

Franklab®
notre expertise l'Ultra-Propreté

DCE 020
F5v2_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:

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



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contact@sterifrance.com



3 avenue des Frênes - 78180 Montigny le Bretonneux - France
S.A.S. au capital de 1 000 000 € - RCS Versailles 76 B 635 - SIRET 306 563 206 00052 - APE 2041Z - TVA FR3306563206

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,

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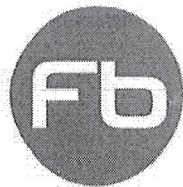


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DCE 050
F2v2_09.2022_EN

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)

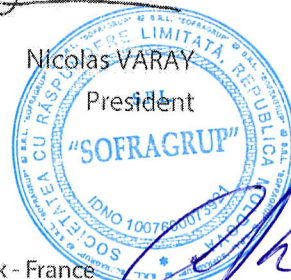
- 5L Can Ref. : 1031705
- Low 5L Can Ref. : 1031705B
- 10L Can Ref. : 1031711
- 20L Can Ref. : 1031721
- 200L Drum Ref. : 1031789
- 200L Drum with drumtainer Ref. : 10317128

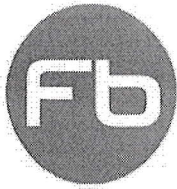
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Franklab®
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DCE 271
F1v2_09.2022_EN

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

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