Vulketan 2.5 mg/g Gel

Authorised

- Ketanserin tartrate

Product identification

Medicine name:
Vulketan 2.5 mg/g Gel
Vulketan 2.5 mg/g Gel
Vulketan 2.5 mg/g Gel

Active substance:
Ketanserin tartrate

Target species:
Horse

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Ketanserin tartrate
3.45 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Gel

Withdrawal period by route of administration:

Cutaneous use:

- **Horse**
  - Meat and offal. no withdrawal period 0 days
  - Milk. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD03AX90

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Vulketan Tube with 75 g Gel

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:
Audevard

Marketing authorisation date:
9/09/1991

Manufacturing sites for batch release:
Sanochemia Pharmazeutika AG

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V155242

Date of authorisation status change:
18/12/2019

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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