

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

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:**

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Cattle (calf)

- Meat and offal. 14 day

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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:

Poland

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:

8/09/2014

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2384

Date of authorisation status change:

8/09/2014

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

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents