

Febrivac DE Vet. pulver og solvens til injektionsvæske, suspension

Not authorised

- Canine distemper virus, strain Lederle D84/1, Live
- Mink enteritis virus, strain E-MINK F1, Inactivated

Product identification

Medicine name:

Febrivac DE Vet. pulver og solvens til injektionsvæske, suspension

Active substance:

Canine distemper virus, strain Lederle D84/1, Live
Mink enteritis virus, strain E-MINK F1, Inactivated

Target species:

Fur animals

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine distemper virus, strain Lederle D84/1, Live
5012.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Mink enteritis virus, strain E-MINK F1, Inactivated
10000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:

Powder and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CH01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Denmark

Package description:

Available only in [Danish](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

IDT Biologika GmbH

Marketing authorisation date:

12/11/1997

Manufacturing sites for batch release:

IDT Biologika GmbH

Responsible authority:

Danish Medicines Agency

Authorisation number:15553

Date of authorisation status change:29/04/2024

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.