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ACTI-STREPTO

Authorised

- Dihydrostreptomycin sulfate

Product identification

Medicine name:
ACTI-STREPTO

Active substance:
Dihydrostreptomycin sulfate

Target species:
Cattle
Pig
Rabbit
Cattle (calf)

Route of administration:
Oral use

Product details

Active substance and strength:
Dihydrostreptomycin sulfate
626.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Milk. no withdrawal period

La spécialité n'est pas destinée aux femelles laitières productrices de lait de consommation.

- **Pig**
  - Meat and offal. 14 day

- **Rabbit**
  - Meat and offal. 14 day

- **Cattle (calf)**
  - Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA07AA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in French
Available only in French
Available only in French
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:
Laboratoires Biove

Marketing authorisation date:
24/07/1992

Manufacturing sites for batch release:
Laboratoires Biove

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

Authorisation number:
FR/V/2441413 9/1992

Date of authorisation status change:
24/07/2012

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

Documents

Summary of Product Characteristics

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