

This is to certify that the quality system of:

Fast Plast A/S
Vilhelmsborgvej 2
DK-7700 Thisted
Denmark

has been approved in conformity with the requirements of

Annex V, section 3.2 - Production quality assurance

of Council Directive 93/42/EEC concerning medical devices as transposed into Danish law.

The certificate covers the following activities:

Manufacture and final testing of medicine cups in class I with a measuring function

The EC certificate is valid provided that the quality system continues to conform to the above-mentioned scope and provided that the company does not introduce substantial changes to the quality system without the approval of Presafe Denmark A/S. This EC certificate is issued pursuant to the Presafe Denmark A/S terms and conditions for the certification of medical devices and entitles the manufacturer to affix the CE mark. The certificate is based on successful audit of the manufacturer. The manufacturer is subject to periodical audits as per the Directive.



Bent Buus
Authorized person

For Presafe Denmark A/S

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Presafe Denmark A/S
Notified Body, Identification No. 0543
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