

EC Certificate

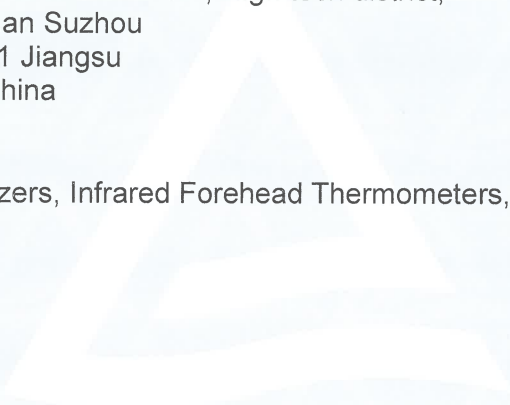


Production Quality Assurance Directive 93/42/EEC on Medical Devices, Annex V

Registration No.: DD 2017980-1

Manufacturer: Vapo Healthcare Co., Ltd.
Southern unit of third floor, building B
No. 99 Yudai West Rd, High tech district,
KunShan Suzhou
215301 Jiangsu
P.R. China

Products: Nebulizers, Infrared Forehead Thermometers, Electrical Thermometers


TÜVRheinland

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC 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 V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Report No.: 17061663 010

Effective date: 2021-03-02

Expiry date: 2024-05-26

Issue date: 2021-03-02



Dipl.-Ing. W. Hsu
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

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The scope of certification includes the following manufacturing sites:

No.	Location	Product groups manufactured
/01	Vapo Healthcare Co., Ltd. Room 1308, Industry Technology Institution, 1699 Zuchongzhi South BLVD, Kunshan, 215300 Jiangsu P.R. China	Nebulizers, Infrared Forehead Thermometers, Electrical Thermometers

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