

Declaration of Conformity

ACON Laboratories, Incorporated
10125 Mesa Rim Road
San Diego, CA 92121, USA

**We, the manufacturer, declare under our sole responsibility that
the *in vitro* diagnostic device:**

On Call® Plus II Blood Glucose Monitoring System
(G113-10C, G113-11C, G113-12C)
On Call® Plus II Blood Glucose Test Strips (G133-10C, G133-11C)
On Call® Plus II Glucose Control Solution (G123-12C)

classified as *Annex II List B* of the directive 98/79/EC,


**meets all the provisions of the directive 98/79/EC on *in vitro*
diagnostic medical devices which apply to it**

**The declaration according to Annex IV of the Directive
is based on approval by the notified body
TÜV SÜD Product Service GmbH,
Ridlerstraße 65,
80339 MÜNCHEN, Germany,
notified under No. 0123 to the EC Commission.**

This declaration is valid until expiration of EC Certificate
No. V1 080997 0017 Rev. 01
Expiration Date: 2022-09-12

Authorized Representative:
Medical Device Safety Service GmbH
Schiffgraben 41
30175 Hannover, Germany

Signed this 13th day of June, 2019
in San Diego, CA, USA



Qi Yi Xie, MD, MPH
Senior Staff, Regulatory Affairs & Clinical Affairs
ACON Laboratories, Inc.

