Clavamox LC Intramammary Suspension for Lactating Cattle

Not authorised

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Synulox LC Plus Intramammary suspension for lactating cattle
Clavamox LC 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:
Expired

Authorised in:
United Kingdom (Northern Ireland)

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 (board) 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 UK Limited

Marketing authorisation date:
17/10/2013

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 42058/4017

Date of authorisation status change:
24/05/2022

Reference member state:
Germany

Procedure number:
DE/V/0315/001

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