

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

LongActon 0,07 mg/ml Injektionslösung für Rinder und Schweine

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:

Germany

Available in:

Germany

Package description:

(ID3) 60 millilitre(s): unspecified outer container with 6 Vial (glass) each with 10 millilitre(s)

(ID2) 600 millilitre(s): unspecified outer container with 12 Vial (glass) each with 50 millilitre(s)

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s)

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:

23/02/2000

Manufacturing sites for batch release:

Wirtschaftsgenossenschaft deutscher Tieraerzte e G

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

400323.00.00

Date of authorisation status change:

12/05/2010

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

Combined File of all Documents

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