

## EC DECLARATION OF CONFORMITY

Issued according to Annex II to the Directive 93/42/EEC on Medical Devices as amended by the Directive 2007/47/EC

Manufacturer:

Stevenage

**BTL Industries Limited** 

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161 Cleveland Way

United Kingdom

Authorised Representative:

**BTL ITALIA S.r.l.** Via San Leonardo 120 84131, Salerno Italy

The **BTL Industries Limited** issues this Declaration of Conformity under its sole responsibility and herewith declares that the products

Product Description: Product Name: Product Models: **Combined Therapy Device** 

BTL-5000 Series / BTL-5000 Combi

BTL-5820S (BTL-5820S Combi) BTL-5825S (BTL-5825S Combi) BTL-5820SL (BTL-5820SL Combi) BTL-5825SL (BTL-5825SL Combi) BTL-5816SLM (BTL-5816SLM Combi) BTL-5818SLM (BTL-5818SLM Combi) BTL-5825L (BTL-5825L Combi) BTL-5825M2 (BTL-5825M2 Combi) BTL-5800SL (BTL-5800SL Combi) BTL-5800LM2 (BTL-5800LM2 Combi)

Electrotherapy Device BTL-5000 Series / BTL-5000 Puls

BTL-5620 Puls BTL-5625 Puls BTL-5640 Puls BTL-5645 Puls

Ultrasound Therapy Device BTL-5000 Series / BTL-5000 Sono BTL-5710 Sono BTL-5720 Sono

Laser Therapy Device BTL-5000 Series / BTL-5000 Laser BTL-5110 Laser



Product Description: Product Name: Product Models:

Product Description: Product Name: Product Models:

Product Description: Product Name: Product Models:



Product Description: Product Name: Product Models:

Magnetotherapy Device BTL-5000 Series / BTL-5000 Magnet BTL-5920 Magnet BTL-5940 Magnet

Risk Classification:

Class IIb According to Annex IX of MDD

are in conformity with requirements of Annex I to the Directive 93/42/EEC on Medical Devices as amended by the Directive 2007/47/EC

and bear the CE mark:

Notified Body:

EC Certificate No.:

C E<sub>2460</sub>

DNV Product Assurance AS 10416-2017-CE-CZS-NA-PS

Date of Issue: May 24, 2021 Place of Issue: **Stevenage** 

Signature on behalf of BTL Industries Ltd.



Page 2 of 2