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Vulketan 2.5 mg/g Gel

- Ketanserin tartrate

Authorised

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

Product information

Package Leaflet

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