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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 : Isolette 8000
Isolette 8000 plus
C2000 Isolette Infant Incubator

Further properties are described in the annexes:

- Description T8533 revision 2;
- Documentation folder T8533-1.

Valid until 11 September 2034

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Remark This revision replaces the earlier versions, except for its documentation folder.

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Certification Board

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