No: JH-125 DoC

EC Declaration of Conformity

Kinetik Medical Devices Ltd, herewith declare that the products

Product: Hearing Amplifier
Model: JH-125
DMDNS/UMDNS code: 17253

meets the provisions of the Council Directive 93/42/EEC concerning Medical Device, as amended by 2007/47/EC.

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

CE 1639

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

SGS Belgium NV
SGS House Noorderlaan 87 2030 Antwerp Belgium
Issue date: 2020-06-02

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