

Erysin Single Shot, emulzija za injekciju, za svinje

Authorised

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

Product identification

Medicine name:

Erysin Single Shot, emulzija za injekciju, za svinje

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated

Erysipelothrix rhusiopathiae, serotype 1, strain 203, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated

Target species:

Pig

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated
1.00 relative potency / 1.00 Dose

Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
1.00 relative potency / 1.00 Dose

Erysipelothrix rhusiopathiae, serotype 1, strain 203, Inactivated
1.00 relative potency / 1.00 Dose

Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
1.00 relative potency / 1.00 Dose

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

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:

2/11/2023

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

Documents

Summary of Product Characteristics

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