kinetik WELLBEING

No: TD3 DoC

EC Declaration of Conformity

Kinetik Medical Devices Ltd, herewith declare that the products

TENS Pain Reliever: TD3

meets the provisions of Directive 93/42/EEC which apply to them.

The medical devices has been assigned to class IIa according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE 0197

The product concerned has been designed and manufactured under a quality management system according to Annex II of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Certificate No.: HD 60147729 0001
Issue date: 2018-08-24
Expiry date: 2023-01-27

following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

This Declaration of conformity is valid in connection with the release document fo the respective batch of produced devices.

Place / Date: Surrey / November 3, 2020

Signed: JRGrover
Name: James Grover
Position: Sales Director