

Hypersol 500 mg/g Powder for Use in Drinking Water

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:

HYPERSOL 500 MG/G POWDER FOR USE IN DRINKING WATER

Hypersol 500 mg/g Powder for Use in Drinking Water

Active substance:

Oxytetracycline hydrochloride

Target species:

Chicken

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Oxytetracycline hydrochloride

540.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:**Oral use:****• Chicken**

- Meat and offal. 7 day
- Eggs. no withdrawal period

Do not use in laying birds producing eggs intended for human consumption.

• Pig

- Meat and offal. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

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:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Huvepharma S.A.

Marketing authorisation date:

10/05/2013

Manufacturing sites for batch release:

Huvepharma

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 41623/4000

Date of authorisation status change:

15/08/2022

Reference member state:

France

Procedure number:

FR/V/0251/001

Concerned member states:

Greece Hungary Ireland Italy Poland Portugal Romania Spain

United Kingdom (Northern Ireland)

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000043441>