

# EC Certificate

**Full Quality Assurance System  
Directive 98/79/EC on In Vitro Diagnostic Medical Devices,  
Annex IV excluding (4, 6)**

Registration No.: HL 2135127-1

Manufacturer: VivaChek Biotech (Hangzhou) Co., Ltd.  
Level 2, Block 2, 146  
East Chaofeng Rd., Yuhang Economy Development Zone,  
Hangzhou,  
311100 Zhejiang  
P.R. China

Products: Blood Glucose Monitoring Systems (Blood Glucose Meters, Blood Glucose Test Strips, Blood Glucose Control Solutions); Multi-function Monitoring Systems (including Multi-function Meters; Blood Ketone Test Strips, Blood Ketone Control Solutions, Uric Acid Test Strips, Uric Acid Control Solutions, Blood Glucose Test Strips, and Blood Glucose Control Solutions)

Replaces Approval, Registration No.: HL 60145622 0001

  
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The Notified Body hereby declares that the requirements of Annex IV, excluding sections 4 and 6 of the directive 98/79/EC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex IV, section 5 of the aforementioned directive. For placing on the market of List A devices covered by this certificate an EC design-examination certificate according to Annex IV, section 4 and a verification of manufactured products according to section 6 is required.

Report No.: 244411842-200

Effective date: 2022-03-23

Expiry date: 2025-05-26

Issue date: 2022-03-23



Fuxiu Sheng  
TÜV Rheinland LGA Products GmbH  
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 98/79/EC concerning in vitro diagnostic medical devices with the identification number 0197.

# EC Certificate

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**Annex IV excluding (4, 6)**

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Hangzhou,  
311100 Zhejiang  
P.R. China

The scope of certification includes the following manufacturing sites:

No.	Location
/01	VivaChek Biotech (Hangzhou) Co., Ltd. 1/2/3 F, Building 1, 16 East Zhenxing Rd., Yuhang Economy Development Zone, Hangzhou, 311100 Zhejiang P.R. China



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TÜV Rheinland LGA Products GmbH • 51105 Köln

VivaChek Biotech (Hangzhou) Co., Ltd.  
Level 2, Block 2, 146 East Chaofeng Rd.,  
Yuhang Economy Development Zone,  
Hangzhou,  
311100 Zhejiang  
P.R. China

Contact

Tel. +49 911 655-5225  
Mail: [medical-products@de.tuv.com](mailto:medical-products@de.tuv.com)

Date August 08, 2023

**Application for: QMS**

Certificate No. : HL 2135127-1  
Requirement : Directive 98/79/EC  
Confirmation letter ID : DOC\_2023-08-09\_ HL 2135127-1  
Report no. : 244508348-200

Dear Madame or Sir,

**Update of information to Certificate no. HL 2135127-1, issued on 23.03.2022**

The change notification received on 21.09.2022 related to the information stipulated on the above mentioned certificate was assessed and information confirmed.

We confirm that the change notification is not considered a significant change in design or intended purpose under Regulation (EU) 2017/746 on in vitro diagnostic medical devices (IVDR), Article 110(3).

With this document we would like to confirm the following updated information to the afore mentioned certificate

**Revised Manufacturer address**

Old Manufacturer address: 1/2/3 F, Building 1, 16 East Zhenxing Rd., Yuhang  
Economy Development Zone, Hangzhou,  
311100 Zhejiang, P.R. China

New Manufacturer address: Building 2 and Building 3, No.17 Jianxing Road,  
Taozhu street, Zhuji, 311800, Zhejiang, P.R.  
China

Best regards,



Herbert Zhong  
Certification body

TÜV Rheinland  
LGA Products GmbH

Am Grauen Stein  
51105 Köln  
Germany

Headquarter

Tillystraße 2  
90431 Nuremberg

Phone. +49 911 655 5225  
Fax +49 911 655 5226  
[service@de.tuv.com](mailto:service@de.tuv.com)  
[www.tuv.com/safety](http://www.tuv.com/safety)

Board of Management

Dipl.-Ing.  
Jörg Mähler, Spokesman

Dipl.-Kfm.  
Dr. Jörg Schlösser

Nuremberg HRB 26013  
VAT No.: DE 811835490

Chairman of the  
Supervisory Board

Dr.-Ing. Michael Fübi