

# HIDROCOL, 4000000 IU/ML SOLUTION FOR USE IN DRINKING WATER/MILK

Authorised

- COLISTIN SULFATE

## Product identification

**Medicine name:**

HIDROCOL, 4000000 IU/ML SOLUTION FOR USE IN DRINKING WATER/MILK

Hidrocol, 4 000 000 NE/ml oldat ivóvízbe vagy tejbe keveréshez

**Active substance:**

COLISTIN SULFATE

**Target species:**

Turkey

Pig

Sheep (lamb)

Cattle (calf)

Chicken

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

COLISTIN SULFATE

4000000.00 international unit(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for use in drinking water/milk

---

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

•

**Turkey**

- Meat and offal. 1 day
- Eggs. 0 day

•

**Pig**

- Meat and offal. 1 day

•

**Sheep (lamb)**

- Meat and offal. 1 day

•

**Cattle (calf)**

- Meat and offal. 1 day

•

**Chicken**

- Meat and offal. 1 day
- Eggs. 0 day

---

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

QA07AA10

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Hungary

---

**Available in:**

Hungary

---

**Package description:**

1 L white high density polyethylene bottles sealed by induction and closed by a high density polyethylene screw cap.

5 L white high density polyethylene barrels sealed by induction and closed by a high density polyethylene screw cap.

---

## 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:**

S P Veterinaria S.A.

---

**Marketing authorisation date:**

4/07/2016

---

**Manufacturing sites for batch release:**

S P Veterinaria S.A.

---

**Responsible authority:**

Directorate Of Veterinary Medicinal Products

---

**Authorisation number:**

**Date of authorisation status change:**

4/07/2016

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0294/001

---

**Concerned member states:**

Bulgaria Cyprus Greece Hungary Ireland Italy Malta Poland Portugal  
Romania Spain United Kingdom (Northern Ireland)

---

To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)