# 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:**
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

English (PDF)
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