VERSICAN PLUS BB ORAL LYOPHILISAT AND SOLVENT FOR ORAL SUSPENSION FOR DOGS

Authorised

- Water, purified
- Bordetella bronchiseptica, strain 92 B, Live

Product identification

Medicine name:
VERSICAN PLUS BB ORAL LYOPHILISAT AND SOLVENT FOR ORAL SUSPENSION FOR DOGS
Versican Plus Bb Oral, Lyophilisat und Lösungsmittel zur Herstellung einer Suspension zur oralen Applikation für Hunde

Active substance:
Water, purified
Bordetella bronchiseptica, strain 92 B, Live

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:

Water, purified
1.00 millilitre(s) / 1.00 Dose

Bordetella bronchiseptica, strain 92 B, Live
140000000.00 Colony forming unit / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for oral suspension

Withdrawal period by route of administration:
Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AE01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Plastic box containing 5, vials of 1 dose of lyophilisate and 5, vials of 1ml of solvent
Plastic box containing 25 vials of 1 dose of lyophilisate and 25 vials of 1ml of solvent
Plastic box containing 10 vials of 1 dose of lyophilisate and 10 vials of 1ml of solvent

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 Osterreich GmbH

Marketing authorisation date:
8/07/2019

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
839005

Date of authorisation status change:
8/07/2019

Reference member state:
France

Procedure number:
FR/V/0401/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.
