# Profexx 50 mg/ml solution for injection for cattle

Authorised

- Carprofen
- Carprofen

## Product identification

**Medicine name:**
Profexx 50 mg/ml solution for injection for cattle
Profexx 50 mg/ml инжекционен разтвор за говеда

**Active substance:**
Carprofen
Carprofen

**Target species:**
Cattle
Cattle

**Route of administration:**
Subcutaneous use
Intravenous use

## Product details

**Active substance and strength:**
Carprofen

50.00 milligram(s) / 1.00 millilitre(s)

Carprofen
50.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **Cattle**
  - Meat and offal. 21 day
  - Milk. no withdrawal period zero hours

**Intravenous use:**

- **Cattle**
  - Meat and offal. 21 day
  - Milk. no withdrawal period zero hours

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QM01AE91

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Bulgaria

**Package description:**
Cardboard box containing one clear glass (type II) vial of 100 ml with a grey bromobutyl rubber stopper and aluminium cap

Cardboard box containing one clear glass (type II) vial of 250 ml with a grey bromobutyl rubber stopper and aluminium cap
Cardboard box containing one clear glass (type II) vial of 50 ml with a grey bromobutyl rubber stopper and aluminium cap

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 18 of Regulation (EU) 2019/6)

**Marketing authorisation holder:**
Alfasan Nederland B.V.

**Marketing authorisation date:**
13/02/2024

**Manufacturing sites for batch release:**
Alfasan Nederland B.V.

**Responsible authority:**
Bulgarian Food Safety Authority

**Authorisation number:**
0022-3242

**Date of authorisation status change:**
13/02/2024

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0409/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland

France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

**Generic of:**
600000047468

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.

NLV0409001DC Profexx cattle final PuAR.pdf