

Febrivac 3-Plus

Not
authorised

- Mink enteritis virus, Inactivated
- Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
- Clostridium botulinum, type C, strain Stockholm, toxoid
- Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
- Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated

Product identification

Medicine name:

Febrivac 3-Plus

Active substance:

Mink enteritis virus, Inactivated

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated

Clostridium botulinum, type C, strain Stockholm, toxoid

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated

Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated

Target species:

Mink

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Mink enteritis virus, Inactivated

10000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated

100000000.00 cells / 1.00 millilitre(s)

Clostridium botulinum, type C, strain Stockholm, toxoid

0.50 relative unit(s) / 1.00 millilitre(s)

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated

100000000.00 cells / 1.00 millilitre(s)

Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated

100000000.00 cells / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Germany

Package description:

Available only in German

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

IDT Biologika GmbH

Marketing authorisation date:

15/06/1995

Manufacturing sites for batch release:

IDT Biologika GmbH

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

137a/92

Date of authorisation status change:

18/03/2026

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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