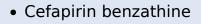
# Cefa Safe 300 mg Intramammary Suspension for dairy cows at drying-off

Authorised



# Product identification

#### Medicine name:

Cefa Safe 300 mg Intramammary Suspension for dairy cows at drying-off Cefa-Safe 300 mg, intramaminė suspensija pieninėms karvėms užtrūkinimo metu

Active substance: Cefapirin benzathine

**Target species:** Cattle (dairy cow at drying-off)

**Route of administration:** Intramammary use

### **Product details**

#### Active substance and strength:

Cefapirin benzathine 383.30 milligram(s) / 1.00 Applicator

#### **Pharmaceutical form:**

Intramammary suspension

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

#### Cattle (dairy cow at drying-off)

- Meat and offal. 14 day
- Milk. 24 hour

24 hours after calving if the interval between treatment and calving is 32 days or longer.

- Milk. 33 day

33 days after treatment if the interval between treatment and calving is less than 32 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51DB08

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### Authorisation status:

Valid

#### Authorised in:

Lithuania

#### Package description:

(ID2) 1440 millilitre(s): Behältnis (Plastic) with 144 Syringe (Low Density PolyEthylene) each with 10 millilitre(s), closed with Lid (Low Density PolyEthylene)(ID1) 200 millilitre(s): Box (Cardboard) with 20 Syringe (Low Density PolyEthylene) each with 10 millilitre(s), closed with Lid (Low Density PolyEthylene)

# Additional information

#### **Entitlement type:** Marketing Authorisation

#### Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

#### Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

7/11/2021

#### Manufacturing sites for batch release:

Intervet International B.V.

#### **Responsible authority:**

State Food And Veterinary Service

#### Authorisation number:

LT/2/21/2688/001-002

#### Date of authorisation status change:

7/11/2021

#### **Reference member state:**

Germany

### Procedure number:

DE/V/0339/001

#### **Concerned member states:**

Austria Belgium Croatia Cyprus Czechia Estonia Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Portugal Romania Slovakia

United Kingdom (Northern Ireland)

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

### Documents

Summary of Product Characteristics

Combined File of all Documents

RV2688.pdf

Source URL: https://medicines.health.europa.eu/veterinary/60000061585