

EC Certificate
Directive 93/42/EEC Annex VI
Product Quality Assurance
Medical Devices

Registration No.: ED 60110388 0001

Report No.: 21248408 001

Manufacturer: SENSOPLAST Packmitteltechnik GmbH
Auf dem Höhchen 1-5
56587 Oberhonnefeld-Gierend
Deutschland

Products: (see attachment for products included)

Replaces Approval, Registration No.: ED 60038582 0001

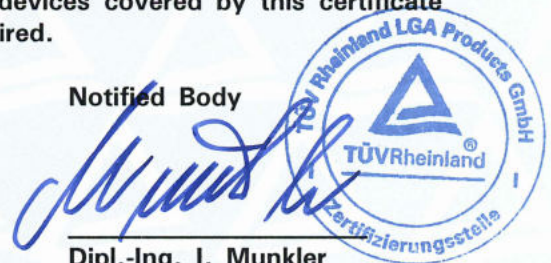
Expiry Date: 2021-05-03

The Notified Body hereby declares that the requirements of Annex VI of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex VI, section 4 of the aforementioned directive. For placing on the market of class IIb devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2016-05-04

Date: 2016-05-04

Notified Body



Dipl.-Ing. I. Munkler

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Doc. 1/1, Rev. 0

**Attachment to
Certificate**

Registration No.: ED 60110388 0001
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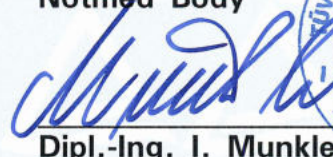
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For the following devices the scope covers only
the aspects of manufacture concerned with conformity
of the products with the metrological requirements:

- Measuring Cups
- Measuring Spoons

Date: 2016-05-04

Notified Body



Dipl.-Ing. I. Munkler

