

Porcilis Ery+Parvo suspensija injekcijām cūkām

Authorised

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

Product identification

Medicine name:

Porcilis Ery+Parvo suspensija injekcijām cūkām

Active substance:

Porcine parvovirus, strain 014, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Target species:

Pig (sow, nullipar)

Pig (sow)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Porcine parvovirus, strain 014, Inactivated

≥ 552 EU

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

>1 aizsargājoša deva cūkām (ppd)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

• **Pig (sow, nullipar)**

- Not specified. 0 day

• **Pig (sow)**

- Not specified. 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:

Latvia

Package description:

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in [Latvian](#)

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:

This information is not available for this product.

Manufacturing sites for batch release:

INTERVET INTERNATIONAL B.V.

Responsible authority:

PVD

Authorisation number:

V/NRP/96/0378

Date of authorisation status change:

21/03/1996

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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