Doxatib 500 mg/g powder for use in drinking water for pigs and chickens

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
Doxatib 500 mg/g powder for use in drinking water for pigs and chickens
Doxatib, 500 mg/g, milteliai naudoti su geriamuoju vandeniu, kiaulėms ir vištoms

Active substance:
Doxycycline hyclate

Target species:
Pig
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Doxycycline hyclate
500.00 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. 4 day /

- **Chicken**
  - Meat and offal. 3 day

Following a dose rate of 10 mg/kg body weight for 4 days. Do not use within 4 weeks of onset of the laying period.

- Meat and offal. 9 day

Following a dose rate of 20 mg/kg body weight for 4 days. Do not use within 4 weeks of onset of the laying period.

- Egg. no withdrawal period

Not authorised for use in laying birds producing eggs for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

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

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Alu quadriplex (PET/Al/PET/PE) bags. Pack sizes of 100 g.
Alu quadriplex (PET/Al/PET/PE) bags. Pack sizes of 1 kg.

Alu quadriplex (PET/Al/PET/PE) bags. Pack sizes of 5 kg.
Alu triplex (PET/Al/PE) bags. Pack sizes of 5 kg.
Alu triplex (PET/Al/PE) bags.Pack sizes of 1 kg.
Alu triplex (PET/Al/PE) bags. Pack sizes of 100 g.

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
9/09/2016

Manufacturing sites for batch release:
TAD Pharma GmbH

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/16/2369/001-003

Date of authorisation status change:
16/08/2021

Reference member state:
Slovenia

Procedure number:
SI/V/0001/001

Concerned member states:
Belgium Bulgaria Croatia Czechia Estonia France Germany Hungary Ireland

Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

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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