PARVOERYSIN emulsione iniettabile

Authorised

- Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated
- Porcine parvovirus, strain CAPM V198 S-27, Inactivated

Product identification

Medicine name:
PARVOERYSIN emulsione iniettabile

Active substance:
Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, 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, strain 1-203, Inactivated
1.00 relative potency / 2.00 millilitre(s)

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

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:

- **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:

Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

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:
2/07/2018

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
2/07/2018

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

Documents

Combined File of all Documents

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