Erysin Single Shot, emulzija za injekciju, za svinje

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
- Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Product identification

Medicine name:
Erysin Single Shot, emulzija za injekciju, za svinje

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Target species:
Pig

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Croatia

Package description:

Available only in Croatian

Available only in Croatian

Available only in Croatian
Available only in Croatian
Available only in Croatian

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:
13/06/2018

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/13-01/381

Date of authorisation status change:
5/01/2026

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.