

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

Authorised

- Cefapirin benzathine

Product identification

Medicine name:

Cefa Safe 300 mg Intramammary Suspension for dairy cows at drying-off
CEFA SAFE 300 MG/10ML ΕΝΔΟΜΑΣΤΙΚΟ ΕΝΑΙΩΡΗΜΑ

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:**

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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:

Greece

Package description:

(ID2) 1440 millilitre(s): Behälter (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:

26/12/2022

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

National Organization For Medicines

Authorisation number:

144677/27-12-2022/K-0246701

Date of authorisation status change:

26/12/2022

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
www.adrreports.eu/vet

Documents

Combined File of all Documents

Source URL: <https://medicines.health.europa.eu/veterinary/600000061604>