Parvoruvax

Authorised

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

Product identification

Medicine name:
Parvoruvax

Active substance:
Porcine parvovirus, strain K22, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine parvovirus, strain K22, Inactivated
1.00 unit(s) / 1.00 Dose

Erysipelothrix rhusiopathiae, serotype 2, Inactivated
1.00 unit(s) / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. no withdrawal period — Withdrawal period = zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AL01

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:

23/01/2004

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.
Boehringer Ingelheim Animal Health France

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 4217

Date of authorisation status change:
29/06/2021

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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