

EU Certificate

Quality Management System

REGULATION (EU) 2017/746 on In Vitro Diagnostic Medical Devices Annex IX Chapter I, Section 2 and 3 and Chapter III

Registration No.: HX 2333085-1
Manufacturer: Dia.Metra Srl
Via Pozzuolo 14
06038 Spello (PG)
Italy

EUDAMED Single
Registration No.: IT-MF-000027206

Products: Products of class B:

IMMUNOCHEMISTRY (IMMUNOLOGY)
IVR 0602 - Devices intended to be used for screening,
determination or monitoring of physiological markers for
a specific disease
W01021199 - RHEUMATOID / INFLAMMATORY DISEASE
MARKERS – OTHER


The Notified Body hereby declares that the requirements of Annex IX, Chapter I, Section 2 and 3 of the REGULATION (EU) 2017/746 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.

If class D devices are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4 is required before placing them on the market.

If class B, C or D devices for self-testing or near-patient testing are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 5.1 is required before placing them on the market.

If companion diagnostics are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 5.2 is required before placing them on the market.

Report No.: 84954672-60
Effective date: 2024-06-25
Expiry date: 2029-06-24
Issue date: 2024-06-25



Rafał Byczkowski
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

This certificate can be validated on <https://www.certipedia.com>

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/746 concerning medical devices with the identification number 0197.



Benannt durch/Designated by
Zentralstelle der Länder
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bei Arzneimitteln und
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Precisely Right.

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Annex IX Chapter I, Section 2 and 3 and Chapter III

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IMMUNOCHEMISTRY (IMMUNOLOGY)

IVR 0608 - Devices intended to be used for screening,
determination or monitoring of physiological markers

W01020410 - THYROID STIMULATING HORMONE


W01020501 - FERTILITY FUNCTION HORMONES /
PROTEINS

W01020602 - RENAL METABOLISM ASSAYS

Authorized representative(s): Not applicable.

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2024-06-25

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