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We, the company

Ernst Kratz GmbH, Goerzallee 263, D-14167 Berlin

in our capacity as manufacturer declare in own responsibility that the medical device

CANNULA, NON-STERILE (Injection cannula, Puncture cannula, Irrigation cannula)

meets all the provisions of the Directive 93/42/EEC Annex VII which applied to him. The product is designed and manufactured in compliance with Directive 93/42/EEC, the essential requirements (appendix I) are fulfilled. Applicable harmonized standards were used. This declaration is valid for all supplies after the date of issue.

The technical documentation is kept by the manufacturer.

Berlin, 29th October 2019

Silja KLUGE

(Managing Director, Safety officer for Medical Devices)