Engineering for clinical success

Eurocor

# Letter of Authorization

03/02/2022

To whom it may concern:

We, Eurocor Tech GmbH (In den Dauen 6 a, 53117 Bonn, Germany), as the Product Owner, hereby authorize Farmina Med, Chisinau MD -2 as Distributor and Agent for Registration of Eurocor Tech Products in Moldova.

For the purpose of complying with the regulatory registration the Distributor is authorized to prepare and submit applications for the evaluation and registration of medical devices to the competent authority on our behalf.

This authorization shall apply to the following medical devices:

Cardiovascular Products:

- DIOR - Paclitaxel eluting PTCA Balloon Catheter
- Eurolimus – Sirolimus Eluting Coronary Stent System

Endovascular Products:

- Freeway 014 - Paclitaxel eluting PTA balloon catheter
- Freeway 035 - Paclitaxel eluting PTA balloon catheter (PTA / AV-Shunt)
- Joker 014 - Balloon Dilatation Catheter
- Joker 035 - Balloon Dilatation Catheter

Eurocor Tech GmbH is a Manufacturer of coronary and peripheral balloon dilatation catheters for use in interventional cardiology and peripheral applications.

This letter of Authorization is valid till February 2024 or in case of subject to review.

Either party can terminate the contract within 60 days written notice.

Any information provided by Eurocor Tech GmbH regarding the device/s is only to be used to support the registration in Moldova and is otherwise to be kept strictly confidential.

Signed on behalf of Eurocor Tech GmbH, Bonn

Eurocor Tech GmbH
In den Dauen 6a
53117 Bonn - Germany
eurocor.de

Katja Hausner
Vice President

Digitally signed by Guțan Ghenadie
Date: 2022.02.03 16:08:06 EET
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