Levaveto 750 mg/g Powder for use in drinking water

Authorised

- Levamisole hydrochloride

Product identification

Medicine name:
Levaveto 750 mg/g Powder for use in drinking water

Active substance:
Levamisole hydrochloride

Target species:
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Levamisole hydrochloride
884.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Pig**

  - Meat and offal. 21 day 21 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

Levaveto 750 mg/g pwdr. for drinking water 10 x 100 g

Levaveto 750 mg/g pwdr. for drinking water 1000 g

Levaveto 750 mg/g pwdr. for drinking water 100 g

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 8(3) of Directive No 2001/83/EC)

Marketing authorisation holder:

V.M.D.

Marketing authorisation date:

14/06/2018

Manufacturing sites for batch release:
V.M.D.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 122516

Date of authorisation status change:
2/02/2022

Reference member state:
Belgium

Procedure number:
BE/V/0034/001

Concerned member states:
Estonia France Germany Hungary Latvia Lithuania Netherlands Poland Romania

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
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