Ecoporc Shiga (--) - Suspension for injection

Authorised

- Escherichia coli, recombinant Shiga toxin 2e

Product identification

Medicine name:
Ecoporc Shiga (--) - Suspension for injection

Active substance:
Escherichia coli, recombinant Shiga toxin 2e

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Escherichia coli, recombinant Shiga toxin 2e

Presentation_strength:≥ 3.2 x 10^6 ELISA units Reference:HSE Index:0

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:

QI09AB02

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)

Package description:

Packaging:Bottle (PET), Package_size:1 bottle, Content:100 ml

Packaging:Bottle (PET), Package_size:1 bottle, Content:50 ml

Packaging:Bottle (LDPE), Package_size:1 bottle, Content:100 ml

Packaging:Bottle (LDPE), Package_size:1 bottle, Content:50 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:
10/04/2013

Manufacturing sites for batch release:
IDT Biologika GmbH
Ceva-Phylaxia Veterinary Biologicals Co. Ltd

Responsible authority:
European Commission

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

Date of authorisation status change:
10/04/2013

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: 14/01/2026
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