# CERTIFICATE

## The Certification Body TÜV Rheinland Italia S.r.l.

certifies, in accordance with the TÜV Rheinland Group procedures, that the Company

## Aries S.r.l.

### Via XXV Luglio, 43

### IT – 41037 Mirandola (MO)

has established and applies a quality management system
for the following scope:

**Design, manufacturing, placing on the market of sterile and non-sterile non-active devices for anaesthesia, emergency and intensive care, sterile and non-sterile non-active devices for injection, infusion, transfusion and dialysis and sterile non-active medical devices with measuring function.**
**Distribution of sterile and non-sterile non-active devices for anaesthesia, emergency and intensive care, sterile and non-sterile non-active devices for injection, infusion, transfusion and dialysis and active medical devices for infusion.**

Through an Audit, Report No. 7992241010FES20, proof has been furnished that the
quality management system fulfils the requirements of the standard

**UNI CEI EN ISO 13485:2021**

Please refer to the Quality Manual for the details about
the exclusions with respect to the requirements of the standard.

Certificate Registration No. **39 05 1492410**

This Certificate is valid from 2024/11/26 to 2027/03/31

The reference date for all the next audits is (day-month): 28/02

**Milan, 2024/11/26.** First Certification: 2024/04/01

The certification responsible: Massimiliano Testi
TÜV Rheinland Italia S.r.l., Via E. Mattei, 3 - I - 20005 Pogliano Milanese (MI)

This certificate does not represent proof that the statutory requirements of the Directives 93/42/EEC, 90/385/EEC, 98/79/EC or Regulations (UE) 2017/745, (UE) 2017/746 have been fulfilled

ACCREDIA
L'ENTE ITALIANO DI ACCREDITAMENTO
MS N° 0083
Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC.
Signatory of EA, IAF and ILAC Mutual Recognition Agreements.
www.tuvitalia.com

TÜVRheinland CERTIFIED
Management System
EN ISO 13485:2016
www.tuv.com
ID 9000035506

TÜVRheinland®