

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

ZeoBent Handels GmbH

Weißer 2, 07407 Uhlstädt-Kirchhasel, 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

development, manufacture and distribution of zeolite and bentonite products

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

875-21-812

Registered under

Z/21/04750E

Valid until

September 17th, 2024

Valid as of: September 18th, 2021

A handwritten signature in blue ink, appearing to read 'N. Devans', is written over a horizontal line.

Certification Body