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Nobivac DP Plus (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Canine distemper virus, strain Onderstepoort, Live
- Canine parvovirus, strain 630a (recombinant), Live

Product identification

Medicine name:
Nobivac DP Plus (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Canine distemper virus, strain Onderstepoort, Live

Canine parvovirus, strain 630a (recombinant), Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, strain Onderstepoort, Live

Presentation_strength:$10^{5.1}$ – $10^{6.5}$ $TCID_{50}$ Index:0

Canine parvovirus, strain 630a (recombinant), Live

Presentation_strength:$10^{5.1}$ – $10^{6.7}$ $TCID_{50}$ Index:8

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:Lyophiliisate: 25 vials + solvent: 25 vials, Content:Lyophilisate: 1 dose + solvent: 1 ml

Packaging:Vial (glass), Package_size:Lyophiliisate: 5 vials + solvent: 5 vials, Content:Lyophilisate: 1 dose + solvent: 1 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
9/12/2020

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

Responsible authority:
European Commission

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

Date of authorisation status change:
9/12/2020

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
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