

# Synulox Ready To Use Injection

Not authorised

- Potassium clavulanate
- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Synulox Ready To Use Injection

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**Active substance:**

Potassium clavulanate

Amoxicillin trihydrate

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**Target species:**

Cattle

Dog

Cat

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Potassium clavulanate

64.21 milligram(s) / 1.00 millilitre(s)

Amoxicillin trihydrate

160.70 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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

QJ01CR02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Ireland

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**Package description:**

Type III glass vials of 100 ml containing an off-white sterile, non-aqueous suspension. The vials are sealed with a rubber bung and an aluminium seal and packed in cardboard containers, 6 x 100ml. Each vial contains a sterile off-white to pale buff coloured, smooth fluid, readily dispersable suspension.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Belgium S.A.

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**Marketing authorisation date:**

2/05/2014

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**Manufacturing sites for batch release:**

Haupt Pharma Latina S.r.l.

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10387/075/002

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**Date of authorisation status change:**

28/07/2025

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

## Documents

Summary of Product Characteristics