

CERTIFICATE

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

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

Reactivos para Diagnostico S.L.

C/ Joseph Tura, 9H 11D

ES – 08181 Sentmenat (Barcelona)

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

Production of culture media and reagents for microbiology (in vitro diagnostic medical devices).

Through an Audit, Report No. 7988790010SR29, proof has been furnished that the
quality management system fulfils the requirements of the standard

EN ISO 13485:2016

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 0931111**.

This Certificate is valid from 2024/05/02 to 2027/05/01.

The reference date for all the next audits is (day-month): 13/01

Milan, 2024/05/02. First Certification: 2012/05/02



The certification responsible: Cesare Gentile
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



MS N° 0083

Membro degli Accordi di Mutuo
Riconoscimento EA, IAF e ILAC.
Signatory of EA, IAF and ILAC
Mutual Recognition Agreements.



Management
System
EN ISO 13485:2016
ISO 9001:2015

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