TÜVRheinland®

EC Certificate

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

Registration No.: DD 60147567 0001

Report No.: 16803978 008

Manufacturer:
SHENYANG AERTI TECH CO., LTD.
No.30 Huahai Road, Shenyang Economic &
Technological Development Area
Shenyang City
110141 Liaoning
P.R. China

Products: Medical Oxygen Concentrators

Replaces Approval, Registration No.: DD 60100205 0001

Expiry Date: 2024-05-26

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.

Effective Date: 2020-03-31

Date: 2020-03-31

Notified Body

Wenxiang Zhang

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.