SEPONVER

Authorised

- Closantel sodium dihydrate

Product identification

Medicine name:
SEPONVER

Active substance:
Closantel sodium dihydrate

Target species:
Cattle
Sheep (ewe lamb)
Sheep
Sheep (ewe)

Route of administration:
Oral use

Product details

Active substance and strength:
Closantel sodium dihydrate
54.36 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
Oral use:

- **Cattle**
  - Meat and offal. 55 day
  - Milk. no withdrawal period

Ne pas administrer aux bovins producteurs de lait pour la consommation humaine y compris durant la période de tarissement. Ne pas utiliser durant la seconde moitié de la gestation chez les génisses qui sont destinées à la production de lait pour la consommation humaine.

- **Sheep (ewe lamb)**
  - Milk. 34 day

34 jours après l’agnelage si la période entre le traitement et l’agnelage est d’au moins 90 jours.

  - Milk. 120 day

4 mois après le traitement si la période entre le traitement et l’agnelage est inférieure à 90 jours.

- **Sheep**
  - Meat and offal. 55 day

- **Sheep (ewe)**
  - Milk. 34 day

34 jours après l’agnelage si la période sèche est d’au moins 90 jours.

  - Milk. 120 day

4 mois après le traitement si la période sèche est inférieure à 90 jours.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AG09

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

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
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:
Elanco GmbH

Marketing authorisation date:
27/09/1989

Manufacturing sites for batch release:
Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.
Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.

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

Authorisation number:

FR/V/5195635 8/1989

Date of authorisation status change:
27/09/2009

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

Documents

Summary of Product Characteristics

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