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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 Levaveto, 750 mg/g pulber joogivees manustamiseks sigadele

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:

Estonia

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:

28/06/2018

Manufacturing sites for batch release: V.M.D.
Responsible authority: State Agency Of Medicines
Authorisation number: 2100
Date of authorisation status change: 28/06/2018
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
Summary of Product Characteristics
This document does not exist in this language (English). You can find it in another

language below.

Combined File of all Documents		