ULTRADIAZINE

Authorised

- Sulfadiazine sodium
- Trimethoprim

Product identification

Medicine name:
ULTRADIAZINE

Active substance:
Sulfadiazine sodium
Trimethoprim

Target species:
Cattle
Sheep
Goat
Pig (for fattening)
Pig (piglet)

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Sulfadiazine sodium
217.58 milligram(s) / 1.00 millilitre(s)

Trimethoprim
40.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. no withdrawal period — Carne: 6 días (IM). 15 días (IV)
- **Sheep**
  - Meat and offal. no withdrawal period — Carne: 3 días (IM). 15 días (IV)
- **Goat**
  - Meat and offal. no withdrawal period — Carne: 5 días (IM). 15 días (IV)
- **Pig (for fattening)**
  - Meat and offal. no withdrawal period — Carne: 7 días (IM). 15 días (IV)
- **Pig (piglet)**
  - Meat and offal. no withdrawal period — Carne: 7 días (IM). 15 días (IV)
- **Sheep**
  - Milk. no withdrawal period — Leche: 60 horas (2,5 días)
- **Goat**
  - Milk. no withdrawal period — Leche: 60 horas (2,5 días)

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period: Carne: 6 días (IM). 15 días (IV)
- **Sheep**
  - Meat and offal. no withdrawal period: Carne: 3 días (IM). 15 días (IV)
- **Goat**
  - Meat and offal. no withdrawal period: Carne: 5 días (IM). 15 días (IV)
- **Pig (for fattening)**
  - Meat and offal. no withdrawal period: Carne: 7 días (IM). 15 días (IV)
- **Pig (piglet)**
  - Meat and offal. no withdrawal period: Carne: 7 días (IM). 15 días (IV)
- **Cattle**
  - Milk. no withdrawal period: Leche (exclusivamente IV): 48 horas (2 días)
- **Sheep**
  - Milk. no withdrawal period: Leche: 60 horas (2,5 días)
- **Goat**
  - Milk. no withdrawal period: Leche: 60 horas (2,5 días)

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EW10

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
S P Veterinaria S.A.

Marketing authorisation date:
12/01/1994

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
857 ESP

Date of authorisation status change:

12/01/1994

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

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

Labelling

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