



Franklab®
notre expertise l'Ultra-Propreté

DCE 474
F1v2_05.2021_EN

DECLARATION OF CONFORMITY

The manufacturer: **FRANKLAB**
Z.A. De L'Observatoire
3 Avenue de Frênes
78180 Montigny-Le-Bretonneux
FRANCE

Hereby declares that, the following product:

PERALEX 9 Hecto +,
Class IIb Medical device according to rule 15 of annex IX of Medical
Device Directive 93/42/EEC

Is manufactured and delivered in accordance with the following directives:

Medical Device Directive 93/42/EEC (14 June 1993)
Public Health Code: Book II

This statement of conformity is based on Technical File (DT Générique 2) constituted according annex II.3 of Medical Device Directive 93/42/EEC and the EC Certificate delivered by the GMED (certificate n°28791 rev. 13 of the 7th May 2021), notified body n°0459.

The product is placed on the market with following packaging:

PERALEX 9 Hecto +

- 5L can

Ref.: 1047405

7th May 2021,

Julien CHARRAT
General Director

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