

Nobivac DHPPi liofilizāts un šķīdinātājs suspensijas injekcijām pagatavošanai suņiem

Authorised

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

Product identification

Medicine name:

Nobivac DHPPi liofilizāts un šķīdinātājs suspensijas injekcijām pagatavošanai suņiem

Active substance:

Canine distemper virus, strain Onderstepoort, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

Canine parainfluenza virus, strain Cornell, Live

Canine parvovirus, strain 154, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine distemper virus, strain Onderstepoort, Live
10000.00 50% tissue culture infectious dose / 1.00 unit(s)

Canine adenovirus 2, strain Manhattan LPV3, Live
10000.00 50% tissue culture infectious dose / 1.00 unit(s)

Canine parainfluenza virus, strain Cornell, Live
316228.00 50% tissue culture infectious dose / 1.00 unit(s)

Canine parvovirus, strain 154, Live
10000000.00 50% tissue culture infectious dose / 1.00 unit(s)

Pharmaceutical form:

Lyophilisate and solvent for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Available in:

Latvia

Package description:

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

14/10/1993

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/NRP/93/0047

Date of authorisation status change:

14/10/1993

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

Documents

Combined File of all Documents

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