EC Declaration of Conformity

Kinetik Medical Devices Ltd, herewith declare that the products

Pulse Oximeter – JPD500E

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

The medical devices has been assigned to class IIb rule 10 according to Annex IX without section 4 of the Directive 93/42/EEC. It bears the mark

CE 0482

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

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg - Germany
Issue date: 2019-12-20
Expiry date: 2024-05-27

following the procedure relating to the EC Declaration of Conformity set out in Annex IX without section 4 of the 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