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, suukaudse suspensiooni lüofilisaat ja lahusti koertele

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 14000000.00 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:

Estonia

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 Belgium

Marketing authorisation date: 28/08/2019
Manufacturing sites for batch release: Zoetis Belgium
Responsible authority: State Agency Of Medicines
Authorisation number: 2197
Date of authorisation status change: 28/08/2019
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

Summary of Product Characteristics

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