TETRANEOMAST F

Authorised

- Prednisolone acetate
- NEOMYCIN SULFATE
- Tetracycline hydrochloride

Product identification

Medicine name:
TETRANEOMAST F

Active substance:
Prednisolone acetate
NEOMYCIN SULFATE
Tetracycline hydrochloride

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Prednisolone acetate
10.00 milligram(s) / 1.00 Syringe
NEOMYCIN SULFATE

250.00 milligram(s) / 1.00 Syringe

Tetracycline hydrochloride
200.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (lactating cow)**
  - Meat and offal. 14 day
  - Milk. 5 day 5 дни (10 издоявания)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RV01

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Farma Vet OOD

Marketing authorisation date:
28/09/2017

Manufacturing sites for batch release:
Farma Vet OOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2760

Date of authorisation status change:
25/01/2022

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

Documents

Summary of Product Characteristics

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Package Leaflet

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