

# Cyductin TriclaMox 1 mg/ml + 50 mg/ml Oral Solution for Sheep

Authorised

- Triclabendazole
- Moxidectin

## Product identification

**Medicine name:**

CYDECTIN TRICLAMOX 1 MG/ML + 50 MG/ML ORAL SOLUTION FOR SHEEP

Cyductin TriclaMox 1 mg/ml + 50 mg/ml Oral Solution for Sheep

**Active substance:**

Triclabendazole

Moxidectin

**Target species:**

Sheep

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Triclabendazole

50.00 milligram(s) / 1.00 millilitre(s)

Moxidectin

1.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral solution

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**Withdrawal period by route of administration:**

**Oral use:**

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**Sheep**

- Milk. no withdrawal period

Not authorised for use in ewes producing milk intended for human consumption including during the dry period. Do not use within 1 year prior to the first lambing in ewes intended to produce milk for human consumption.

- Meat and offal. 31 day

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

QP54AB52

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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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**Available in:**

United Kingdom (Northern Ireland)

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**Package description:**

1 L bottle

5 L bottle

2.5 L bottle

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Fixed combination application (Article 13b of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Belgium S.A.

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**Marketing authorisation date:**

18/12/2009

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**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 60021/3058

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**Date of authorisation status change:**

6/07/2024

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**Reference member state:**

France

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**Procedure number:**

FR/V/0201/001

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**Concerned member states:**

Austria Belgium Germany Iceland Ireland Italy Luxembourg Netherlands  
Portugal Spain Sweden 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)