

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/ Josep Tura, 9H

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. 28300442-007, 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 2023-11-24 to 2024-05-01.

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

Milan, 2023-11-24. 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



Management
System
EN ISO
13485:2016

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