PARVORUVAX, injekcinė suspensija kiaulėms

Authorised

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

Product identification

Medicine name:
PARVORUVAX, injekcinė suspensija kiaulėms

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, 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 / 1.00 Dose

Porcine parvovirus, strain K22, Inactivated

2.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AL01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

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:
19/03/2002

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/02/1375/001-003

Date of authorisation status change:
10/06/2007

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

Documents

RV1375.pdf