# 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 tissue culture infective dose 50 / 1.00 millilitre(s)

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

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

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

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**Pharmaceutical form:**

Lyophilisate and solvent for suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Dog**

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI07AD04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria

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**Package description:**

Available only in German

Available only in German

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## 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

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.

Summary of Product Characteristics

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

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