





Product Service

EC Certificate

Full Quality Assurance System
Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6)
(List A and B and devices for self-testing)

No. V1 104507 0003 Rev. 06

Manufacturer: ACON Laboratories, Inc.

5850 Oberlin Drive, #340 San Diego CA 92121

USA

Product Category(ies): Blood glucose measuring systems for self testing

and self-testing devices for clinical chemistry, hematology and pregnancy and ovulation

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device families in accordance with IVDD Annex IV. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of List A devices an additional Annex IV (4) certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:V1 104507 0003 Rev. 06

Report no.: SH22743EXT01

 Valid from:
 2022-05-04

 Valid until:
 2025-05-26

Date, 2022-05-04

Christoph Dicks

Head of Certification/Notified Body



Full Quality Assurance System Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6) (List A and B and devices for self-testing)

No. V1 104507 0003 Rev. 06

On Call Plus Blood Glucose Monitoring System, Model(s):

On Call Plus Blood Glucose Test Strips,

On Call EZ II Blood Glucose Monitoring System.

On Call Advanced Blood Glucose Monitoring System,

On Call Advanced Blood Glucose Test Strips.

On Call Chosen Blood Glucose Test Strips,

On Call Vivid Blood Glucose Monitoring System (OGM-101),

On Call Vivid Blood Glucose Test Strips (OGS-101),

On Call Sharp Blood Glucose Monitoring System (OGM-

121),

On Call Sharp Blood Glucose Test Strips (OGS-121)

On Call Plus II Blood Glucose Monitoring System (OGM-

On Call Plus II Blood Glucose Test Strips (OGS-171),

On Call Extra Blood Glucose Monitoring System (OGM-191).

On Call Extra Blood Glucose Test Strips (OGS-191),

On Call GK Dual Blood Glucose & Ketone Monitoring

System (OGM-161),

On Call Blood Ketone Test Strips (OGS-161),

Urinalysis Reagent Strips (Urine),

UTI Urinary Tract Infection Test Strips.

Cholesterol Monitoring System (CCM-111),

CHOL Total Cholesterol Test Devices (CCS-111).

TRIG Triglycerides Test Devices (CCS-112),

HDL High Density Lipoprotein Test Devices (CCS-113),

3-1 Lipid Panel Test Devices (CCS-114),

Cholesterol CTRL Control Devices,

Cholesterol Monitoring System (CCM-101),

CHOL Total Cholesterol Test Strips (CCS-101).

PT/INR Monitoring System (CCM-151),

PT/INR Test Strips (CCS-151),

Hemoglobin Testing System (CCM-141),

Hemoglobin Test Strips (CCS-141),

hCG Pregnancy Rapid Test Cassette (Urine),

Pregnancy Rapid Test Midstream,

On Call Extra Mobile Blood Glucose Monitoring System

(OGM-281),

On Call Sure Blood Glucose Monitoring System (OGM-211),

On Call Sure Sync Blood Glucose Monitoring System (OGM-

212),

On Call Sure Blood Glucose Test Strips (OGS-211),

GIMA Blood Glucose Monitoring System,

GIMA Bluetooth Blood Glucose Monitoring System.

GIMA Blood Glucose Test Strips,

On Call GU Dual Blood Glucose & Uric Acid Monitoring





Full Quality Assurance System
Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6)
(List A and B and devices for self-testing)

No. V1 104507 0003 Rev. 06

System (OGM-201),

On Call Blood Uric Acid Test Strips (OGS-201),

LH Ovulation Rapid Test Cassette (Urine),

Ovulation Rapid Test Midstream,

Ovulation & Pregnancy Test Combo Pack,

On Call Extra Voice Blood Glucose Monitoring System (OGM-291).

Early Detection Pregnancy Test,

Digital Pregnancy Test,

Go-Keto Blood Glucose & Ketone Monitoring System (OGM-161).

Go-Keto Blood Ketone Test Strips (OGS-161),

Go-Keto Blood Glucose Test Strips,

On Call Extra GM Blood Glucose Monitoring System(OGM-191)

On Call Extra GM Blood Glucose Test Strips (OGS-191),

On Call Plus GM Blood Glucose Monitoring System,

On Call Plus GM Blood Glucose Test Strips,

Go-Keto Urinalysis Reagent Strips

Facility(ies): ACON Laboratories, Inc.

