Soludox 500 mg/g powder for use in drinking water for turkeys

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
Soludox 500 mg/g powder for use in drinking water for turkeys
Soludox 500 mg/g Pulver zum Eingeben über das Trinkwasser für Puten

Active substance:
Doxycycline hyclate

Target species:
Turkey

Route of administration:
In drinking water use

Product details

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

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Turkey**
  - Meat and offal. 12 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
100 g sachet Polyester, polyethylene, aluminium sachets with an inner layer of ionomer (surlyn) packed per 10 in an carton box
1000 g bag: polyethylene terephtalic acid, aluminium, polyamide and an inner layer of polyethylene.
1000 g bag: polyester, polyethylene, aluminium, polyethylene and an inner layer of polyethylene.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Eurovet Animal Health B.V.

Marketing authorisation date:

31/07/2012

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401644.00.00

Date of authorisation status change:
15/11/2017

Reference member state:
Netherlands

Procedure number:
NL/V/0285/001

Concerned member states:
France Germany Hungary Italy Poland United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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