

Synergal Inj. (140 mg + 35 mg)/ml Zawiesina do wstrzykiwań

Not authorised

- Amoxicillin
- Potassium clavulanate

Product identification

Medicine name:

Synergal Inj. (140 mg + 35 mg)/ml Zawiesina do wstrzykiwań

Active substance:

Amoxicillin

Potassium clavulanate

Target species:

Cat

Dog

Cattle

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Amoxicillin

140.00 milligram(s) / 1.00 millilitre(s)

Potassium clavulanate

35.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Cattle

- Milk. 36 hour
- Meat and offal. 42 day

Subcutaneous use:

-

Cattle

- Milk. 36 hour
- Meat and offal. 42 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CR02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Poland

Package description:

Available only in Polish

Available only in Polish

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:

Scanvet Poland Sp. z o.o.

Marketing authorisation date:

29/04/2004

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

1524

Date of authorisation status change:

20/06/2025

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet

Documents

Package Leaflet

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Labelling

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.