5850 Oberlin Drive, #340, San Diego CA 92121, USA

ACON Laboratories, Inc.

10125 Mesa Rim Road, San Diego CA 92121, USA

AZURE Institute, Inc.

10125 Mesa Rim Road, San Diego CA 92121, USA

Acon Laboratories Inc.

Guerrero Negro 9942 Parque Industrial Pacifico IV, 22644 Tijuana

B.C. CP, MEXICO





Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 104507 0002 Rev. 01

Manufacturer: ACON Laboratories, Inc.

5850 Oberlin Drive, #340 San Diego CA 92121

USA

Product Category(ies): Lancets, Safety Lancets

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: SH1974310

Valid from: 2019-10-24 **Valid until:** 2023-09-06

Date, 2019-10-24

Stefan Preiß
Head of Certification/Notified Body

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Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 104507 0002 Rev. 01

Facility(ies): ACON Laboratories, Inc.

5850 Oberlin Drive, #340, San Diego CA 92121, USA

ACON Laboratories, Inc.

10125 Mesa Rim Road, San Diego CA 92121, USA

AZURE Institute, Inc.

10125 Mesa Rim Road, San Diego CA 92121, USA

Declaration of Conformity

ACON Laboratories, Incorporated 5850 Oberlin Drive #340 San Diego, CA 92121 USA

We, the manufacturer, declare under our sole responsibility that the medical device:

Mission® Lancets (C121-3041)
On Call® Lancets (G124-10A)
Insight® Lancets (C121-3045)
Swiss Point of Care Lancets (G124-90AA)

of class IIA according to Annex IX rule 6 of the directive 93/42/EEC,

meets all the provisions of the directive 93/42/EEC as amended by directive 2007/47/EC concerning medical devices which apply to it.

This declaration is according to Annex II of the Directive and thus 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. G1 104507 0002 Rev. 01
Expiration Date: 2023-09-06

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

Signed this 17 day of August, 2021 in San Diego, CA USA

Qiyi Xie, MD, MPH

Senior Staff, Regulatory Affairs & Clinical Affairs ACON Laboratories, Inc.



Declaration of Conformity

ACON Laboratories, Incorporated 5850 Oberlin Drive #340 San Diego, CA 92121 USA

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

Device Name	REF Number
On Call® Plus Blood Glucose	G113-111
Monitoring System	
On Call® Plus Blood Glucose Meter	G113-211, G113-214
On Call® Plus Blood Glucose Test	G133-111, G133-112, G133-
Strips	114, G133-115, G133-117,
	G133-118, G133-119, G133-
	211
On Call® Plus Glucose Control	G123-311
Solution	

classified for *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 104507 0003 Rev. 06
Expiration Date: 2025-05-26

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



Signed this 25 day of May, 2022 in San Diego, CA USA

Qiyi Xie, MD, MPH

Senior Staff, Regulatory Affairs & Clinical Affairs ACON Laboratories, Inc.

Letter of Declaration

To whom it may concern:

We *Acon Laboratories,Inc.*, who is the legal manufacturer of Blood Glucose Monitoring System (Including Glucose Meter, Glucose test strip, Control Solution, Lancet and lancing device etc, to test the glucose level of human blood),have registered office at 10125 Mesa Rim Road, San Diego, CA 92121 USA, here to declare that:

- On Call® Plus Strips correspond with On Call® Plus Blood Glucose Monitoring System.
- We currently have in stock the tender required quantity of Meters, Strips and Lancets (1000/50000/50000).

This clarification letter will only be used for product registration, tender submission, sales and marketing of *On Call® Plus* Blood Glucose Monitoring System in **Moldova** it should not be used for any other business or non-business purposes.

Sincerely yours,

Eddie.SA

International Sales Warketing Sales Manager

Diabetes Care

Acon Laboratories,Inc.







Certificate

No. Q5 104507 0001 Rev. 03

Holder of Certificate: ACON Laboratories, Inc.

