ERYGAL P – injekčná emulzia

Authorised

- Erysipelothrix rhusiopathiae, Inactivated
- Porcine parvovirus, Inactivated

Product identification

Medicine name:
ERYGAL P – injekčná emulzia

Active substance:
Erysipelothrix rhusiopathiae, Inactivated

Porcine parvovirus, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, Inactivated
1.00 Protective Dose / 2.00 millilitre(s)

Porcine parvovirus, Inactivated
7.00 log2 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - All relevant tissues. 0 day zero days

Subcutaneous use:

- **Pig**
  - All relevant tissues. 0 day zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AL03

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharmagal Bio spol. s r.o.

Marketing authorisation date:
2/05/2003

Manufacturing sites for batch release:
Pharmagal Bio spol. s r.o.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/ 025/ 03-S

Date of authorisation status change:
29/11/2022

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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