Rhiniseng (--) - Suspension for injection

Authorised

- Bordetella bronchiseptica, strain 833CER, Inactivated
- Pasteurella multocida, serotype D, dermonecrotic toxin, recombinant

Product identification

Medicine name:
Rhiniseng (--) - Suspension for injection

Active substance:
Bordetella bronchiseptica, strain 833CER, Inactivated

Pasteurella multocida, serotype D, dermonecrotic toxin, recombinant

Target species:
Pig (sow)
Pig (sow, nullipar)
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Bordetella bronchiseptica, strain 833CER, Inactivated

Presentation_strength:9.8 BbCC Reference:Ph Eur Index:0

Pasteurella multocida, serotype D, dermonecrotic toxin, recombinant

Presentation_strength:≥ 1 MED63 Reference:Ph Eur Index:1

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (sow)**
  - Not applicable. 0 day Zero days
- **Pig (sow, nullipar)**
  - Not applicable. 0 day Zero days
- **Pig**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (PET), Package_size:1 bottle, Content:250 ml

Packaging:Vial (PET), Package_size:1 bottle, Content:100 ml

Packaging:Vial (PET), Package_size:1 bottle, Content:50 ml

Packaging:Vial (PET), Package_size:10 bottles, Content:20 ml

Packaging:Vial (PET), Package_size:1 bottle, Content:20 ml
Packaging:Vial (glass), Package_size:1 vial, Content:100 ml
Packaging:Vial (glass), Package_size:1 vial, Content:50 ml
Packaging:Vial (glass), Package_size:10 vials, Content:20 ml
Packaging:Vial (glass), Package_size:1 vial, Content:20 ml

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:
Laboratorios Hipra, S.A.

Marketing authorisation date:
16/09/2010

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
European Commission

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

Date of authorisation status change:
16/09/2010

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

Documents

Combined File of all Documents

English (PDF)
Published on: 5/02/2024
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