Nobivac SHPPi, Lyophilisat und Lösungsmittel zur Herstellung einer Injektionssuspension für Hunde

Authorised

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

Product identification

Medicine name:
Nobivac SHPPi, Lyophilisat und Lösungsmittel zur Herstellung einer Injektionssuspension für Hunde

Active substance:
Canine distemper virus, strain Onderstepoort, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

Canine parvovirus, strain 154, Live

Canine parainfluenza virus, strain Cornell, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, strain Onderstepoort, Live
6.00 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine adenovirus 2, strain Manhattan LPV3, Live
6.50 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parvovirus, strain 154, Live
8.40 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parainfluenza virus, strain Cornell, Live
7.00 log10 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:
Lyophilisate and solvent for suspension 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:
Austria

Package description:
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet Ges.m.b.H.

Marketing authorisation date:
1/02/1994

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-20124

Date of authorisation status change:
1/02/1994

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

Documents

Labelling

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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.