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.