kinetik WELLBEING

No: AG-607 DoC

EC Declaration of Conformity

Kinetik Medical Devices Ltd, herewith declare that the products

Blood Glucose Monitor
EDMS-Code: 21.07.10.01
Model: AG-607

meets the provisions of Directive IVDD 98/79/EC Annex IV which apply to them.

The medical device has been assigned to List B according to IVDD 98/79/EC Annex II. It bears the mark

CE 0197

The product concerned has been designed and manufactured under a quality management system according to IVDD 98/79/EC Annex IV.

Compliance of the designated product with the Directive IVDD 98/79/EC Annex IV has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Issue date: 2019-08-08
Expiry date: 2024-05-27

following the procedure relating to the EC Declaration of Conformity set out in IVDD 98/79/EC Annex IV.

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