Nobivac DHPPi vet. injektiokuiva-aine ja liuotin, suspensiota varten

Authorised

- Canine parainfluenza virus, strain Cornell, Live
- Canine parvovirus, strain 154, Live
- Canine adenovirus 2, strain Manhattan LPV3, Live
- Canine distemper virus, strain Onderstepoort, Live

Product identification

Medicine name:
Nobivac DHPPi vet. injektiokuiva-aine ja liuotin, suspensiota varten

Active substance:
Canine parainfluenza virus, strain Cornell, Live

Canine parvovirus, strain 154, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

Canine distemper virus, strain Onderstepoort, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Canine parainfluenza virus, strain Cornell, Live
5.50 log10 50% tissue culture infectious dose / 1.00 dose

Canine parvovirus, strain 154, Live
7.00 log10 50% tissue culture infectious dose / 1.00 dose

Canine adenovirus 2, strain Manhattan LPV3, Live
4.00 log10 50% tissue culture infectious dose / 1.00 dose

Canine distemper virus, strain Onderstepoort, Live
4.00 log10 50% tissue culture infectious dose / 1.00 dose

Pharmaceutical form:

Powder and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
  - Not applicable. no withdrawal period — no period

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:
Available only in Finnish
Available only in Finnish

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:
Intervet International B.V.

Marketing authorisation date:
11/04/1996

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
Finnish Medicines Agency

Authorisation number:
13483

Date of authorisation status change:
11/04/1996

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.

Package Leaflet

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