

FEBRIVAC 3-PLUS

Not
authorised

- PSEUDOMONAS AERUGINOSA IMMUNOTYPE 5 ANTIGENS
- PSEUDOMONAS AERUGINOSA IMMUNOTYPE 6 ANTIGENS
- PSEUDOMONAS AERUGINOSA IMMUNOTYPE 7,8 ANTIGENS
- Clostridium botulinum, type C, toxoid
- Mink enteritis virus, Inactivated

Product identification

Medicine name:

FEBRIVAC 3-PLUS

Active substance:

PSEUDOMONAS AERUGINOSA IMMUNOTYPE 5 ANTIGENS

PSEUDOMONAS AERUGINOSA IMMUNOTYPE 6 ANTIGENS

PSEUDOMONAS AERUGINOSA IMMUNOTYPE 7,8 ANTIGENS

Clostridium botulinum, type C, toxoid

Mink enteritis virus, Inactivated

Target species:

Mink

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

PSEUDOMONAS AERUGINOSA IMMUNOTYPE 5 ANTIGENS

100.00 million cells / 1.00 millilitre(s)

PSEUDOMONAS AERUGINOSA IMMUNOTYPE 6 ANTIGENS

100.00 million cells / 1.00 millilitre(s)

PSEUDOMONAS AERUGINOSA IMMUNOTYPE 7,8 ANTIGENS

100.00 million cells / 1.00 millilitre(s)

Clostridium botulinum, type C, toxoid

0.50 relative potency / 1.00 millilitre(s)

Mink enteritis virus, Inactivated

4.00 log₁₀ 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Mink

- Unspecified. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Italy

Package description:

Available only in [Italian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

IDT Biologika GmbH

Marketing authorisation date:

15/02/1999

Manufacturing sites for batch release:

IDT Biologika GmbH

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

19/04/2024

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.