SPUTOLOSIN

Authorised

- Dembrexine hydrochloride monohydrate

Product identification

Medicine name:
SPUTOLOSIN

Active substance:
Dembrexine hydrochloride monohydrate

Target species:
Horse (mare)
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Dembrexine hydrochloride monohydrate
5.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:
Oral use:

- **Horse (mare)**
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR05CB90

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
1/10/1990

Manufacturing sites for batch release:

Klocke Pharma-Service GmbH

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/3226426 5/1990

Date of authorisation status change:
1/10/2010

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

Documents

Summary of Product Characteristics

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