CRIMECTIN INJ - BG

Authorised

- Ivermectin

Product identification

Medicine name:
CRIMECTIN INJ - BG

Active substance:
Ivermectin

Target species:
Cattle
Pig
Sheep
Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 49 day

Мляко: не се разрешава за употреба при животни, чието мляко е предназначено за консумация от хора.

- **Pig**
  - Meat and offal. 28 day

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

Мляко: не се разрешава за употреба при животни, чието мляко е предназначено за консумация от хора.

- **Goat**
  - Meat and offal. 28 day

Мляко: не се разрешава за употреба при животни, чието мляко е предназначено за консумация от хора.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

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Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:

Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

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:
Crida Pharm S.R.L.

Marketing authorisation date:
10/12/2014

Manufacturing sites for batch release:
Crida Pharm S.R.L.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2441

Date of authorisation status change:
24/10/2019

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.