FIXR RHINI suspension for injection for pigs

Authorised

- Pasteurella multocida, toxoid
- Pasteurella multocida, serogroup A, Inactivated
- Bordetella bronchiseptica, strain F236, Inactivated
- Pasteurella multocida, serogroup D, Inactivated

Product identification

Medicine name:
FIXR RHINI suspension for injection for pigs
FIXR RHINI suspension injectable pour porcins
FIXR RHINI suspensie voor injectie voor varkens
FIXR RHINI Injektionssuspension für Schweine

Active substance:
Pasteurella multocida, toxoid
Pasteurella multocida, serogroup A, Inactivated
Bordetella bronchiseptica, strain F236, Inactivated
Pasteurella multocida, serogroup D, Inactivated

Target species:
Pig

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Pasteurella multocida, toxoid
2.00 microgram(s) / 1.00 Dose

Pasteurella multocida, serogroup A, Inactivated
80.00 percentage protection / 1.00 Dose

Bordetella bronchiseptica, strain F236, Inactivated
32.00 titre / 1.00 Dose

Pasteurella multocida, serogroup D, Inactivated
80.00 percentage protection / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Belgium

Package description:

20 ml injection vial of hydrolytic class I glass closed with a rubber stopper and an aluminium seal in a cardboard box

20 ml injection vial of hydrolytic class I glass closed with a rubber stopper and an aluminium seal in a polystyrene case
100 ml injection vial of hydrolytic class I glass closed with a rubber stopper and an aluminium seal in a cardboard box
100 ml injection vial of hydrolytic class I glass closed with a rubber stopper and an aluminium seal in a polystyrene case
20 ml polypropylene bottle closed with a rubber stopper and an aluminium seal in a cardboard box
20 ml polypropylene bottle closed with a rubber stopper and an aluminium seal in a polystyrene case
100 ml polypropylene bottle closed with a rubber stopper and an aluminium seal in a cardboard box
100 ml polypropylene bottle closed with a rubber stopper and an aluminium seal in a polystyrene case

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 19(1) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Kernfarm B.V.

Marketing authorisation date:
23/02/2023

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
23/02/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0393/001

Concerned member states:
Belgium

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

Documents

Combined File of all Documents