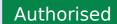
WELLICOX 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND HORSES



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:

OM01AG90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

Available only in French

Available only in French

Available only in $\underline{\mathsf{French}}$

Available only in $\underline{\mathsf{French}}$

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.

Source URL: https://medicines.health.europa.eu/veterinary/600000029098