Doxylin 100%, powder for use in drinking water/milk for calves and pigs

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
Doxylin 100%, powder for use in drinking water/milk for calves and pigs
Doxylin 100% pulveris lietošanai ar dzeramo ūdeni/pienu teļiem un cūkām

Active substance:
Doxycycline hyclate

Target species:
Cattle (calf)
Pig

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
Doxycycline hyclate
1000.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water/milk

Withdrawal period by route of administration:
In drinking water/milk use:

- **Cattle (calf)**
  - Meat and offal. 14 day

- **Pig**
  - Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
White polypropylene container (bucket) of 5 kg, covered with a polypropylene closure.
White polypropylene container (bucket) of 2 kg, covered with a polypropylene closure.
White polypropylene container of 100 grams, covered with a low-density polyethylene closure.
White polypropylene container (bucket) of 1 kg, covered with a polypropylene closure.
White polypropylene container of 1 kg, covered with a low-density polyethylene closure.

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:
Dopharma Research B.V.

Marketing authorisation date:
28/12/2013

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/MRP/13/0053

Date of authorisation status change:
29/12/2013

Reference member state:
Netherlands

Procedure number:
NL/V/0184/001

Concerned member states:
Belgium Denmark Estonia France Germany Greece Hungary Italy Latvia Lithuania Poland Romania

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 19/03/2024
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Package Leaflet

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Labelling

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