

TEROVAXEC

Suspended

- Clostridium perfringens, type B and C, beta toxoid
- Clostridium perfringens, type B and D, epsilon toxoid
- Clostridium septicum, toxoid

Product identification

Medicine name:

TEROVAXEC

Active substance:

Clostridium perfringens, type B and C, beta toxoid

Clostridium perfringens, type B and D, epsilon toxoid

Clostridium septicum, toxoid

Target species:

Sheep

Goat

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Clostridium perfringens, type B and C, beta toxoid

10.00 Toxic unit(s) / 2.00 millilitre(s)

Clostridium perfringens, type B and D, epsilon toxoid

5.00 Toxic unit(s) / 2.00 millilitre(s)

Clostridium septicum, toxoid
2.50 Toxic unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

• **Sheep**

- Meat and offal. 0 day

• **Goat**

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI03AB

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Suspended

Authorised in:

Spain

Package description:

Available only in Spanish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Aquatreck Animal Health S.L.

Marketing authorisation date:

17/12/1982

Manufacturing sites for batch release:

Cz Veterinaria S.A.

Responsible authority:

(AEMPS)

Authorisation number:

3003 ESP

Date of authorisation status change:

25/12/2021

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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