

CERTIFICATE



EN ISO 13485:2016

DEKRA Certification GmbH hereby certifies that the company

PAJUNK GmbH Medizintechnologie

Scope of certification:

Development, manufacturing, contract manufacturing and distribution of sterile and non-sterile, active and non-active, disposable and reusable medical devices

Certified location:

Karl-Hall-Straße 1, 78187 Geisingen, Germany

(further locations see annex)

has established and maintains a quality management system according to the above mentioned standard. The conformity was adduced with audit report no. 51268-Z3-00.

This certificate is valid from 2018-03-22 to 2021-03-21

Registration No.: 51268-14-00


Ruth Delbeck-Bayer



DEKRA Certification GmbH Stuttgart; 2018-03-20



Annex to the Certificate No. 51268-14-00

Revision status: 0

valid from 2018-03-22 to 2021-03-21

The following locations belong to the certificate above:

	Headquarters	Certified location	Scope of certification
	PAJUNK GmbH Medizintechnologie	Karl-Hall-Straße 1 D-78187 Geisingen	Development, manufacturing, contract manufacturing and distribution of sterile and non-sterile, active and non-active, disposable and reusable medical devices
	Subsidiaries	Certified locations	Scope of certification
1.	PAJUNK GmbH Medizintechnologie	Tuttlinger Straße 7 D-78187 Geisingen	Development, manufacturing, contract manufacturing and distribution of sterile and non-sterile, active and non-active, disposable and reusable medical devices



Ruth Delbeck-Bayer
DEKRA Certification GmbH, Stuttgart, 2018-03-20

