VENTIPULMIN SOLUTION INJECTABLE

Authorised

- Clenbuterol hydrochloride

Product identification

Medicine name:
VENTIPULMIN SOLUTION INJECTABLE

Active substance:
Clenbuterol hydrochloride

Target species:
Horse (mare)
Horse

Route of administration:
Intravenous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse (mare)**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR03CC13

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

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Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
14/02/1986

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/0486110 8/1986

Date of authorisation status change:
14/02/2011

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.