

Doxylin 50% WSP

Authorised

- Doxycycline hyclate

Product identification

Medicine name:

Doxylin 50% WSP

Active substance:

Doxycycline hyclate

Target species:

Chicken

Cattle (calf)

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Doxycycline hyclate

500.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for oral solution

Withdrawal period by route of administration:**Oral use:**

-

Chicken

- Meat and offal. 5 day

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

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Cattle (calf)

- Meat and offal. 7 day

Not permitted for use in cattle producing milk for human consumption.

-

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:

Denmark

Available in:

Denmark

Package description:

Bucket: white polypropylene bucket provided with a polypropylene lid. The bucket contains 5 kg of product.

Bucket: white polypropylene bucket provided with a polypropylene lid. The bucket contains 2,5 kg of product.

Bucket: white polypropylene bucket provided with a polypropylene lid. The bucket contains 1 kg of product.

Securitainer: white polypropylene container , covered with a low-density polyethylene lid. The securitainer contains 1 kg of product.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Dopharma Research B.V.

Marketing authorisation date:

8/10/2002

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

Danish Medicines Agency

Authorisation number:

18227

Date of authorisation status change:

8/10/2002

Reference member state:

Netherlands

Procedure number:

NL/V/0171/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Denmark Estonia Finland Greece Hungary

Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal
Romania Slovakia Sweden

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

Documents

Summary of Product Characteristics

English (PDF)

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Package Leaflet and Labelling