RINI-SUIVAX T sospensione iniettabile per suini

Authorised

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

Product identification

Medicine name:
RINI-SUIVAX T sospensione iniettabile per suini

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

Target species:
Pig (female)
Pig (young female)

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Pasteurella multocida, toxoid
2.00 microgram(s) / 2.00 millilitre(s)

Pasteurella multocida, serogroup A, strain PMSA, Inactivated
80.00 percentage protection / 2.00 millilitre(s)

Pasteurella multocida, serogroup D, strain PMSD, Inactivated
80.00 percentage protection / 2.00 millilitre(s)

Bordetella bronchiseptica, strain F236, Inactivated
32.00 slow agglutination test unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Pig (female)**
  - Meat and offal. 0 day

- **Pig (young female)**
  - Meat and offal. 0 day

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:

Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

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:
Fatro S.p.A.

Marketing authorisation date:
25/08/2003

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

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
25/08/2003

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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