File downloaded on 2026-11-24
Source URL: https://medicines.health.europa.eu/veterinary/en/600000036801

DFV DOXIVET 500 mg/g, powder for use in drinking water for pigs and chickens

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
DFV DOXIVET 500 mg/g, powder for use in drinking water for pigs and chickens

Active substance:
Doxycycline hyclate

Target species:
Chicken
Pig

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:

- **Chicken**
  - Meat and offal. 12 day
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

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

Authorisation status:
Valid

Authorised in:
Norway

Package description:
Bag formed of polyester/aluminium/polyethylene laminate. Bag 1 kg
Bag formed of polyester/aluminium/polyethylene laminate.Carton box 10 x 100 g
Bag formed of polyester/aluminium/polyethylene laminate.Carton box 250 x 100 g
Bag formed of polyester/aluminium/polyethylene laminate.Carton box 50 x 100 g
Bag formed of polyester/aluminium/polyethylene laminate. Bag 2.5 kg

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Divasa Farmavic S.A.

Marketing authorisation date:
1/08/2011

Manufacturing sites for batch release:
Divasa Farmavic S.A.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
10-7848

Date of authorisation status change:
27/06/2016

Reference member state:
Netherlands

Procedure number:
NL/V/0152/002

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Germany Greece Hungary Ireland Italy Lithuania Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 20/06/2022
Updated on: 14/03/2026
Download

Combined File of all Documents

Package Leaflet

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

108007 - par.pdf