PARVOVAX

Authorised

- Porcine parvovirus, strain K22, Inactivated

Product identification

Medicine name:
PARVOVAX

Active substance:
Porcine parvovirus, strain K22, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine parvovirus, strain K22, Inactivated
2.00 haemagglutinating units / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AA02

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Type 1 glass bottle;Nitrile elastomer closure; Aluminium cap; 25-dose bottle, 1-bottle package.
Type 1 glass bottle;Nitrile elastomer closure; Aluminium cap; 5-dose bottle, 1-bottle package.
Type 1 glass bottle;Nitrile elastomer closure; Aluminium cap; 1-dose bottle, 10-bottle package.
Type 1 glass bottle;Nitrile elastomer closure; Aluminium cap; 1-dose bottle, 1-bottle package.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
6/11/2002

Manufacturing sites for batch release:

Merial
Ceva-Phylaxia Veterinary Biologicals Co. Ltd.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10815/053/001

Date of authorisation status change:
6/11/2002

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

Documents

Summary of Product Characteristics

Source URL: *https://medicines.health.europa.eu/veterinary/600000064555*