Parvoruvax Vet. injektionsvæske, suspension

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, Inactivated
- Porcine parvovirus, strain K22, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, Inactivated
- Porcine parvovirus, strain K22, Inactivated
- Porcine parvovirus, strain K22, Inactivated

Product identification

Medicine name:
Parvoruvax Vet. injektionsvæske, suspension

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, Inactivated
Porcine parvovirus, strain K22, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, Inactivated
Erysipelothrix rhusiopathiae, serotype 2, Inactivated
Porcine parvovirus, strain K22, Inactivated
Porcine parvovirus, strain K22, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, Inactivated
1.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Porcine parvovirus, strain K22, Inactivated
2.00 unit(s) / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, Inactivated
1.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, Inactivated
1.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Porcine parvovirus, strain K22, Inactivated
2.00 unit(s) / 2.00 millilitre(s)

Porcine parvovirus, strain K22, Inactivated
2.00 unit(s) / 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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AL01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

Package description:
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:
Ceva Sante Animale

Marketing authorisation date:
3/06/1996

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Danish Medicines Agency

Authorisation number:
14969

Date of authorisation status change:
3/06/1996

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

Documents

Summary of Product Characteristics

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