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Synulox Lactating Cow Intramammary suspension

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate
- Prednisolone

Product identification

Medicine name:
Synulox Lactating Cow Intramammary suspension

Active substance:
Amoxicillin trihydrate

Potassium clavulanate

Prednisolone

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Amoxicillin trihydrate

230.00 milligram(s) / 1.00 Syringe

Potassium clavulanate
59.56 milligram(s) / 1.00 Syringe

Prednisolone
10.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 7 day
  - Milk. 84 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RV01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
HISTORICAL Low density polyethylene syringes packed in cartons containing 3 syringes.
Low density polyethylene syringes packed in cartons containing 300 syringes.
Low density polyethylene syringes packed in cartons containing 24 syringes.
Low density polyethylene syringes packed in cartons containing 12 syringes.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Animal Health ApS

Marketing authorisation date:
24/11/1997

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
94-638

Date of authorisation status change:
13/12/2021

Reference member state:
Ireland

Procedure number:
IE/V/0605/001

Concerned member states:
Austria Bulgaria Cyprus Czechia France Greece Hungary Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain 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: 8/03/2026
Updated on: 13/03/2026
Download

Package Leaflet

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Labelling

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