Tetra-Delta (105 mg + 100 000 j.m. + 100 mg + 100 mg + 10 mg) /10 ml Zawiesina dowymieniowa

Not authorised

- Prednisolone
- Benzylpenicillin procaine
- Dihydrostreptomycin sulfate
- Neomycin
- Novobiocin sodium

Product identification

Medicine name:
Tetra-Delta (105 mg + 100 000 j.m. + 100 mg + 100 mg + 10 mg) /10 ml Zawiesina dowymieniowa

Active substance:
Prednisolone

Benzylpenicillin procaine

Dihydrostreptomycin sulfate

Neomycin

Novobiocin sodium

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Prednisolone
10.00 milligram(s) / 10.00 millilitre(s)

Benzylpenicillin procaine
100000.00 international unit(s) / 1.00 international unit(s)

Dihydrostreptomycin sulfate
100.00 milligram(s) / 10.00 millilitre(s)

Neomycin
105.00 milligram(s) / 10.00 millilitre(s)

Novobiocin sodium
100.00 milligram(s) / 10.00 millilitre(s)

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 5 week

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RV01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Surrendered

Authorised in:

Poland

Package description:
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Polska Sp. z o.o.

Marketing authorisation date:
12/06/1996

Manufacturing sites for batch release:
Zoetis Belgium
Norbrook Laboratories Limited

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
0259

Date of authorisation status change:
20/04/2022

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

Documents

Package Leaflet

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