

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

Borrelia afzelii, strain BR33, Inactivated

1.00 relative potency / 1.00 Dose

Borrelia 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:

Slovenia

Package description:

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

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:

Bioveta a.s.

Marketing authorisation date:

8/01/2013

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

MR/V/0420/001

Date of authorisation status change:

8/01/2013

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

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

Package Leaflet

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.