Porcilis Ery Parvo Vet. injektionsvæske, suspension

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
- Porcine parvovirus, strain 014, Inactivated
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- Porcine parvovirus, strain 014, Inactivated
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- Porcine parvovirus, strain 014, Inactivated
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- Porcine parvovirus, strain 014, Inactivated

Product identification

Medicine name:
Porcilis Ery Parvo Vet. injektionsvæske, suspension

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
Porcine parvovirus, strain 014, Inactivated
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Porcine parvovirus, strain 014, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Porcine parvovirus, strain 014, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
1.00 unit(s) / 2.00 millilitre(s)

Porcine parvovirus, strain 014, Inactivated
552.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
1.00 unit(s) / 2.00 millilitre(s)

Porcine parvovirus, strain 014, Inactivated
552.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
1.00 unit(s) / 2.00 millilitre(s)

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Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
1.00 unit(s) / 2.00 millilitre(s)

Porcine parvovirus, strain 014, Inactivated
552.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Available in:
Denmark

Package description:
Available only in Danish
Available only in Danish
Available only in Danish
Available only in Danish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
10/03/2000

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
19404

Date of authorisation status change:
10/03/2000

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

Documents

Summary of Product Characteristics

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