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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:

Luxembourg

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:
30/11/2022

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:
V 110/22/11/0001

Date of authorisation status change:
30/11/2022

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

Combined File of all Documents