

Porcilis Ery Parvo Vet. injektionsvæske, suspension

Authorised

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

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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:

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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

This document does not exist in this language (English). You can find it in another language below.