# Doramax 5 mg/ml Pour-on Solution for Cattle

Authorised

- Doramectin

## Product identification

**Medicine name:**
Doramax 5 mg/ml Pour-on Solution for Cattle

**Active substance:**
Doramectin

**Target species:**
Cattle

**Route of administration:**
Pour-on use

## Product details

**Active substance and strength:**
Doramectin
5.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Pour-on solution

**Withdrawal period by route of administration:**

**Pour-on use:**

- **Cattle**
  - Meat and offal. 35 day

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

QP54AA03

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**

Valid

**Authorised in:**

Germany

**Package description:**

The product will be supplied in:8L (5 L + 3 L) high-density polyethylene bottles with a tamper evident cap in a carton box
The product will be supplied in:6L (5L+1L) high-density polyethylene bottles with a tamper evident cap in a carton box
The product will be supplied in:5 L high-density polyethylene bottles with a tamper evident cap in a carton box
The product will be supplied in:3 L high-density polyethylene bottles with a tamper evident cap in a carton box
The product will be supplied in:2.5 L high-density polyethylene bottles with a tamper evident cap in a carton box
The product will be supplied in:1 L high-density polyethylene bottles with a tamper evident cap in a carton box

## 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:**
Chanelle Pharmaceuticals Manufacturing Limited

**Marketing authorisation date:**
8/09/2022

**Manufacturing sites for batch release:**
Chanelle Pharmaceuticals Manufacturing Limited

**Responsible authority:**
Federal Office Of Consumer Protection And Food Safety

**Authorisation number:**
V7007518.00.00

**Date of authorisation status change:**
8/09/2022

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0669/001

**Concerned member states:**
Belgium Czechia France Germany Hungary Netherlands Poland Romania Slovenia

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

# Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics