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

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

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Escherichia coli, LT toxoid

Presentation_strength:≥10.9 log2 Ab titre Reference:Hse Index:0

Clostridium perfringens, type C, strain 578, beta toxoid

Presentation_strength:≥20 IU Reference:Hse Index:1

Escherichia coli, fimbrial adhesin F5

Presentation_strength:≥8.4 log2 Ab titre Reference:Hse Index:2

Escherichia coli, fimbrial adhesin F6

Presentation_strength:≥7.8 log2 Ab titre Reference:Hse Index:3

Escherichia coli, fimbrial adhesin F4ac

Presentation_strength:≥8.1 log2 Ab titre Reference:Hse Index:4

Escherichia coli, fimbrial adhesin F4ab

Presentation_strength:≥9.7 log2 Ab titre Reference:Hse Index:5

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB08

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)

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)

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

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:
Intervet International B.V.

Marketing authorisation date:
14/06/2012

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:
14/06/2012

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: 8/04/2025
Download

ema-puar-v2011-porciliscoliclos-vra0018g-en.pdf

ema-puar-porcilic-coliclos-v-2011-par-en.pdf