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

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:
Netherlands

Available in:
Netherlands

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 B.V.

Marketing authorisation date:

4/04/2016

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 117278

Date of authorisation status change:
8/02/2022

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

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics

2613562-paren-20251101.pdf