EC Certificate Production Quality Assurance System: Certificate ES20/87302

SGS

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

SGS House Noorderlaan 87 2030 Antwerp Belgium t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

LPMD5008 - Certificate CE1639 AnnexV_EN rev, 01

Page 1 of 1





This document is issued by the Company subject to its General Conditions of Certification Services, unless otherwise agreed, accessible at www.sgs.com/terms_and_conditions.htm. Attention is drawn to the limitations of fability, indemnification and jurisdictional issues established therein. The authenticity of this document may be verified at https://www.sgs.com/en/certified-clients-and-products/certified-client-directory. Any unauthonized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.