

Amoxicillin 20% LA pro inj.

Authorised

- Amoxicillin

Product identification

Medicine name:

Amoxicillin 20% LA pro inj.

Active substance:

Amoxicillin

Target species:

Cattle (calf)

Cattle

Pig

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Amoxicillin

200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

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

-

Cattle (calf)

- Meat and offal. 14 day

-

Cattle

- Meat and offal. 14 day

- Milk. 4 day

-

Pig

- Meat and offal. 14 day

Subcutaneous use:

-

Cattle (calf)

- Meat and offal. 14 day

-

Cattle

- Meat and offal. 14 day

- Milk. 4 day

-

Pig

- Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Available in:

Bulgaria

Package description:

Available only in Bulgarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Alfasan International B.V.

Marketing authorisation date:

4/11/2004

Manufacturing sites for batch release:

Alfasan International B.V.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-1438

Date of authorisation status change:

4/11/2004

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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