

Procactive 300 mg/ml Suspension for injection for cattle, sheep and pigs

Authorised

- Benzylpenicillin procaine monohydrate

Product identification

Medicine name:

Procactive 300 mg/ml Suspension for injection for cattle, sheep and pigs

Active substance:

Benzylpenicillin procaine monohydrate

Target species:

Cattle
Sheep
Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Benzylpenicillin procaine monohydrate
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

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Cattle

- Meat and offal. 6 day Treatment duration 3-5 days.

- Meat and offal. 8 day Treatment duration 6-7 days.

- Milk. 96 hour

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Sheep

- Meat and offal. 4 day Treatment duration 3-5 days.

- Meat and offal. 6 day Treatment duration 6-7 days.

- Milk. 156 hour

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Pig

- Meat and offal. 6 day Treatment duration 3-5 days.

- Meat and offal. 8 day Treatment duration 6-7 days.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Norway

Package description:

Polyethylene terephthalate (PET) colourless bottles (250 ml) with type I bromobutyl rubber stoppers and flip-off caps.Pack size: Carton box with 12 boxes containing 1 bottle of 250 ml

Polyethylene terephthalate (PET) colourless vials (100 ml) with type I bromobutyl rubber stoppers and flip-off caps.Pack size: Carton box with 30 boxes containing 1 vial of 100 ml

Polyethylene terephthalate (PET) colourless vials (100 ml) with type I bromobutyl rubber stoppers and flip-off caps.Pack size: Carton box with 10 boxes containing 1 vial of 100 ml

Polyethylene terephthalate (PET) colourless bottle (250 ml) with type I bromobutyl rubber stoppers and flip-off caps.Pack size: Carton box with 1 bottle of 250 ml

Polyethylene terephthalate (PET) colourless vial (100 ml) with type I bromobutyl rubber stoppers and flip-off caps.Pack size: Carton box with 1 vial of 100 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratorios Syva S.A.

Marketing authorisation date:

26/04/2019

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:

17-11936

Date of authorisation status change:

26/04/2019

Reference member state:

Ireland

Procedure number:

IE/V/0397/001

Concerned member states:

Belgium Bulgaria Denmark Estonia Finland France Germany Greece
Hungary Italy Netherlands Norway Portugal Spain Sweden
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 1/02/2026

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Package Leaflet

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Combined File of all Documents