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Porcilis AR-T DF (--) - Suspension for injection

Authorised

- Pasteurella multocida, protein dO (non-toxic derivative of dermonecrotic toxin), recombinant
- Bordetella bronchiseptica, Inactivated

Product identification

Medicine name:
Porcilis AR-T DF (--) - Suspension for injection

Active substance:
Pasteurella multocida, protein dO (non-toxic derivative of dermonecrotic toxin), recombinant

Bordetella bronchiseptica, Inactivated

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

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Pasteurella multocida, protein dO (non-toxic derivative of dermonecrotic toxin), recombinant

Presentation_strength:≥ 6.2 log2 TN. titre Reference:Hse Index:0

Bordetella bronchiseptica, Inactivated

Presentation_strength:≥ 5.5 log2 Aggl.titre Reference:Hse 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

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:

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)

Available in:

Austria , Belgium , Bulgaria , Cyprus , Denmark , France , Germany , Greece , Hungary , Ireland , Italy , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (polyethylene terephthalate (PET)), Package_size:1 vial, Content:100 ml (50 doses)
Packaging:Vial (polyethylene terephthalate (PET)), Package_size:1 vial, Content:250 ml (125 doses)
Packaging:Vial (type I glass), Package_size:1 vial, Content:20 ml (10 doses)
Packaging:Vial (type I glass), Package_size:1 vial, Content:50 ml (25 doses)
Packaging:Vial (polyethylene terephthalate (PET)), Package_size:1 vial, Content:20 ml (10 doses)
Packaging:Vial (polyethylene terephthalate (PET)), Package_size:1 vial, Content:50 ml (25 doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
16/11/2000

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
16/11/2000

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: 22/07/2024
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