PREMELANGE MEDICAMENTEUX Z 30

Authorised

- Trimethoprim
- Sulfadiazine

Product identification

Medicine name:
PREMELANGE MEDICAMENTEUX Z 30

Active substance:
Trimethoprim
Sulfadiazine

Target species:
Cattle
Pig
Sheep (lamb)
Sheep
Cattle (calf)
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
Trimethoprim
12.50 milligram(s) / 1.00 gram(s)

Sulfadiazine
62.50 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:
Oral use:

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

Voir rubrique « 3.1 - Espèce-cible », ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

- **Pig**
  - Meat and offal. 12 day

- **Sheep (lamb)**
  - Meat and offal. 12 day

- **Sheep**
  - Milk. no withdrawal period

Voir rubrique « 3.1 - Espèce-cible », ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine.

- **Cattle (calf)**

- Meat and offal. 12 day

- **Poultry**
  - Meat and offal. 12 day
  - Eggs. no withdrawal period

En l'absence de LMR pour les œufs, ne pas utiliser chez les oiseaux pondeurs d’œufs ou destinés à la ponte d’œufs pour la consommation humaine, 4 semaines avant le démarrage de la ponte et pendant celle-ci.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW10

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

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:

Huvepharma S.A.

Marketing authorisation date:
6/08/1992

Manufacturing sites for batch release:
Huvepharma S.A.

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

Authorisation number:
FR/V/3585483 8/1992

Date of authorisation status change:
6/08/2012

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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