No: NB-222C DoC

EC Declaration of Conformity

Kinetik Medical Devices Ltd, herewith declare that the products

Compressor Nebuliser – NB-222C

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

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

CE 0123

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 SÜD Product Service GmbH
Ridlerstr, 65, 80339, München, Germany
Issue date: 2019-11-25
Expiry date: 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex IV 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