Moxodex 1 mg/ml oral solution for sheep

Authorised

- Moxidectin

Product identification

Medicine name:
Moxodex 1 mg/ml oral solution for sheep
Moxidectin Chanelle 1 mg/ml Lösung zum Eingeben für Schafe

Active substance:
Moxidectin

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Moxidectin
1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Sheep**
  - Meat and offal. 14 day
  - Milk. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Package description:

White HDPE flexi containers containing 1L of product. The containers are closed with an aluminium foil seal and polypropylene tamper-evident caps.The product is marketed in a cardboard outer carton.
White HDPE flexi containers containing 2.5 L of product. The containers are closed with an aluminium foil seal and polypropylene tamper-evident caps.The product is marketed in a cardboard outer carton.
White HDPE flexi containers containing 3 L of product. The containers are closed with an aluminium foil seal and polypropylene tamper-evident caps.The product is marketed in a cardboard outer carton.
White HDPE flexi containers containing 5 L of product. The containers are closed with an aluminium foil seal and polypropylene tamper-evident caps.The product is marketed in a cardboard outer carton.

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:
12/08/2020

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402524.00.00

Date of authorisation status change:
12/08/2020

Reference member state:
Ireland

Procedure number:
IE/V/0404/001

Concerned member states:
Austria Belgium France Germany Italy Luxembourg Netherlands 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)
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