

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

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

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**Active substance:**

Canine distemper virus, strain Lederle, Live

Water for injection

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**Target species:**

Mink

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Canine distemper virus, strain Lederle, Live

3.00 log<sub>10</sub> unit(s) / 1.00 millilitre(s)

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

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**Pharmaceutical form:**

Suspension for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI20CD01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

United Vaccines Holding B.V.

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**Marketing authorisation date:**

7/06/2018

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**Manufacturing sites for batch release:**

Coooperative Federation Of Nobel Animal Keepers Nederasselt U.A.

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

80863/08-07-2024/K-0229701

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**Date of authorisation status change:**

7/07/2024

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0238/001

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**Concerned member states:**

Denmark Finland Greece Italy Latvia Lithuania Norway Poland Romania  
Spain Sweden

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.

## Package Leaflet

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