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Dilaterol 25 micrograms/ml syrup for horses

Authorised

- Clenbuterol hydrochloride

Product identification

Medicine name:
Dilaterol 25 micrograms/ml syrup for horses

Active substance:
Clenbuterol hydrochloride

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Clenbuterol hydrochloride
25.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Syrup

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR03CC13

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

Authorisation status:
Valid

Authorised in:
Denmark

Package description:
355 ml HDPE bottle sealed with an aluminium/PE heat seal. The product is supplied in a carton box with a multi-component mechanical pump dispenser capable of delivering 4 ml of the product
355 ml HDPE bottle sealed with a transparent HDPE cap. The product is supplied in a carton box with a multi-component mechanical pump dispenser capable of delivering 4 ml of the product.

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:
Le Vet. Beheer B.V.

Marketing authorisation date:
21/02/2013

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
49677

Date of authorisation status change:
21/02/2013

Reference member state:
Ireland

Procedure number:
IE/V/0619/001

Concerned member states:
Austria Belgium Czechia Denmark Finland France Hungary Iceland Italy Luxembourg Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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