CERTIFICATE

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

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

Medical Products Research S.r.l.

Via Novara, 68

IT - 20025 Legnano (MI)

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

Design, manufacture and placing on the market of active and non-active medical devices for the treatment of respiratory pathologies. Service of active medical devices for the treatment of respiratory pathologies.

Through an Audit, Report No. 7986724010CV12, 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 0711110.**

This Certificate is valid from 2024/01/22 to 2027/01/22.

The reference date for all the next audits is (day-month): 11/11

Milan, 2024/01/22. First Certification: 2012/01/23

The certification responsible: Daniele Ricchi
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.

TÜVRheinland CERTIFIED
Management System
EN ISO 13485:2016
www.tuv.com
ID 9105066786

www.tuvitalia.com

TÜVRheinland®