



**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 for the respective batch of produced devices.

Place / Date:

Surrey / November 3, 2020

Signed:  
Name:  
Position:

*JRGrover*  
James Grover  
Sales Director