Franklab®
notre expertise l'Ultra-Propreté

DCE 020
FSv2_09.2022_EN

DECLARATION OF CONFORMITY

The manufacturer: FRANKLAB
Z.A. De L'Observatoire 2
3 Avenue de Frênes
78180 Montigny-Le-Bretonneux
FRANCE

Digitally signed by Bunic Gheorghe
Date: 2026.04.05 12:43:26 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

Hereby declares that the following product:

TFD7
Class I Medical device according to rule 1 of annex VIII of the European Medical Device Regulation 2017/745.

Is manufactured and delivered in accordance with the following regulations:

European Medical Device Regulation 2017/745 (April 5th, 2017)
Public Health Code: Part 5 Book II

This statement of conformity is based on Technical File (DT Générique 3) constituted according to the annex II of the European Medical Device Regulation 2017/745.The product is placed on the market with following packaging:

TFD7 (Basic UDI-ID: 37013875020ML)

• 5L Can	Ref. : 1020805
• Low 5L Can	Ref. : 1020805B
• 10L Can	Ref. : 1020811
• 20L Can	Ref. : 1020821
• 200L Drum	Ref. : 1020889
• 200L Drum with drumtainer	Ref. : 10208128

26th September 2022,

FRANKLAB SAS
3 av. des Frênes - ZA de l'Observatoire
78180 MONTIGNY LE BRETONNEUX
Tél. 01 39 44 93 40 - Fax 01 39 44 93 41
@internet : http://www.franklab.com
RC Versailles 306 563 206 - APE 2041Z

Nicolas VARAY
President

S.R.L. "SOFRAGRUP"

Franklab®
notre expertise l'Ultra-Propreté

DCE 1060
F3v2_09.2022_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:

RINCE L7
Class I Medical device according to rule 1 of annex VIII of the European Medical Device Regulation 2017/745.

Is manufactured and delivered in accordance with the following regulations:

European Medical Device Regulation 2017/745 (April 5th 2017)
Public Health Code: Part 5 Book II

This statement of conformity is based on Technical File (DT RINCE L7) constituted according to the annex II of the European Medical Device Regulation 2017/745.

The product is placed on the market with following packaging:

RINCE L7 (Basic UDI-ID: 3701387510608P)

- 5L Can — Ref. : 1032905
- Low 5L Can — Ref. : 1032905B
- 10L Can — Ref. : 1032911
- 20L Can — Ref. : 1032921
- 200L Drum — Ref. : 1032989
- 200L Drum with drumtainer — Ref. : 10329128

26th September 2022,

Nicolas VARAY
President

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Franklab®
notre expertise l'Ultra-Propreté

DCE 050
F2v2_09.2022_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:

PHOSPHAX
Class I Medical device according to rule 1 of annex VIII of the European Medical Device Regulation 2017/745.

Is manufactured and delivered in accordance with the following regulations:

European Medical Device Regulation 2017/745 (April 5th, 2017)
Public Health Code: Part 5 Book II

This statement of conformity is based on Technical File (DT PHOSPHAX) constituted according to the annex II of the European Medical Device Regulation 2017/745.

The product is placed on the market with following packaging:

PHOSPHAX (Basic UDI-DI : 37013875050MV)

- 5L Can — Ref. : 1031705
- Low 5L Can — Ref. : 1031705B
- 10L Can — Ref. : 1031711
- 20L Can — Ref. : 1031721
- 200L Drum — Ref. : 1031789
- 200L Drum with drumtainer — Ref. : 10317128

26th September 2022,

Nicolas VARAY
President

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Franklab®
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DCE 271
F1v2_09.2022_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:

ENZYMEX LD
Class I Medical device according to rule 1 of annex VIII of the European Medical Device Regulation 2017/745.

Is manufactured and delivered in accordance with the following regulations:

European Medical Device Regulation 2017/745 (April 5th 2017)
Public Health Code: Part 5 Book II

This statement of conformity is based on Technical File (DT ENZYMEX LD) constituted according to the annex II of the European Medical Device Regulation 2017/745.

The product is placed on the market with following packaging:

ENZYMEX LD (Basic UDI-ID : 37013875271NF)

- 1L Dosing bottle Ref. : 1027101
- 5L Can Ref. : 1027105
- 10L Can Ref. : 1027111

26th September 2022,

Nicolas VARAY
President

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S.R.L. "SOFRAGRUP"