B. Braun Medical
Société par actions simplifiée
Etablissement de CHASSENEUIL-DU-POITOU
30 Avenue des Temps Modernes
86360 CHASSENEUIL-DU-POITOU

To the competent Health Authority of COLOMBIA

Letter of Authorization

This letter will serve to authorize: **B. BRAUN MEDICAL S.A.**
Carrera 19 # 100-45, Piso 6
Bogotá, Colombia

who is the representative in Colombia for medical devices, which are manufactured and distributed, under the legal responsibility according to the medical device directive 93/42/EEC, by:

B. BRAUN MEDICAL
26 rue Armengaud
92210 Saint-Cloud, France

to register, market, import and distribute the following product groups:

Implantable medical devices and accessories:

- Celsite access ports,
- Celsite IMPLANTOFIX access ports,
- Celsite Discreet access ports,
- Surecan and Cytocan needles,
- Surecan Safety II needles.

For which the design and regulatory affairs activities are under the responsibility of:
For which the manufacturing site for all products mentioned above except needles are:

B.Braun Medical
30 Avenue des Temps Modernes
86360 Chasseneuil du Poitou, France

For which manufacturing site for needles is: **B.Braun Medical Industries Sdn. Bhd.**
Bayan Lepas Free, Industrial zone
11900 Penang, Malaysia

This Authorization Letter is effective for a period of 5 years from its date of signature.

Yours faithfully,

Date: 15/06/2022

Manuelle SCHNEIDER-PONSOT
Director of Regulatory and Pharmaceutics Operations
General Manager
B. BRAUN Medical

Catherine BOISMENU
Deputy Director in charge of Quality
and delegated Regulatory Affairs
B.BRAUN Medical

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