Nobivac CEPPi, liofilizzato per sospensione iniettabile per cani

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

Product identification

Medicine name:
Nobivac CEPPi, liofilizzato per sospensione iniettabile per cani

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

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 unit(s)

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

Canine distemper virus, strain Onderstepoort, Live
4.00 log10 50% tissue culture infectious dose / 1.00 unit(s)

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

Pharmaceutical form:

Lyophilisate for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
  - Unspecified. 0 day

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:

Italy

Available in:

Italy

Package description:

Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

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 International B.V.

Marketing authorisation date:
30/07/1993

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

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
16/10/1997

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.