Oxytetraciclin hydrochloride-NGP 7/ 0,340 g, 11/ 0,510 g, 22/ 1,020 g comprettae spumescentes

Authorised

- Oxytetracycline hydrochloride
- Oxytetracycline hydrochloride
- Oxytetracycline hydrochloride

Product identification

Medicine name:
Окситетрациклин хидрохлорид-NGP 7/ 0,340 g, 11/ 0,510 g, 22/ 1,020 g пенообразуващи компрети
Oxytetraciclin hydrochloride-NGP 7/ 0,340 g, 11/ 0,510 g, 22/ 1,020 g comprettae spumescentes

Active substance:
Oxytetracycline hydrochloride
Oxytetracycline hydrochloride
Oxytetracycline hydrochloride

Target species:
Pig (sow for reproduction)
Goat
Sheep

Route of administration:
Intrauterine use

Product details

Active substance and strength:
Oxytetracycline hydrochloride
0.34 gram(s) / 1.00 Tablet

Oxytetracycline hydrochloride
0.51 gram(s) / 1.00 Tablet

Oxytetracycline hydrochloride
1.02 gram(s) / 1.00 Tablet

Pharmaceutical form:
Intrauterine tablet

Withdrawal period by route of administration:
Intrauterine use:

- **Pig (sow for reproduction)**
  - Meat and offal. no withdrawal period

Месо и вътрешни органи - третираните животни не може да се използват за консумация от хора

- **Goat**
  - Milk. no withdrawal period

Мляко - по време на третирането и най-малко 4-5 дни след това животните отделят коластра, която не се консумира от хора

  - Meat and offal. no withdrawal period

Месо и вътрешни органи - третираните животни не може да се използват за консумация от хора

- **Sheep**

- Meat and offal. no withdrawal period

Месо и вътрешни органи - третираните животни не може да се използват за консумация от хора

- Milk. no withdrawal period

Мляко - по време на третирането и най-малко 4-5 дни след това животните отделят коластра, която не се консумира от хора

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Northern Veterinary Dealer-SVD OOD

Marketing authorisation date:
10/06/2007

Manufacturing sites for batch release:
Northern Veterinary Dealer-SVD OOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1649-03-11-2011

Date of authorisation status change:
2/11/2011

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.

Package Leaflet and Labelling

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