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

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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.