Certificate

Quality Management System

ecm Zertifizierungsgesellschaft für Medizinprodukte in Europa mbH, Bismarckstr. 106, 52066 Aachen, Germany, hereby declares that an examination of the undermentioned quality assurance system has been carried out following the requirements of DIN EN ISO 13485:2016.

Through an audit performed on behalf of

C. Hedenkamp GmbH & Co. KG
Schierbusch 1, 33161 Hövelhof, Germany

it could be demonstrated that a quality management system

according to **DIN EN ISO 13485:2016**
"Medical devices – Quality management systems – Requirements for regulatory purposes"

for the **design, development, manufacturing and contract manufacturing of medical devices as capsules, chewable tablets, tablets to swallow, lozenges and effervescent tablets and devices in powder form**

has been established and implemented.

This certificate is only valid under the conditions stated in the audit report mentioned hereafter. Any substantial changes of the quality assurance have to be notified to ECM and are subject to a separate assessment.

Report Number	Registered under	Valid until
656-21-915	**Z/21/04760E**	**November 14th, 2024**

Valid as of: November 15th, 2021

Certification Body