Fielmann

Zentrale

CE

Name of the manufacturer:	Fielmann AG Weidestraße 118a 22083 Hamburg
Single Registration Number (SRN)	DE-MF-000006245

We declare under our sole responsibility the conformity with Regulation (EU) 2017/745 of the European Parliament and of the Council for the following product:

Basic UDI DI according to Annex VI MDR:	4066209metfuladu6Z
Product name:	BD 333 CL
Risk class:	1 (according to rules set out in Annex VIII MDR)

The above-mentioned product is a medical device according to article 2 (1) of the Regulation. It fulfils the basic safety and performance requirements according to Annex I of the Regulation and complies with the state of the art and applicable harmonized standards.

Hamburg, 31.Mai 2021

Place, date

Fielmann
Fielmann AG
Weidestraße 118a
22083 Hamburg
Telefon: 040 27076-0

Christian Daiber
Qualified Person (Art. 15 MDR)