

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

Authorised

This information is not available for this product.

## Product identification

### Medicine name:

WELICOX 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND HORSES  
Allevinix 50 mg/ml Solution for Injection for Cattle, Pigs and Horses

### 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

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AG90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

United Kingdom (Northern Ireland)

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Animal Health Limited

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**Marketing authorisation date:**

16/04/2013

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**Manufacturing sites for batch release:**

Ceva Sante Animale

Vetem S.p.A.

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**Responsible authority:**

European Medicines Agency

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**Authorisation number:**

Vm 15052/4144

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**Date of authorisation status change:**

15/08/2022

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**Reference member state:**

France

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**Procedure number:**

FR/V/0241/001

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**Concerned member states:**

Belgium Bulgaria Czechia Denmark Germany Hungary Italy Netherlands  
Poland Portugal Romania Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

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