# Borrelym 3, Suspension for injection

Authorised

- Borreliella afzelii, Inactivated
- Borreliella garinii, Inactivated
- Borreliella burgdorferi, Inactivated

## Product identification

**Medicine name:**
Borrelym 3, Suspension for injection
BORRELYM 3, injekcinė suspensija šunims

**Active substance:**
Borreliella afzelii, Inactivated
Borreliella garinii, Inactivated
Borreliella burgdorferi, Inactivated

**Target species:**
Dog

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Borreliella afzelii, Inactivated
1.00 relative potency / 1.00 Dose

Borreliella garinii, Inactivated
1.00 relative potency / 1.00 Dose

Borreliella burgdorferi, Inactivated
1.00 relative potency / 1.00 Dose

**Pharmaceutical form:**
Suspension for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Dog**

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI07AB04

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

**Authorisation status:**
Valid

**Authorised in:**
Lithuania

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

**Marketing authorisation date:**
This information is not available for this product.

**Manufacturing sites for batch release:**
Bioveta a.s.

**Responsible authority:**
State Food And Veterinary Service

**Authorisation number:**
LT/2/13/2163/001-005

**Date of authorisation status change:**
6/08/2019

**Reference member state:**
Czechia

**Procedure number:**
CZ/V/0114/001

**Concerned member states:**
Austria Belgium Denmark Estonia France Germany Hungary Ireland Italy 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

RV2163.pdf