

# Taneven LC 3 g intramammary suspension for cattle

Authorised

- Benzylpenicillin procaine monohydrate

## Product identification

**Medicine name:**

Taneven LC 3 g intramammary suspension for cattle

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**Active substance:**

Benzylpenicillin procaine monohydrate

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**Target species:**

Cattle

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Benzylpenicillin procaine monohydrate

3.00 gram(s) / 1.00 Applicator

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**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:****Intramammary use:**

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**Cattle**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ51CE09

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**Available in:**

Ireland

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

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**Marketing authorisation date:**

24/07/2020

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**Manufacturing sites for batch release:**

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10660/004/001

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**Date of authorisation status change:**

24/07/2020

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0338/001

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**Concerned member states:**

Austria Bulgaria Hungary Ireland Luxembourg Netherlands Poland Portugal  
Romania Sweden

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents