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# Synulox LC Plus Intramammary suspension for lactating cattle

Authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

## Product identification

**Medicine name:**
Synulox LC Plus Intramammary suspension for lactating cattle

**Active substance:**
Prednisolone

Potassium clavulanate

Amoxicillin trihydrate

**Target species:**
Cattle (dairy cow)

**Route of administration:**
Intramammary use

## Product details

**Active substance and strength:**
Prednisolone

10.00 milligram(s) / 1.00 Applicator

Potassium clavulanate
59.60 milligram(s) / 1.00 Applicator

Amoxicillin trihydrate
229.60 milligram(s) / 1.00 Applicator

**Pharmaceutical form:**
Intramammary suspension

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

- **Cattle (dairy cow)**
  - Meat and offal. 7 day
  - Milk. 84 hour

84 hours. With cows milked twice daily, milk for human consumption may only be taken the 7th milking after the last treatment. Where any other milking routine is followed, milk may be taken for human consumption only after the same period from the last treatment (e.g. with 3 times a day milking, milk may be taken for human consumption at the 11th milking).

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ51RV01

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

**Authorisation status:**
Valid

**Authorised in:**
Germany

**Available in:**
Germany

**Package description:**

(ID4) 900 gram(s): unspecified outer container with 300 Applicator (Low Density PolyEthylene) each with 3 gram(s)
(ID3) 72 gram(s): unspecified outer container with 24 Applicator (Low Density PolyEthylene) each with 3 gram(s)
(ID2) 36 gram(s): unspecified outer container with 12 Applicator (Low Density PolyEthylene) each with 3 gram(s)
(ID1) 9 gram(s): Box (Cardboard) with 3 Applicator (Low Density PolyEthylene) each with 3 gram(s)

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Zoetis Deutschland GmbH

**Marketing authorisation date:**
29/10/2013

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

**Responsible authority:**
Federal Office Of Consumer Protection And Food Safety

**Authorisation number:**
401895.00.00

**Date of authorisation status change:**
19/12/2018

**Reference member state:**
Germany

**Procedure number:**

DE/V/0315/001

**Concerned member states:**
Belgium Luxembourg United Kingdom (Northern Ireland)

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

## Documents

Combined File of all Documents

English (PDF)
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