File downloaded on 2026-11-17
Source URL: https://medicines.health.europa.eu/veterinary/en/600000050229

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:

- **Cattle**
  - Meat and offal. 6 day — Treatment duration 3-5 days.
  - Meat and offal. 8 day — Treatment duration 6-7 days.
  - Milk. 96 hour
- **Sheep**
  - Meat and offal. 4 day — Treatment duration 3-5 days.
  - Meat and offal. 6 day — Treatment duration 6-7 days.
  - Milk. 156 hour
- **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:
Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

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:
10/04/2019

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
57035

Date of authorisation status change:
10/04/2019

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
Updated on: 14/03/2026
Download

Package Leaflet

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

Labelling

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

Combined File of all Documents