EC Certificate

TÜVRheinland®

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

Registration No.: HD 1497647-1

Manufacturer: VELDER
G. Rondio, J. Grządko Spółka Jawna
ul. Siekierkowska 8
00-907 Warszawa
Poland

Products: - Medical gases pipeline installations

Replaces Certificate, Registration No.: HD 60138236 0001

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 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 II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II section 4 is required.

Report No.: 84949829-20

Effective date: 2020-09-16

Expiry date: 2024-05-26

Issue date: 2020-09-16

TÜV Rheinland LGA Products GmbH Zertifizierstelle

Maciej Ściera
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.