Ventipulmin granulat; 0,016 mg/g granulat dla koni

Authorised

- Clenbuterol hydrochloride

Product identification

Medicine name:
Ventipulmin granulat; 0,016 mg/g granulat dla koni

Active substance:
Clenbuterol hydrochloride

Target species:
Horse

Route of administration:
In-feed use

Product details

Active substance and strength:
Clenbuterol hydrochloride
0.02 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Granules

Withdrawal period by route of administration:

In-feed 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:
Poland

Available in:
Poland

Package description:
Available only in Polish

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:
14/11/2001

Manufacturing sites for batch release:
Klocke Pharma-Service GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1206

Date of authorisation status change:
14/11/2001

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

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