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 solutie injectabila pentru bovine, porcine, cabaline

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:
Romania

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 Romania S.R.L.

Marketing authorisation date:
8/08/2017

Manufacturing sites for batch release:
Ceva Sante Animale
Vetem SPA

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
180037

Date of authorisation status change:
21/08/2025

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

Combined File of all Documents

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