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.

- **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

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Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Securitainer: white polypropylene container , covered with a low-density polyethylene lid.The securitainer contains 1 kg of product.
Bucket: white polypropylene bucket provided with a polypropylene lid. The bucket contains 1 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 5 kg of product.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
22/03/2013

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
22/03/2013

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

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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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Package Leaflet and Labelling

English (PDF)
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Labelling

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