

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
10000.00 50% tissue culture infectious dose / 1.00 Dose

Canine adenovirus 2, strain Manhattan LPV3, Live
10000.00 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, strain 154, Live
10000000.00 50% tissue culture infectious dose / 1.00 Dose

Canine parainfluenza virus, strain Cornell, Live
316228.00 50% tissue culture infectious dose / 1.00 Dose

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:

Germany

Available in:

Germany

Package description:

Available only in [German](#)

Available only in [German](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet Deutschland GmbH

Marketing authorisation date:

12/09/2002

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

205a/97

Date of authorisation status change:

17/05/2011

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

Documents

Combined File of all Documents

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