

Distemink Vet, lyophilisate and solvent for suspension for injection for minks

Not
authorised

- Canine distemper virus, strain Lederle, Live
- Water for injection

Product identification

Medicine name:

Distemink Vet, lyophilisate and solvent for suspension for injection for minks

Active substance:

Canine distemper virus, strain Lederle, Live

Water for injection

Target species:

Mink

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine distemper virus, strain Lederle, Live

3.00 log₁₀ unit(s) / 1.00 millilitre(s)

Water for injection
1.00 millilitre(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Italy

Package description:

Type 1 glass vials closed with a rubber stopper and sealed with an aluminum cap.
Pack containing 1 vial of 250 doses of lyophilisate (per dose of 1 ml).
Type 1 glass vials closed with a rubber stopper and sealed with an aluminum cap.
Pack containing 1 vial of 100 doses of lyophilisate (per dose of 1 ml).
1 vial of 250 ml of solvent
1 vial of 100 ml of solvent

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Similar biological application (Article 13(4) of Directive No 2001/82/EC)

Marketing authorisation holder:

United Vaccines Holding B.V.

Marketing authorisation date:

27/06/2018

Manufacturing sites for batch release:

Coooperative Federation Of Nobel Animal Keepers Nederasselt U.A.

Responsible authority:

Ministry Of Health

Authorisation number:

105149

Date of authorisation status change:

22/06/2024

Reference member state:

Netherlands

Procedure number:

NL/V/0238/001

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

Documents

Combined File of all Documents

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