

The management system of

Neutroplast, Industria de Embalagens Plasticas, S.A.

Zona Industrial, Casal da Espinheira, Lote 10,
2590-057 Sobral de Monte Agraço, Portugal

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

Restricted to the aspects of manufacture concerned with the conformity of the devices with metrological requirements

For the following products

**Dosage Cup and Dosage Spoon
for the administration of oral medicine.**

**Copos doseadores e colheres doseadoras
para administração oral de medicamentos**

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 16 February 2020 until 24 May 2024 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 27 March 2014 and first certified by SGS Belgium NV since 16 February 2020.

Certification is based on reports numbered ES/MAD/ 230843

Authorised by

SGS Belgium NV, Notified Body 1639

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