Synulox Lactating Cow Intramammary suspension.

Authorised

- Prednisolone
- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
Synulox Lactating Cow Intramammary suspension.
Avuloxil, suspensie voor intramammair gebruik voor melkgevende runderen

Active substance:
Prednisolone

Amoxicillin trihydrate

Potassium clavulanate

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Prednisolone
10.00 milligram(s) / 1.00 Syringe

Amoxicillin trihydrate
200.00 milligram(s) / 1.00 Syringe

Potassium clavulanate
59.56 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:
Netherlands

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.
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 B.V.

Marketing authorisation date:
20/11/2000

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 9427

Date of authorisation status change:
25/07/2022

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

Combined File of all Documents

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Summary of Product Characteristics
