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Febrivac 3-PLUS injekcinė suspensija audinėms

Authorised

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

Product identification

Medicine name:
Febrivac 3-PLUS injekcinė suspensija audinėms

Active substance:
Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated
Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
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, serotype 7/8, strain EP1, Inactivated
100000000.00 cells / 1.00 Dose

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
100000000.00 cells / 1.00 Dose

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
100000000.00 cells / 1.00 Dose

Clostridium botulinum, type C, toxoid
0.50 relative unit(s) / 1.00 Dose

Mink enteritis virus, Inactivated
10000.00 50% tissue culture infectious dose / 1.00 Dose

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:

Valid

Authorised in:

Lithuania

Package description:

Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

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:
11/09/2000

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/00/1166/001-003

Date of authorisation status change:
21/09/2010

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

Documents

Summary of Product Characteristics

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