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

Authorised

This information is not available for this product.

Product identification

Medicine name:
WELLICOX 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND HORSES
Wellicox 50 mg/ml Oplossing voor injectie
Wellicox 50 mg/ml Solution injectable
Wellicox 50 mg/ml Injektionslösung

Active substance:
This information is not available for this product.

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
This information is not available for this product.

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. 20 day
- **Horse**
  - Meat and offal. 10 day
  - Milk. no withdrawal period

Not authorised for use in lactating animals producing milk for human consumption.

Intravenous use:

- **Cattle**
  - Meat and offal. 10 day
  - Milk. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised 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

Summary of Product Characteristics

English (PDF)
Published on: 15/02/2024
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Package Leaflet

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