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:

Greece

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:
17/09/2015

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
National Organization For Medicines

Authorisation number:
61176/18-09-2015/K-0205202

Date of authorisation status change:
17/09/2015

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

Package Leaflet and Labelling