WELLICOX 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, HORSES, PIGS

Authorised

- Flunixin meglumine

Product identification

Medicine name:
WELLICOX 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, HORSES, PIGS
WELLICOX 50 MG/ML SOLUTION INJECTABLE POUR BOVINS, CHEVAUX, PORCINS

Active substance:
Flunixin meglumine

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:

Flunixin meglumine
83.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 31 day
  - Milk. 36 hour
- **Pig**
  - Meat and offal. 24 day

Intravenous use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 5 day
  - Milk. no withdrawal period

Not authorised for use in mares producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

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
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
7/02/2013

Manufacturing sites for batch release:
Ceva Sante Animale
Vetem S.p.A.

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

Authorisation number:
FR/V/6664786 1/2013

Date of authorisation status change:
17/01/2018

Reference member state:
France

Procedure number:
FR/V/0241/001

Concerned member states:
Belgium Bulgaria Czechia Denmark Germany Hungary Italy Netherlands Poland Portugal Romania Sweden 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.