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Borrelym 3, suspension for injection for dogs

Authorised

- Borreliella garinii, strain BR14, Inactivated
- Borreliella afzelii, strain BR33, Inactivated
- Borreliella burgdorferi, strain DSM 4681, Inactivated

Product identification

Medicine name:
Borrelym 3, suspension for injection for dogs

Active substance:
Borreliella garinii, strain BR14, Inactivated

Borreliella afzelii, strain BR33, Inactivated

Borreliella burgdorferi, strain DSM 4681, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Borreliella garinii, strain BR14, Inactivated

1.00 relative potency / 1.00 Dose

Borreliella afzelii, strain BR33, Inactivated
1.00 relative potency / 1.00 Dose

Borreliella burgdorferi, strain DSM 4681, Inactivated
1.00 relative potency / 1.00 Dose

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:
Norway

Package description:
Glass Vial 10 x 1.0 millilitre(s)
Glass Vial 2 x 1.0 millilitre(s)
Glass Vial 20 x 1.0 millilitre(s)
Glass Vial 100 x 1.0 millilitre(s)
Glass Vial 50 x 1.0 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health Denmark A/S

Marketing authorisation date:
26/03/2013

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
12-9180

Date of authorisation status change:
16/04/2019

Reference member state:
Czechia

Procedure number:
CZ/V/0114/001

Concerned member states:
Austria Belgium Denmark Estonia France Germany Hungary Ireland Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia 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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Package Leaflet

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