

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

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:

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Cattle

- Meat and offal. 31 day

- Milk. 36 hour

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Pig

- Meat and offal. 24 day

Intravenous use:

•

Cattle

- Meat and offal. 4 day

- Milk. 24 hour

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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 S.p.A.

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

English (PDF)

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