

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

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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

N 
Nicolas VARAY
President

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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 FR38306563206



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 |

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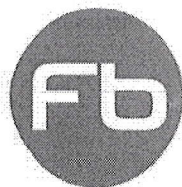


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DCE 050
F2v2_09.2022_EN

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)

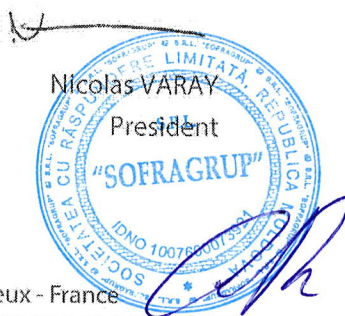
- | | |
|-----------------------------|----------------|
| • 5L Can | Ref.: 1031705 |
| • Low 5L Can | Ref.: 1031705B |
| • 10L Can | Ref.: 1031711 |
| • 20L Can | Ref.: 1031721 |
| • 200L Drum | Ref.: 1031789 |
| • 200L Drum with drumtainer | Ref.: 10317128 |

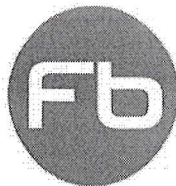
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DCE 271
F1v2_09.2022_EN

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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