

Parvosin ERY, emulzija za injekciju, za svinje

Authorised

- Erysipelothrix rhusiopathiae, serotype 1 and 2, Inactivated
- Porcine parvovirus, strain CAPM V198 S-27, Inactivated

Product identification

Medicine name:

Parvosin ERY, emulzija za injekciju, za svinje

Active substance:

Erysipelothrix rhusiopathiae, serotype 1 and 2, Inactivated
Porcine parvovirus, strain CAPM V198 S-27, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 1 and 2, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Porcine parvovirus, strain CAPM V198 S-27, Inactivated
4.00 log2 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:**Intramuscular use:**

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Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AL01

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:

11/06/2021

Manufacturing sites for batch release:

Bioveta, a.s.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/21-01/447

Date of authorisation status change:

23/08/2023

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

Documents

Summary of Product Characteristics

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