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FIXR HP ERY emulsion for injection for pigs

Authorised

- Haemophilus parasuis, serotype 13, Inactivated
- Haemophilus parasuis, serotype 5, Inactivated
- Haemophilus parasuis, serotype 1, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
- Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Product identification

Medicine name:
FIXR HP ERY emulsion for injection for pigs

Active substance:
Haemophilus parasuis, serotype 13, Inactivated

Haemophilus parasuis, serotype 5, Inactivated

Haemophilus parasuis, serotype 1, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated

Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Haemophilus parasuis, serotype 13, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Haemophilus parasuis, serotype 5, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Haemophilus parasuis, serotype 1, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain 2-II, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 1, strain 1-203, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain 2-5, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated
1.00 unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
60 ml vial plastic vials (HDPE) containing 50 ml of the vaccine. Vials are hermetically sealed with puncturable chlorobutyl rubber stoppers, fitted with aluminium caps or flip off caps and placed in paper cardboard.
120 ml vial plastic vials (HDPE) containing 100 ml of the vaccine. Vials are hermetically sealed with puncturable chlorobutyl rubber stoppers, fitted with aluminium caps or flip off caps and placed in paper cardboard.

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Kernfarm B.V.

Marketing authorisation date:
30/03/2020

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 124851

Date of authorisation status change:
1/02/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0328/001

Concerned member states:
Belgium Germany

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

Documents

Combined File of all Documents

English (PDF)
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