



EU-type examination certificate

Number **T12101** revision 1
Project number 4162104
Page 1 of 1

Issued by

NMi Certin B.V.,
designated and notified by the Netherlands to perform tasks with respect to
conformity assessment procedures mentioned in Article 13 of Directive
2014/31/EU, after having established that the measuring instrument meets
the applicable requirements of Directive 2014/31/EU, to:

Manufacturer

Dräger Medical Systems, Inc.
3135 Quarry Road
Telford, PA 18969
United States of America

Measuring instrument

A **Non-automatic weighing instrument**
Type : TN300

Further properties are described in the annexes:

- Description T12101 revision 1;
- Evaluation report ER12101 revision 1;
- Documentation folder T12101-2.

Valid until

12 May 2032

Initially issued

12 May 2022

Remarks

This revision replaces the previous version, including its documentation
folder.

Issuing Authority

NMi Certin B.V., Notified Body number 0122
11 September 2026

Certification Board

NMi Certin B.V.
Thijssseweg 11
2629 JA Delft
The Netherlands
T +31 88 636 2332
certin@nmi.nl
www.nmi.nl

This document is issued under the provision that
no liability is accepted and that the
manufacturer shall indemnify third-party
liability.

The designation of NMi Certin B.V. as Notified
Body can be verified at
<https://webgate.ec.europa.eu/single-market-compliance-space/home>

Reproduction of the complete
document only is permitted.

This document is digitally signed
and sealed. The digital signature
can be verified in the blue ribbon at
the top of the electronic version of
this certificate.

