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 1,4x10^8-5.5x10^9 cfu/dose ΛΥΟΦΙΛΟΠΟΙΗΜΕΝΟ ΥΛΙΚΟ ΚΑΙ ΔΙΑΛΥΤΗΣ ΓΙΑ ΠΟΣΙΜΟ ΕΝΑΙΩΡΗΜΑ
VERSICAN PLUS BB ORAL LYOPHILISAT AND SOLVENT FOR ORAL SUSPENSION FOR DOGS

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:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Plastic box containing 10 vials of 1 dose of lyophilisate and 10 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 5, vials of 1 dose of lyophilisate and 5, 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 Hellas S.A.

Marketing authorisation date:
24/02/2020

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
National Organization For Medicines

Authorisation number:
24213/24-02-2020/K-0240301

Date of authorisation status change:
24/02/2020

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

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