

Versican Plus DP lyophilisate and solvent for suspension for injection for dogs

Authorised

- Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live
- Canine distemper virus, strain CDV Bio 11/A, Live

Product identification

Medicine name:

Versican Plus DP lyophilisate and solvent for suspension for injection for dogs

Versican Plus DP Liofilizat (żywy, atenuowany):-Wirus nosówki psów, szczep CDV Bio 11/A $\geq 10^{3,1}$ TCID₅₀*, $\leq 10^{5,1}$ TCID₅₀,-Parwovirus psów, typ 2b, szczep CPV-2b-Bio 12/B $\geq 10^{4,3}$ TCID₅₀*, $\leq 10^{6,6}$ TCID₅₀.(*) Dawka zakaźna dla 50 % hodowli tkankowych Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

Active substance:

Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live

Canine distemper virus, strain CDV Bio 11/A, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine parvovirus, type 2b, strain CPV-2b Bio 12/B, Live
19952.00 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, strain CDV Bio 11/A, Live
1258.00 50% tissue culture infectious dose / 1.00 Dose

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:

Poland

Available in:

Poland

Package description:

(ID2) 50 Dose; 50 millilitre(s): Box (plastics) with 50 Vial (glass) each with 1 Dose and 50 Vial (glass) each with 1 millilitre(s)

(ID1) 25 Dose; 25 millilitre(s): Box (plastics) with 25 Vial (glass) each with 1 Dose and 25 Vial (glass) each with 1 millilitre(s)

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:

Zoetis Polska Sp. z o.o.

Marketing authorisation date:

31/10/2016

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2570

Date of authorisation status change:

31/10/2016

Reference member state:

Germany

Procedure number:

DE/V/0266/001

Concerned member states:

Belgium Bulgaria Cyprus Greece Hungary Italy Lithuania Luxembourg
Netherlands Poland Portugal Romania Spain
United Kingdom (Northern Ireland)

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

Documents

2613562-paren-20251101.pdf