# 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

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**Pharmaceutical form:**

Suspension for injection

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AB04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**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)

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**Package description:**
Packaging:Vial (polyethylene terephthalate (PET)), 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 (type I glass), Package_size:1 vial, Content:50 ml (25 doses)
Packaging:Vial (type I glass), Package_size:1 vial, Content:20 ml (10 doses)
Packaging:Vial (polyethylene terephthalate (PET)), Package_size:1 vial, Content:250 ml (125 doses)
Packaging:Vial (polyethylene terephthalate (PET)), Package_size:1 vial, Content:100 ml (50 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:**
23/09/2010

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

## Documents

Combined File of all Documents

English (PDF)
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