Nobivac DHPPi vet. Pulver och vätska till injektionsvätska, suspension

Authorised

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

Product identification

Medicine name:
Nobivac DHPPi vet. Pulver och vätska till injektionsvätska, suspension

Active substance:
Canine parvovirus, strain 154, Live

Canine parainfluenza virus, strain Cornell, Live

Canine distemper virus, strain Onderstepoort, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

Water

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

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

Canine parainfluenza virus, strain Cornell, Live
5.50 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

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

Water
1.00 millilitre(s) / 1.00 Dose

Pharmaceutical form:

Powder and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
24/02/1995

Manufacturing sites for batch release:
Oriola Sweden AB
Intervet International B.V.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
12612

Date of authorisation status change:
24/02/1995

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

Documents

Package Leaflet

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Summary of Product Characteristics

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