti sotto elencati con etichetta aggiornata, firmando la specifica Lettera di Designazione prevista da MHRA. I prodotti sono registrati nel Data Base Pubblico di Registrazione dei DM.

To have commissioned the company shown here to act as UKRP in his name and on behalf regarding the items **listed below;** that Letter of Designation that includes standard text provided by the MHRA was signed. All devices are registered with Public Access Database for Medical Device Registration.

Emergo Consulting (UK) Limited
c/o Cr360 – UL International
Compass House, Vision Park Histon
Cambridge CB24 9BZ
United Kingdom

Elenco dei prodotti (Codice Gima + descrizione)
List of products (Gima code + description)

Codice Gima/Gima code	Nome/Name
DEL1	Change-A-Tip Deluxe High
H101	Replacement Tip Fine (10/bx)
H103	Replacement Tip Loop (10/bx)
H105	Replacement Tip 2" Flex (10/bx)

3034 Owen Drive, Antioch, TN 37013
Office 800.251.3000
Fax 615.964.5567
symmetrysurgical.com

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H109	Replacement Tip 5" Loop (10/bx)
H121	Replacement Tip Vasectomy (10/bx)
HIT1	Power Handle High-Temp
UV59	U.V. Woods Light
UV59B	Lamp Flourscent Black

Date, place

May 24, 2022

Antioch, Tennessee 37013

Stamp and signature

Christopher Smith

Senior Director of Regulatory Affairs

symmetry surgical

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