BUTOX 50 mg/ml, concentrat pentru soluție cu aplicare cutanată

Not authorised

- Deltamethrin

Product identification

Medicine name:
BUTOX 50 mg/ml, concentrat pentru soluție cu aplicare cutanată

Active substance:
Deltamethrin

Target species:
Cattle
Sheep

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Deltamethrin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for cutaneous solution

Withdrawal period by route of administration:

Cutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 4 day

nu este permisa administrarea la animale producatoare lapte pentru consum uman

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC11

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:
Intervet International B.V.

Marketing authorisation date:
9/08/2006

Manufacturing sites for batch release:
Intervet Productions S.A.

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

Authorisation number:
160392

Date of authorisation status change:
4/06/2024

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.