EC Certificate

TÜVRheinland®

Production Quality Assurance
Directive 93/42/EEC on Medical Devices, Annex V

Registration No.: DD 1024978-1

Manufacturer: CRYONOVE PHARMA SAS
50 rue de Chanzy
28000 Chartres
France

Products: - Cryotherapy devices for treatment of pigmentary skin lesions

(see attachment for site included)

Replaces EC Certificate. Registration No. DD 60126851 0001

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Report No.: 73070872 400

Effective date: 2020-07-07

Expiry date: 2023-01-10

Issue date: 2020-07-07

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle TÜVRheinland

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TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

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No.	Location	Product groups manufactured
/01	CRYONOVE PHARMA SAS 50 rue de Chanzy 28000 Chartres France	Activity: Cryotherapy devices for treatment of pigmentary skin lesions
/02	CRYONOVE PHARMA SAS 6 Parc des Fontenelles 78870 Bailly France	Activity: Manufacture

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