Vulketan 2.5 mg/g Gel

Authorised

- Ketanserin tartrate

Product identification

Medicine name:
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. 0 day
  - Milk. 0 day

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

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Summary of Product Characteristics

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Labelling

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