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Versican Plus Pi (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Product identification

Medicine name:
Versican Plus Pi (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine parainfluenza virus, strain CPiV-2-Bio 15, Live
Presentation_strength:$10^{3.1}$ - $10^{5.1}$ $TCID_{50}$ Reference:Ph Eur 1955
Comments:Lyophilisate Index:0

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AD08

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:
Belgium , Denmark , Finland , Germany , Luxembourg , Sweden

Package description:
Packaging:Lyophilisate: vial (glass); solvent: vial (glass), Package_size:Lyophilisate: 25 vials + solvent: 25 vials, Content:Lyophilisate: 1 dose; solvent: 1 ml
Packaging:Lyophilisate: vial (glass); solvent: vial (glass), Package_size:Lyophilisate: 50 vials + solvent: 50 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:
Zoetis Belgium

Marketing authorisation date:
4/07/2014

Manufacturing sites for batch release:
Bioveta a.s.
Ceva-Phylaxia Zrt.

Responsible authority:
European Commission

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

Date of authorisation status change:
4/07/2014

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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