GESTAVET OXYTOCIN 10 IU/ml Synthetic Oxytocin, solution for injection

Authorised

- OXYTOCIN SYNTHETIC

Product identification

Medicine name:
HIPRACIN 10IU/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ
GESTAVET OXYTOCIN 10 IU/ml Synthetic Oxytocin, solution for injection

Active substance:
OXYTOCIN SYNTHETIC

Target species:
Cattle
Dog
Sheep
Cat
Pig

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:

OXYTOCIN SYNTHETIC
10.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 12 hour
  - Milk. 12 hour
- **Dog**
- **Sheep**
  - Meat and offal. 12 hour
  - Milk. 12 hour
- **Cat**
- **Pig**
  - Meat and offal. 12 hour

Intravenous use:

- **Cattle**
  - Meat and offal. 12 hour
  - Milk. 12 hour
- **Dog**
- **Sheep**
  - Meat and offal. 12 hour
  - Milk. 12 hour
- **Cat**
- **Pig**
  - Meat and offal. 12 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

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

Authorisation status:
Valid

Authorised in:
Greece

Package description:
The veterinary medicinal product is bottled in sterile 10 ml, colourless Type I glass vial, closed with Type II basic polymeric elastomer closure with anodised aluminium cap. Two 10 ml vials are available in a cardboard box.
The veterinary medicinal product is bottled in sterile 50 ml colourless Type II glass vial, closed with Type II basic polymeric elastomer closure with anodised aluminium cap. One vial of 50 ml is available in a cardboard box.
The veterinary medicinal product is bottled in sterile 10 ml, colourless Type I glass vial closed with Type II basic polymeric elastomer closure with anodised aluminium cap. Clinical presentation: 25 x 10
The veterinary medicinal product is bottled in sterile 10 ml, colourless Type I glass vial closed with Type II basic polymeric elastomer closure with anodised aluminium cap. Clinical presentation: 20 x 10

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Biogenesis Global S.L.

Marketing authorisation date:
24/11/2002

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:

72709/08-11-2007/K-0144701

Date of authorisation status change:
16/08/2023

Reference member state:
Ireland

Procedure number:
IE/V/0134/001

Concerned member states:
Austria Greece Spain

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

Documents

Summary of Product Characteristics
