EU Certificate

for the assessment of the quality management system

according to Medical Device Regulation (EU) 2017/745 Annex IX Chapter I+III

As a Notified Body of the European Union DEKRA Certification GmbH certifies, that the manufacturer

elm-plastic GmbH

Single Registration Number (SRN): DE-MF-000005725
Kollenbergstraße 7, 54647 Dudeldorf, Germany

applies a quality management system according to Annex IX Chapter I+III of the Medical Device Regulation (EU) 2017/745 for the medical devices listed in the annex. This certificate is based on the assessments listed in CNo50809-00 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 50809-60-00-00

Certificate valid from: 2024-07-10
Certificate valid to: 2028-01-30

Digitally signed by Stephanie Donner
Date: 2024-07-12 11:19:06+02:00

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-092

DEKRA Certification GmbH, Stuttgart
Notified Body ID number: 0124

Annex to the EU Certificate no. 50809-60-00-00

Following devices/device categories are included in this certificate:

Class Im

For the devices listed below, the review of the quality management system refers exclusively to the aspects relating to the conformity of the devices with the metrological requirements.

MDN 1202, MDS 1010

- Dosing Pipettes
  - 42517534DPI911AD; 42517534DPI921AG; 42517534SFP711E9;
- Measuring beakers, and -spoons
  - 42517534MCS611BA

Change(s) to previous certificate: n.a.