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Taneven LC 3 g intramammary suspension for lactating cows

Authorised

- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:
Taneven LC 3 g intramammary suspension for lactating cows

Active substance:
Benzylpenicillin (procaine) monohydrate

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Benzylpenicillin (procaine) monohydrate
3.00 gram(s) / 1.00 Applicator

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Milk. 120 hour
  - Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51CE09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Sweden

Package description:

(ID4) 1600 gram(s): Box with 80 Syringe (High Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID3) 400 gram(s): Box with 20 Syringe (High Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID2) 240 gram(s): Box with 12 Syringe (High Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID1) 200 gram(s): Box with 10 Syringe (High Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID6) 240 gram(s): Box with 12 Syringe (Low Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID8) 1600 gram(s): Box with 80 Syringe (Low Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID5) 200 gram(s): Box with 10 Syringe (Low Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)
(ID7) 400 gram(s): Box with 20 Syringe (Low Density PolyEthylene) each with 20 gram(s), closed with Lid (Low Density PolyEthylene)

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:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Marketing authorisation date:
2/11/2020

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
60298

Date of authorisation status change:
2/11/2020

Reference member state:
Germany

Procedure number:
DE/V/0338/001

Concerned member states:
Austria Bulgaria Hungary Ireland Luxembourg Netherlands Poland Portugal Romania Sweden

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents