

Covexin 8 vet. injeksjonsvæske, suspensjon Vaksine mot klostridieinfeksjoner til sau og storfe (inaktivert, adsorbent)

Not
authorised

- Clostridium tetani, toxoid
- Clostridium novyi, toxoid
- Clostridium septicum, toxoid
- Clostridium haemolyticum, toxoid
- Clostridium perfringens, type D, epsilon toxoid
- Clostridium perfringens, type B and C, beta toxoid
- Clostridium chauvoei, cells and toxin, Inactivated

Product identification

Medicine name:

Covexin 8 vet. injeksjonsvæske, suspensjon Vaksine mot klostridieinfeksjoner til sau og storfe (inaktivert, adsorbent)

Active substance:

Clostridium tetani, toxoid

Clostridium novyi, toxoid

Clostridium septicum, toxoid

Clostridium haemolyticum, toxoid

Clostridium perfringens, type D, epsilon toxoid

Clostridium perfringens, type B and C, beta toxoid

Clostridium chauvoei, cells and toxin, Inactivated

Target species:

Sheep

Cattle

Sheep (lamb)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Clostridium tetani, toxoid

1.60 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium novyi, toxoid

2.10 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium septicum, toxoid

2.80 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium haemolyticum, toxoid

11.70 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium perfringens, type D, epsilon toxoid

7.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium perfringens, type B and C, beta toxoid

13.10 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Clostridium chauvoei, cells and toxin, Inactivated

90.00 percentage protection / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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Sheep

- All relevant tissues. 0 day

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Cattle

- All relevant tissues. 0 day

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Sheep (lamb)

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AB01

QI04AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Norway

Package description:

Available only in [Norwegian](#)

Available only in [Norwegian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Zoetis Finland Oy

Marketing authorisation date:

16/06/1999

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:

95-3537

Date of authorisation status change:

27/09/2023

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.