Febrivac 3-Plus Zawiesina do wstrzykiwań

Authorised

- Mink enteritis virus, strain E-MINK F1, Inactivated
- Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
- Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
- Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated
- Clostridium botulinum, type C, strain Stockholm, toxoid

Product identification

Medicine name:
Febrivac 3-Plus Zawiesina do wstrzykiwań

Active substance:
Mink enteritis virus, strain E-MINK F1, Inactivated
Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated
Clostridium botulinum, type C, strain Stockholm, toxoid

Target species:
Mink

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Mink enteritis virus, strain E-MINK F1, Inactivated

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated

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

Clostridium botulinum, type C, strain Stockholm, toxoid

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Mink**
  - All relevant tissues. no withdrawal period — Not applicable.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in Polish

Available only in Polish

Available only in Polish

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:
CZ Vaccines S.A.U.

Marketing authorisation date:
16/04/2002

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1262

Date of authorisation status change:
16/04/2002

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

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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