Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten

Baden-Württemberg

Bayern

Berlin

Brandenburg

Bremen

Hamburg

Hessen

Mecklenburg-Vorpommern

RECOGNITION

The Central Authority of the Länder for Health Protection with regard to Medicinal Products and Medical Devices (ZLG) has determined in a recognition procedure that

SAS Hagmann GmbH & Co. KG

Weberstrasse 3
72160 Horb am Neckar
Germany

is competent under the terms of

Council Regulation 2017/745 and EN ISO/IEC 17025

for

chemical testing of medical devices.

This recognition according to § 18 Medical Device Law Implementation Act is valid up to **2028-05-11**.

This document is valid only in conjunction with the recognition notice which contains the binding information on the recognition. The scope of the recognition is specified in the annex in force of the recognition notice and can be found on www.zlg.de.

Registration number **ZLG-PL-MDR.018.23**

Bonn, 2023-05-12

Dr Rainer Edelhäuser
Director of ZLG

Thüringen

Schleswig-Holstein

Sachsen

Sachsen-Anhalt

Saarland

Rheinland-Pfalz

Nordrhein-Westfalen

Niedersachsen

Basis of recognition

Council Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC

Medical Device Law Implementation Act of 12 May 2021 concerning medical devices

EN ISO/IEC 17025 : 2018-03	General requirements for the competence of testing and calibration laboratories

For the use of indications on the status of recognition the document of ZLG 1000 HI02 applies (www.zlg.de).