Vulketan gél

Authorised

- Ketanserin tartrate

Product identification

Medicine name:
Vulketan gél

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD03AX90

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

Authorisation status:
Valid

Authorised in:
Hungary

Package description:
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Janssen Pharmaceutica

Marketing authorisation date:
8/06/2000

Manufacturing sites for batch release:
Janssen Pharmaceutica

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
8/06/2000

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

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