Butox Protect 7.5 mg/ml pour-on suspension for cattle and sheep

Authorised

- Deltamethrin

Product identification

Medicine name:
Butox Protect 7.5 mg/ml pour-on suspension for cattle and sheep
BUTOX PROTECT 7,5 mg/ml kožni poliv, suspenzija za govedo in ovce

Active substance:
Deltamethrin

Target species:
Cattle
Sheep

Route of administration:
Pour-on use

Product details

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

Pharmaceutical form:

Pour-on suspension

Withdrawal period by route of administration:

Pour-on use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 18 day
- **Sheep**
  - Milk. 12 hour
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC11

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Available in:

Slovenia

Package description:

(ID3): 1 unspecified outer container with 1 Bottle (High Density PolyEthylene) with 2500 millilitre(s) (2500 millilitre(s))
(ID2): 1 unspecified outer container with 1 Bottle (High Density PolyEthylene) with 1000 millilitre(s) (1000 millilitre(s))
(ID1): 1 unspecified outer container with 1 Bottle (High Density PolyEthylene) with 250 millilitre(s) (250 millilitre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
17/03/2010

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0052/001

Date of authorisation status change:
17/03/2010

Reference member state:
Germany

Procedure number:
DE/V/0134/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Denmark Finland Greece Luxembourg Netherlands Slovenia Sweden

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.