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

- Flunixin meglumine

Authorised

# Product identification

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

- Belgium

Available in:

- Belgium

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:

- 19/11/2020

Manufacturing sites for batch release:

- Ceva Sante Animale
- Vetem S.p.A.

Responsible authority:

- Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.
Date of authorisation status change:

- 19/11/2020

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

## Product information

Summary of Product Characteristics

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