Febrivac HV Vet. lyofilisat og solvens til injektionsvæske, suspension

Authorised

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

Product identification

Medicine name:
Febrivac HV Vet. lyofilisat og solvens til injektionsvæske, suspension
Febrivac HV Vet. lyofilisat og solvens til injektionsvæske, suspension

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

Target species:
Fur animals

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, strain Lederle D84/1, Live
5012.00 tissue culture infective dose 50 / 1.00 millilitre(s)

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Fur animals**

Subcutaneous use:

- **Fur animals**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI20CD01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

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:
3/08/2007

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
39060

Date of authorisation status change:
3/08/2007

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.
