

TETRA DELTA

Not authorised

- Dihydrostreptomycin sulfate
- NEOMYCIN SULFATE
- Prednisolone
- Benzylpenicillin procaine
- Novobiocin sodium

Product identification

Medicine name:

TETRA DELTA

Active substance:

Dihydrostreptomycin sulfate

NEOMYCIN SULFATE

Prednisolone

Benzylpenicillin procaine

Novobiocin sodium

Target species:

Cattle (dairy cow)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Dihydrostreptomycin sulfate

100.00 milligram(s) / 1.00 Syringe

NEOMYCIN SULFATE

105.00 milligram(s) / 1.00 Syringe

Prednisolone

10.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine

100.00 milligram(s) / 1.00 Syringe

Novobiocin sodium

100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

-

Cattle (dairy cow)

- Meat and offal. 7 day

- Milk. 108 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

This information is not available for this product.

Authorisation status:

Surrendered

Authorised in:

Romania

Package description:

Available only in [Romanian](#)

Available only in [Romanian](#)

Available only in [Romanian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Zoetis Belgium

Marketing authorisation date:

2/05/2005

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

140018

Date of authorisation status change:

27/03/2022

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

Documents

Summary of Product Characteristics

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