

# Porcilis ColiClos (--)- Suspension for injection

Authorised

- Escherichia coli, LT toxoid
- Clostridium perfringens, type C, strain 578, beta toxoid
- Escherichia coli, fimbrial adhesin F5
- Escherichia coli, fimbrial adhesin F6
- Escherichia coli, fimbrial adhesin F4ac
- Escherichia coli, fimbrial adhesin F4ab

## Product identification

**Medicine name:**

Porcilis ColiClos (--)- Suspension for injection

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**Active substance:**

Escherichia coli, LT toxoid

Clostridium perfringens, type C, strain 578, beta toxoid

Escherichia coli, fimbrial adhesin F5

Escherichia coli, fimbrial adhesin F6

Escherichia coli, fimbrial adhesin F4ac

Escherichia coli, fimbrial adhesin F4ab

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**Target species:**

Pig

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**Route of administration:**

Intramuscular use

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## Product details

### **Active substance and strength:**

Escherichia coli, LT toxoid

Presentation\_strength:  $\geq 10.9 \log_2$  Ab titre Reference:Hse Index:0

Clostridium perfringens, type C, strain 578, beta toxoid

Presentation\_strength:  $\geq 20$  IU Reference:Hse Index:1

Escherichia coli, fimbrial adhesin F5

Presentation\_strength:  $\geq 8.4 \log_2$  Ab titre Reference:Hse Index:2

Escherichia coli, fimbrial adhesin F6

Presentation\_strength:  $\geq 7.8 \log_2$  Ab titre Reference:Hse Index:3

Escherichia coli, fimbrial adhesin F4ac

Presentation\_strength:  $\geq 8.1 \log_2$  Ab titre Reference:Hse Index:4

Escherichia coli, fimbrial adhesin F4ab

Presentation\_strength:  $\geq 9.7 \log_2$  Ab titre Reference:Hse Index:5

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### **Pharmaceutical form:**

Suspension for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Pig**

- Not applicable. 0 day Zero days

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

QI09AB08

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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 , Croatia , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Lithuania , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Vial (glass), Package\_size:1 vial, Content:250 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:1 vial, Content:20 ml

Packaging:Vial (PET), Package\_size:1 vial, Content:250 ml

Packaging:Vial (PET), Package\_size:1 vial, Content:200 ml

Packaging:Vial (PET), Package\_size:1 vial, Content:100 ml

Packaging:Vial (PET), Package\_size:1 vial, Content:50 ml

Packaging:Vial (PET), Package\_size:1 vial, Content:20 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

14/06/2012

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

14/06/2012

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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