5850 Oberlin Drive, #340 San Diego CA 92121 USA

Certification Mark:



Scope of Certificate: Design and Development, Manufacture and distribution of In Vitro Diagnostic Test Kits and Reagents for the

Determination of Infectious Diseases, Clinical
Chemistry, Drugs of Abuse, Tumor/Cardiac Marker,
Fertility/Pregnancy and Blood Glucose Monitoring

System, Lancing Devices and Lancets

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5 104507 0001 Rev. 03

Report No.: SH22743A01

 Valid from:
 2022-09-15

 Valid until:
 2025-09-06

Date. 2022-09-15 Christoph Dicks

Head of Certification/Notified Body





Certificate

No. Q5 104507 0001 Rev. 03

Applied Standard(s): EN ISO 13485:2016

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016) DIN EN ISO 13485:2016

Facility(ies): ACON Laboratories, Inc.

5850 Oberlin Drive, #340, San Diego CA 92121, USA

Address holder for registration only

ACON Laboratories, Inc.

10125 Mesa Rim Road, San Diego CA 92121, USA

Manufacture and distribution of

In Vitro Diagnostic Test Kits and Reagents for the Determination of Infectious Diseases, Clinical Chemistry, Drugs of Abuse, Tumor/Cardiac Marker, Fertility/Pregnancy and Blood Glucose Monitoring System, Lancing Devices and Lancets

ACON Laboratories, Inc.

6865 Flanders Dr., Suite B, San Diego CA 92121, USA

Storage of

In Vitro Diagnostic Test Kits and Reagents for the Determination of Infectious Diseases, Clinical Chemistry, Drugs of Abuse, Tumor/Cardiac Marker, Fertility/Pregnancy and Blood Glucose Monitoring System, Lancing Devices and Lancets

AZURE Institute, Inc.

10125 Mesa Rim Road, San Diego CA 92121, USA

Design and Development of

In Vitro Diagnostic Test Kits and Reagents for the Determination of Infectious Diseases, Clinical Chemistry, Drugs of Abuse, Tumor/Cardiac Marker, Fertility/Pregnancy and Blood Glucose Monitoring System, Lancing Devices and Lancets

Acon Laboratories Inc.

Guerrero Negro 9942 Parque Industrial Pacifico IV, 22644 Tijuana B.C. CP, MEXICO

Manufacture of

blood glucose test strips, antigen rapid test and IgG/IgM antibody rapid test for infectious disease.

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Specification

Feature	Specification	MS	
Technology	Biosensor/Electrochemical, Glucose oxidase (GOD)	The state of the s	
Result Calibration	Plasma-equivalent		
Test Time	10 seconds		
Sample Size	0.5 μL		
Sample Type	Fresh capillary whole blood		
Hematocrit Range	25 - 60%		
Glucose Test Range	20 - 600 mg/dL (1.1 - 33.3 mmol/L)		
Memory Storage	300 results with date and time		
Test Averaging	7, 14, 30-day averages		
Data Transfer	USB		
Control Solution	3 levels		
Audio Feature	Optional beep for sample detection, error messages		
Automatic Shutoff	2 minutes after last action		
Battery	One (1) CR 2032 3.0V coin cell battery		
Battery Life	1,000 measurements		
Operating Conditions	41 - 113 °F (5 - 45°C) and 10 - 90% relative humidity		
Strip Storage Temperature	2-35°C		
Expiration Date	24 months (6 months after first opening)		

