SPECTAM SOLUTION INJECTABLE

Authorised

- Spectinomycin dihydrochloride pentahydrate

Product identification

Medicine name:
SPECTAM SOLUTION INJECTABLE

Active substance:
Spectinomycin dihydrochloride pentahydrate

Target species:
Cattle
Pig
Horse
Horse (mare)
Sheep
Goat
Poultry

Route of administration:
Intramuscular use
Subcutaneous use
Endosinusial use

Product details

Active substance and strength:

Spectinomycin dihydrochloride pentahydrate
116.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
    - Meat and offal. 30 day
    - Milk. 3 day
- **Pig**
    - Meat and offal. 30 day
- **Horse**
    - Meat and offal. 30 day
- **Horse (mare)**
    - Milk. 3 day
- **Sheep**
    - Meat and offal. 30 day
    - Milk. 3 day
- **Goat**
    - Meat and offal. 30 day
    - Milk. 3 day
- **Poultry**

- Eggs. no withdrawal period

En l'absence de LMR pour les œufs, ne pas utiliser chez les espèces pondeuses productrices d'œufs de consommation, 4 semaines avant le démarrage de la ponte et pendant celle-ci.

- Meat and offal. 30 day

Subcutaneous use:

- **Poultry**
  - Meat and offal. 30 day
  - Eggs. no withdrawal period

En l'absence de LMR pour les œufs, ne pas utiliser chez les espèces pondeuses productrices d'œufs de consommation, 4 semaines avant le démarrage de la ponte et pendant celle-ci.

Endosinusial use:

- **Poultry**
  - Meat and offal. 30 day
  - Eggs. no withdrawal period

En l'absence de LMR pour les œufs, ne pas utiliser chez les espèces pondeuses productrices d'œufs de consommation, 4 semaines avant le démarrage de la ponte et pendant celle-ci.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01XX04

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

---

Authorisation status:
Valid

Authorised in:
France

Package description:

Available only in French

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:

Ceva Sante Animale

Marketing authorisation date:

24/07/1992

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1512729 5/1992

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

This document does not exist in this language (English). You can find it in another language below.