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LongActon 0.07 mg/ml solution for injection for cattle and pigs

Authorised

- Carbetocin

Product identification

Medicine name:
LongActon 0.07 mg/ml solution for injection for cattle and pigs

Active substance:
Carbetocin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Carbetocin
0.07 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Intravenous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01BB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:
(ID3) 60 millilitre(s): Box (Cardboard) with 6 Vial (Glass) each with 10 millilitre(s), closed with Cap (Aluminium) and Stopper (Rubber)
(ID2) 600 millilitre(s): Box (Cardboard) with 12 Vial (Glass) each with 50 millilitre(s), closed with Cap (Aluminium) and Stopper (Rubber)
(ID1) 50 millilitre(s): Box (Cardboard) with 1 Vial (Glass) with 50 millilitre(s), closed with Cap (Aluminium) and Stopper (Rubber)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
9/12/2002

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

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

Authorisation number:
FR/V/2864058 3/2002

Date of authorisation status change:
9/12/2012

Reference member state:
Germany

Procedure number:
DE/V/0106/001

Concerned member states:
Austria Belgium France Ireland Luxembourg Netherlands Spain United Kingdom (Northern Ireland)

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.

Package Leaflet and Labelling

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

Combined File of all Documents