Virbagen canis B Suspension for injection for dogs

Authorised

- Borreliella afzelii, Inactivated
- Borreliella garinii, Inactivated

Product identification

Medicine name:
Virbagen canis B Suspension for injection for dogs
Virbagen canis B Injektionssuspension für Hunde

Active substance:
Borreliella afzelii, Inactivated
Borreliella garinii, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Borreliella afzelii, Inactivated
1.00 relative unit(s) / 1.00 millilitre(s)

Borreliella garinii, Inactivated
1.00 relative unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AB04

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
(ID4) 20 millilitre(s): Box (Cardboard) with 20 Vial (Glass type I) each with 1 millilitre(s), closed with Stopfen and Kappe (Gummi, Aluminium)
(ID5) 50 millilitre(s): Box (Cardboard) with 50 Vial (Glass type I) each with 1 millilitre(s), closed with Stopfen and Kappe (Gummi, Aluminium)
(ID1) 2 millilitre(s): Box (Cardboard) with 2 Vial (Glass type I) each with 1 millilitre(s), closed with Stopfen and Kappe (Gummi, Aluminium)
(ID2) 5 millilitre(s): Box (Cardboard) with 5 Vial (Glass type I) each with 1 millilitre(s), closed with Stopfen and Kappe (Gummi, Aluminium)
(ID6) 100 millilitre(s): Box (Cardboard) with 100 Vial (Glass type I) each with 1 millilitre(s), closed with Stopfen and Kappe (Gummi, Aluminium)
(ID3) 10 millilitre(s): Box (Cardboard) with 10 Vial (Glass type I) each with 1 millilitre(s), closed with Stopfen and Kappe (Gummi, Aluminium)

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:
Virbac Tierarzneimittel GmbH

Marketing authorisation date:
7/10/2009

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11433.01.1

Date of authorisation status change:
28/04/2014

Reference member state:
Germany

Procedure number:
DE/V/0250/001

Concerned member states:
Austria

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

Documents

Combined File of all Documents

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