Catalog

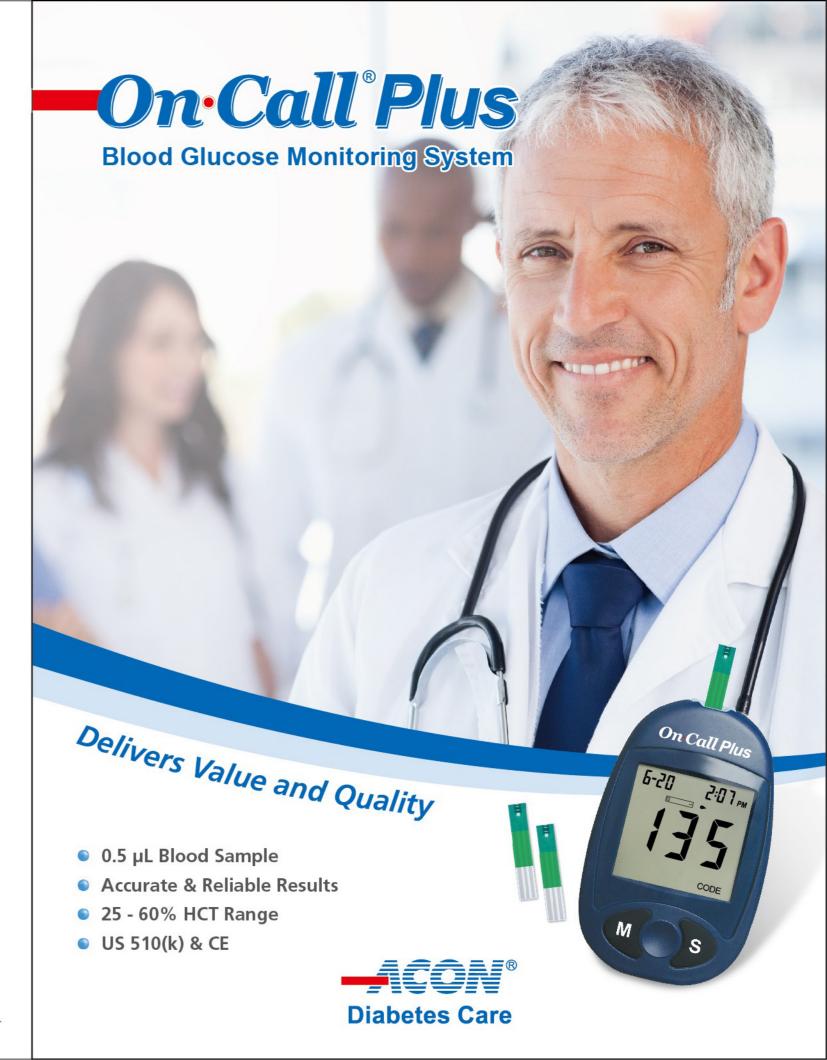
	3.00						
Product Name	Catalog No.			Conte	ents		
On Call® Plus Blood Glucose Monitoring System	G113-111 √ †	1 Meter 1 Manual 10 Lancets	10 Test St 1 Carrying 1 Code Ch	Case		lution 1 erence Guide (for testing on forear	1 Lancing Device 1 Warranty Card m and palm)
On Call® Plus	G113-211 √ †	1 Meter 1 Manual		rol Solution 1 ranty Card		arrying Case uick Reference Guide)
Blood Glucose Meter	G113-214 V	1 Meter 1 Manual 10 Lancets	1 Carry	ing Device ying Case ranty Card	10	ontrol Solution 1 Juick Reference Guide lear Cap (for testing o	e on forearm and palm)
	0400 444 11	50 Test Strips (50/vial) 100 Test Strips (25/vial) 10 Test Strips (10/vial) 25 Test Strips (Individually Foil Wrapped)		1 C	ode Chip	1 Package Insert	
	G133-111 √ †			1 C	ode Chip	1 Package Insert	
	G133-112 √			1 C	ode Chip	1 Package Insert	
On Call® Plus	G133-114 V			1 C	ode Chip	1 Package Insert	
Blood Glucose Test Strips	G133-115 √			10	ode Chip	1 Package Insert	
	G133-117 √			10	ode Chip	1 Package Insert	
	G133-118 V 25 Test Strips (25/vial)		1 C	ode Chip	1 Package Insert		
On Call® Plus Blood Glucose Test Strips and Lancets	G133-211 √	50 Test Strips (25/vial)	50 Lancets (2	25/bag)	1 Code Chip	1 Package Insert
On Call® Plus Blood Glucose Control Solution	G123-311 à	1 Control Solut	tion 0 1	Control Solution	on 1 1	Control Solution 2	1 Package Insert
On Call® Lancets	G124-10A à	100 Lancets (2	5/bag)				
On Call® Lancing Device	G124-11AV	1 Lancing Devi	ce		1 Package In	sert	
On Call® Diabetes Management Software Kit	G124-13A†	1 USB Data Tra	nsfer Cable		1 Installation	n Disk	

