

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

Authorised

- Doramectin

## Product identification

**Medicine name:**

Taurador 5 mg/ml Pour-on Solution for Cattle

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**Active substance:**

Doramectin

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**Target species:**

Cattle

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**Route of administration:**

Pour-on use

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## Product details

**Active substance and strength:**

Doramectin

5.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Pour-on solution

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**Withdrawal period by route of administration:****Pour-on use:**

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**Cattle**

- Meat and offal. 35 day

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

QP54AA03

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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:**

Ireland

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**Available in:**

Ireland

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**Package description:**

The veterinary product will be supplied in:20L white high density polyethylene Jerry cans with high density polyethylene caps.

The veterinary product will be supplied in:10L white high density polyethylene Jerry cans with high density polyethylene caps.

The veterinary product will be supplied in:5L white flat bottomed heavy duty high density polyethylene back-packs with 38mm white polypropylene easy peel caps.

The veterinary product will be supplied in:2.5L white flat bottomed heavy duty high density polyethylene back-packs with 38mm white polypropylene easy peel caps.

The veterinary product will be supplied in:1L white flat bottomed heavy duty high density polyethylene back-packs with 38mm white polypropylene easy peel caps.

The veterinary product will be supplied in:1L standard high density polyethylene bottles with 28mm polypropylene/high density polyethylene caps.

The veterinary product will be supplied in:250mL standard high density polyethylene bottles with 28mm polypropylene/high density polyethylene caps.

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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:**

Norbrook Laboratories (Ireland) Limited

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**Marketing authorisation date:**

29/11/2013

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**Manufacturing sites for batch release:**

Norbrook Laboratories Limited

Norbrook Laboratories (Ireland) Limited

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA22664/111/001

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**Date of authorisation status change:**

29/11/2013

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0617/001

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**Concerned member states:**

Austria Belgium France Luxembourg Netherlands Portugal Spain

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

Summary of Product Characteristics

Combined File of all Documents