DOXINYL, 500 MG/G POWDER FOR USE IN DRINKING WATER FOR PIGS, CHICKENS AND TURKEYS

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
DOXINYL, 500 MG/G POWDER FOR USE IN DRINKING WATER FOR PIGS, CHICKENS AND TURKEYS
Doxinyl 500 mg/g Pulver zum Eingeben über das Trinkwasser für Schweine, Hühner, Puten

Active substance:
Doxycycline hyclate

Target species:
Turkey
Pig
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Doxycycline hyclate
577.00 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
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption. Do not use within 4 weeks before the start of the laying period.

- **Pig**
  - Meat and offal. 4 day

- **Chicken**
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption. Do not use within 4 weeks before the start of the laying period.

- Meat and offal. 5 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:
Germany

Available in:
Germany

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
15/12/2020

Manufacturing sites for batch release:
aniMedica GmbH
Industria Italiana Integratori Trei S.p.A.
Industrial Veterinaria S.A.
aniMedica Herstellungs GmbH
aniMedica Herstellungs GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402722.00.00

Date of authorisation status change:
15/12/2020

Reference member state:
France

Procedure number:
FR/V/0406/001

Concerned member states:
Austria Czechia Germany Greece Hungary Italy Poland Portugal Romania Slovakia Spain

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

Documents

Combined File of all Documents

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