Febrivac DIST, süstesuspensiooni lüofilisaat ja lahusti minkidele ja tuhkrutele

Not authorised

- Canine distemper virus, strain Lederle D84/1, Live

Product identification

Medicine name:
Febrivac DIST, süstesuspensiooni lüofilisaat ja lahusti minkidele ja tuhkrutele

Active substance:
Canine distemper virus, strain Lederle D84/1, Live

Target species:
Mink

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 dose

Pharmaceutical form:

Lyophilisate and solvent for 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:
Estonia

Package description:
Available only in Estonian
Available only in Estonian
Available only in Estonian

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:
IDT Biologika GmbH

Marketing authorisation date:
4/12/2003

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
State Agency Of Medicines

Authorisation number:

1160

Date of authorisation status change:
26/11/2025

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

Documents

Summary of Product Characteristics

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