NEOCIDOL 600 g/L concentrat emulsionabil pentru bovine, ovine, caprine, suine

Authorised

- Dimpylate

Product identification

Medicine name:
NEOCIDOL 600 g/L concentrat emulsionabil pentru bovine, ovine, caprine, suine

Active substance:
Dimpylate

Target species:
Cattle
Goat
Sheep
Pig

Route of administration:
Dipping
Topical use

Product details

Active substance and strength:

Dimpylate
600.00 gram(s) / 1.00 litre(s)

Pharmaceutical form:

Concentrate for dip solution

Withdrawal period by route of administration:

Dipping:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Goat**
  - Meat and offal. 21 day
  - Milk. 21 day
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 21 day

Topical use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 21 day
- **Goat**

- Meat and offal. 21 day
- Milk. 21 day

- **Pig**
  - Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AF03

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

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
ZAGRO Europe GmbH

Marketing authorisation date:
24/03/1998

Manufacturing sites for batch release:
Denka International B.V.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
110046

Date of authorisation status change:
30/03/2011

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.