

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:

Lithuania

Available in:

Lithuania

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 Belgium

Marketing authorisation date:

18/03/1998

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/98/0667/001-003

Date of authorisation status change:

10/04/2008

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

Combined File of all Documents

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