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

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

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

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:
26/05/2021

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:
National Organization For Medicines

Authorisation number:
42693/27-05-2021/K-0242601

Date of authorisation status change:

26/05/2021

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

eu-puar-frv0406001-mr-rpe603-en.pdf