Porcilis Porcoli Diluvac Forte (--) - Suspension for injection

Authorised

- Escherichia coli, fimbrial adhesin F4ac
- Escherichia coli, fimbrial adhesin F5
- Escherichia coli, LT toxoid
- Escherichia coli, fimbrial adhesin F6
- Escherichia coli, fimbrial adhesin F4ab

Product identification

Medicine name:
Porcilis Porcoli Diluvac Forte (--) - Suspension for injection

Active substance:
Escherichia coli, fimbrial adhesin F4ac
Escherichia coli, fimbrial adhesin F5
Escherichia coli, LT toxoid
Escherichia coli, fimbrial adhesin F6
Escherichia coli, fimbrial adhesin F4ab

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

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Escherichia coli, fimbrial adhesin F4ac

Presentation_strength:at least 5.4 log2 Ab titre* Reference:HSE Index:0

Escherichia coli, fimbrial adhesin F5

Presentation_strength:at least 6.8 log2 Ab titre* Reference:HSE Index:1

Escherichia coli, LT toxoid

Presentation_strength:at least 6.8 log2 Ab titre* Reference:HSE Index:2

Escherichia coli, fimbrial adhesin F6

Presentation_strength:at least 7.1 log2 Ab titre* Reference:HSE Index:3

Escherichia coli, fimbrial adhesin F4ab

Presentation_strength:at least 9.0 log2 Ab titre* Reference:HSE Index:4

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:

QI09AB02

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 , Croatia , Denmark , Finland , France , Germany , Hungary , Ireland , Italy , Netherlands , Norway , Poland , Romania , Sweden ,
United Kingdom (Northern Ireland)

Package description:
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
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

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:
29/02/1996

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:
2/05/2002

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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