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IVOMEC Premix

Not authorised

- Ivermectin

Product identification

Medicine name:

IVOMEC Premix

Active substance:

Ivermectin

Target species:

Pig

Route of administration:

In-feed use

Product details

Active substance and strength:

Ivermectin

0.60 percent weight/weight / 1.00 Bag

Pharmaceutical form:

Premix for medicated feeding stuff

Withdrawal period by route of administration:

In-feed use:

-

Pig

- Meat and offal. 12 day Прасета > 100 kg: 12 дни

- Meat and offal. 3 day Прасета < 100 kg: 3 дни

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

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:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

10/06/2013

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2044

Date of authorisation status change:

23/07/2023

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.