Synulox Lactating Cow Intramammary suspension.

Authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Synulox Lactating Cow Intramammary suspension.

Active substance:
Prednisolone

Potassium clavulanate

Amoxicillin trihydrate

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Prednisolone

10.00 milligram(s) / 1.00 Syringe

Potassium clavulanate
59.56 milligram(s) / 1.00 Syringe

Amoxicillin trihydrate
230.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Milk. 60 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RV01

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

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

Marketing authorisation date:
25/07/1996

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1112 ESP

Date of authorisation status change:
25/07/1996

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
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Package Leaflet

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Labelling

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