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Rococlos Vet. injektionsvæske, emulsion

Authorised

- Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated
- Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), Inactivated
- Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), Inactivated
- Porcine rotavirus A, strain OSU 6, Inactivated
- Porcine rotavirus A, strain OSU 6, Inactivated
- Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated
- Clostridium perfringens, type C, beta toxoid
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- Clostridium perfringens, type C, beta toxoid

Product identification

Medicine name:

Rococlos Vet. injektionsvæske, emulsion

Active substance:

Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated
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Clostridium perfringens, type C, beta toxoid

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated

1.00 relative potency / 1.00 Dose

Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), Inactivated

1.00 relative potency / 1.00 Dose

Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), Inactivated

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Clostridium perfringens, type C, beta toxoid

1.00 relative potency / 1.00 Dose

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Emulsion for injection

Intramuscular use:

-

- Meat. 0 day

- [illegible]

- Meat. 0 day
- Meat. 0 day
- Meat. 0 day
- Meat. 0 day
- Meat. 0 day
- Meat. 0 day
- Meat. 0 day

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

Available only in Danish
Available only in Danish
Available only in Danish
Available only in Danish
Available only in Danish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Salfarm Danmark A/S

Marketing authorisation date:

19/12/2025

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Danish Medicines Agency

Authorisation number:

72024

Date of authorisation status change:

19/12/2025

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.