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:
In drinking water 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:

In drinking water 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

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Authorisation status:
Valid

Authorised in:
Ireland

Package description:
1 kg jar
10 kg bag
5 kg bag
5 kg bucket

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:
31/05/2013

Manufacturing sites for batch release:
Huvepharma S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10453/001/001

Date of authorisation status change:
31/05/2013

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

Documents

Combined File of all Documents
