CYDECTIN 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

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)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:
Oral use:

- **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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AB52

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
1 L bottle
5 L bottle
2.5 L bottle

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Portugal Lda.

Marketing authorisation date:
20/10/2009

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain, S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
206/01/09DFVPT

Date of authorisation status change:
29/08/2022

Reference member state:
France

Procedure number:
FR/V/0201/001

Concerned member states:
Austria Belgium Germany Iceland Ireland Italy Luxembourg Netherlands Portugal Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
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