

# Doxx-Sol 500 mg/g Powder for Use in Drinking Water/Milk Replacer for Pre-ruminant Calves, Pigs and Chickens

Authorised

- Doxycycline hyclate

## Product identification

**Medicine name:**

Doxx-Sol 500 mg/g Powder for Use in Drinking Water/Milk Replacer for Pre-ruminant Calves, Pigs and Chickens

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**Active substance:**

Doxycycline hyclate

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**Target species:**

Chicken

Cattle (calf)

Pig

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**Route of administration:**

In drinking water/milk use

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## Product details

### **Active substance and strength:**

Doxycycline hyclate

500.00 milligram(s) / 1.00 gram(s)

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### **Pharmaceutical form:**

Powder for use in drinking water/milk

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### **Withdrawal period by route of administration:**

#### **In drinking water/milk use:**

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#### **Chicken**

- Meat and offal. 5 day

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#### **Cattle (calf)**

- Meat and offal. 7 day

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#### **Pig**

- Meat and offal. 8 day

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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01AA02

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Valid

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### **Authorised in:**

United Kingdom (Northern Ireland)

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### **Package description:**

Bags of 5 kg formed from polyethylene/aluminium/polyethylene terephthalate laminate.

Bags of 1 kg formed from polyethylene/aluminium/polyethylene terephthalate laminate.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

21/01/2015

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 30282/4022

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**Date of authorisation status change:**

11/05/2024

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0185/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France

Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg  
Malta Poland Portugal Romania Slovakia Slovenia  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents