

Allewinix 50 mg/ml Solution for Injection for Cattle, Pigs and Horses

Authorised

- Flunixin meglumine

Product identification

Medicine name:

Allewinix 50 mg/ml Solution for Injection for Cattle, Pigs and Horses

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:

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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:

United Kingdom (Northern Ireland)

Package description:

Available only in [French](#)

Available only in [French](#)

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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:

16/04/2013

Manufacturing sites for batch release:

Ceva Sante Animale

Vetem S.p.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 14966/3000

Date of authorisation status change:

22/06/2024

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