

Ventipulmin Solution for Injection

30 micrograms/ml

Authorised

- Clenbuterol hydrochloride

Product identification

Medicine name:

Ventipulmin Solution for Injection 30 micrograms/ml

Active substance:

Clenbuterol hydrochloride

Target species:

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

- 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:

Ireland

Package description:

50 ml amber glass injection vial (Ph. Eur. Type II), with pink bromobutyl rubberstopper and aluminium crimp cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

1/10/1988

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10454/019/001

Date of authorisation status change:

1/10/1988

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

Documents

Summary of Product Characteristics