

EC CERTIFICATE

Full Quality Assurance System

Certificate No.:
10000371110-PA-NA-DNK Rev 0.0

Project No.:
PRJN-188850-2020-PA-DNK

Valid Until:
26 May 2024

This is to certify that the quality system of:

CODAN Medical ApS

Færgevej 4, 4970 Rødby, Denmark

For design, production and final product inspection/testing of:
SYRINGES

Has been assessed with respect to:

**THE CONFORMITY ASSESSMENT PROCEDURE DESCRIBED IN
ANNEX II EXCLUDING SECTION 4 OF COUNCIL DIRECTIVE
93/42/EEC ON MEDICAL DEVICES, AS AMENDED**

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and date:
Høvik, 18 January 2021

For:
DNV GL PRESAFE AS
Notified Body No.: 2460



Eugenie Winger Husebye

The certificate is digitally verified by blockchain technology. For more info, see www.dnvgl.com/assurance/certificates-in-the-blockchain.html



Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.
NOTIFIED BODY: DNV GL PRESAFE AS, Veritasveien 3, N-1363 Høvik, Norway - Registered Enterprise No: NO 997 067 401 MVA

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Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate history:

Revision	Description	Issue Date
0.0	Transfer of Presafe Denmark A/S (NB 0543) Certificate No. DGM -380 to DNV GL Presafe A/S (NB2460)	19 January 2021

Products covered by this Certificate:

Product Description	Product Name	Class
Syringes	<ul style="list-style-type: none"> • luer CODAN • luer-lock CODAN • luer ONCE 	Im
Syringes	<ul style="list-style-type: none"> • luer CODAN • luer-lock CODAN • Female luer-lock CODAN • Catheter tip CODAN • luer PRIMO • Catheter tip PRIMO • luer ONCE • Catheter tip ONCE • Luer ERSTA 	Is

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

Site Name	Address
CODAN Medical ApS	Færgevej 4, 4970 Rødby, Denmark

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Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

End of Certificate