

Ventipulmin 0.016 mg/g Granulaat

Authorised

- Clenbuterol hydrochloride

Product identification

Medicine name:

Ventipulmin 0.016 mg/g Granulaat

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Active substance:

Clenbuterol hydrochloride

Target species:

Horse

Route of administration:

In-feed use

Product details

Active substance and strength:

Clenbuterol hydrochloride

0.02 gram(s) / 1.00 gram(s)

Pharmaceutical form:

Granules

Withdrawal period by route of administration:**In-feed use:**

-

Horse

- Milk. no withdrawal period

Do not use with animals producing milk for human consumption

- Meat and offal. no withdrawal period

Do not use in animals aimed at human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QR03CC13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

Ventipulmin 500 g Bottle Granules

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:

24/03/1983

Manufacturing sites for batch release:

Klocke Pharma-Service GmbH

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V122543

Date of authorisation status change:

4/12/2020

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

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Labelling

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