v CE Marked for sale in the European Community (6 0123



† US 510(k) Cleared and CLIA Waived

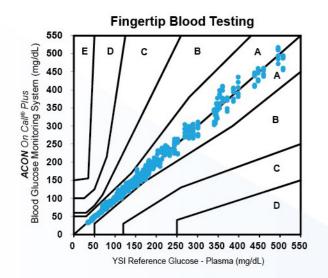




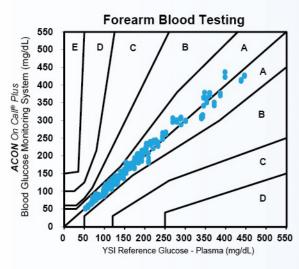


Accurate and Reliable

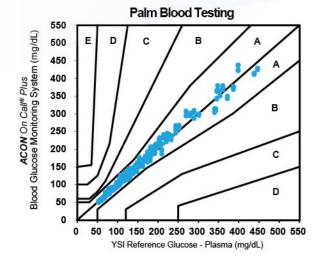
Extensive clinical studies proved the accuracy of *On Call® Plus* Blood Glucose Monitoring System with fresh capillary blood samples, which can comply with EN ISO 15197: 2015.



Clinical Trial -	onsensus Error Grid Ana Fingertip Capillary Bloo lus Blood Glucose Monit	d, by Technican	
System Accuracy Re	sults for Glucose Conce	ntration ≥ 100 mg/dL	
Within ±5%	Within ±10%	Within ±15%	
290 / 462 (62.8%)	432 / 462 (93.5%)	462 / 462 (100.0%)	
System Accuracy Results for Glucose Concentration <100 mg/dL			
Within ±5 mg/dL	Within ±10 mg/dL	Within ±15 mg/dL	
145 / 198 (73.2%)	193 / 198 (97.5%)	198 / 198 (100.0%)	
System Accuracy Results for both Glucose Concentration ≥ 100 mg/dL and < 100 mg/dL			
V	Vithin ±15% or ±15 mg/	dL	
658 / 660 (99.7%)			



Clinical Trial	onsensus Error Grid Ana - Forearm Capillary Bloo lus Blood Glucose Monit	d, by Technican	
System Accuracy Res	sults for Glucose Conce	ntration ≥ 100 mg/dL	
Within ± 5%	Within ± 10%	Within ± 15%	
202 / 444 (45.5%)	375 / 444 (84.5%) 440 / 444 (99		
System Accuracy Re	sults for Glucose Conce	ntration <100 mg/dL	
Within ±5 mg/dL	Within ±10 mg/dL	Within ±15 mg/dL	
110 / 168 (65.5%)	154 / 168 (91.7%)	168 / 168 (100.0%)	
	Results for both Gluco 00 mg/dL and < 100 mg		
V	Vithin ±15% or ±15 mg/c	dL	
	608 / 612 (99.3%)		



Consensus Error Grid Analysis Clinical Trial - Palm Capillary Blood, by Technican ACON On Call® Plus Blood Glucose Monitoring System vs. YSI				
System Accuracy Results for Glucose Concentration ≥ 100 mg/dL				
Within ±5%	Within ±10%	Within ±15%		
219 / 444 (49.3%)	395 / 444 (89.0%)	441 / 444 (99.3%)		
System Accuracy Results for Glucose Concentration < 100 mg/dL				
Within ±5 mg/dL	Within ±10 mg/dL	Within ±15 mg/dL		
130 / 168 (77.4%)	166 / 168 (98.8%)	168 / 168 (100.0%)		
System Accuracy Results for both Glucose Concentration ≥ 100 mg/dL and < 100 mg/dL				
Within ±15% or ±15 mg/dL				
609 / 612 (99.5%)				



Key Features



Authority Certificate







CE certificate

USFDA CFG certificate

Health Canada certificate