

# Parvoruvax Vet. injektionsvæske, suspension

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, Inactivated
- Porcine parvovirus, strain K22, Inactivated
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- Porcine parvovirus, strain K22, Inactivated
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## Product identification

### Medicine name:

Parvoruvax Vet. injektionsvæske, suspension

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### Active substance:

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Porcine parvovirus, strain K22, Inactivated

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Porcine parvovirus, strain K22, Inactivated

Porcine parvovirus, strain K22, Inactivated

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### Target species:

Pig

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### Route of administration:

Intramuscular use

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## 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)

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### **Pharmaceutical form:**

Suspension for injection

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### **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
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  - Meat and offal. 0 day
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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AL01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Denmark

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**Package description:**

Available only in [Danish](#)

Available only in [Danish](#)

Available only in [Danish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

3/06/1996

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

14969

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**Date of authorisation status change:**

3/06/1996

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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