

**PARTICULARS TO APPEAR ON THE OUTER PACKAGE {Cardboard box}**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

RIMADYL Cattle 50 mg/ml Solution for injection.

**2. STATEMENT OF ACTIVE SUBSTANCES**

Carprofen 50 mg/ml

**3. PACKAGE SIZE**

50 ml  
100 ml  
250 ml

**4. TARGET SPECIES**

Cattle.

**5. INDICATIONS**

**6. ROUTES OF ADMINISTRATION**

For subcutaneous or intravenous use.

**7. WITHDRAWAL PERIODS**

Withdrawal periods:  
Meat and offal: 21 days.  
Milk: Zero days.

**8. EXPIRY DATE**

Exp. {mm/yyyy}  
Once broached, use within 28 days.

**9. SPECIAL STORAGE PRECAUTIONS**

Do not store above 30°C.  
Keep the container in the outer carton in order to protect from light.

**10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”**

Read the package leaflet before use.

**11. THE WORDS “FOR ANIMAL TREATMENT ONLY”**

For animal treatment only.

**12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”**

Keep out of the sight and reach of children.

**13. NAME OF THE MARKETING AUTHORISATION HOLDER**

Zoetis UK Limited

**14. MARKETING AUTHORISATION NUMBERS**

Vm 42058/5152

**15. BATCH NUMBER**

Lot {number}

**PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE {100 ml/250 ml}**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

RIMADYL Cattle 50 mg/ml Solution for injection.

**2. STATEMENT OF ACTIVE SUBSTANCES**

Carprofen 50 mg/ml

**3. TARGET SPECIES**

Cattle.

**4. ROUTES OF ADMINISTRATION**

s.c., i.v.

Read the package leaflet before use.

**5. WITHDRAWAL PERIODS**

Withdrawal period:

Meat and offal: 21 days.

Milk: Zero days.

**6. EXPIRY DATE**

Exp. {mm/yyyy}

Once broached, use within 28 days.

Once broached, use by:

**7. SPECIAL STORAGE PRECAUTIONS**

Do not store above 30°C.

Keep the container in the outer carton in order to protect from light.

**8. NAME OF THE MARKETING AUTHORISATION HOLDER**

Zoetis UK Limited

**9. BATCH NUMBER**

Lot {number}

**MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING**  
**UNITS {Vial label – 50 ml}**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

RIMADYL Cattle

**2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES**

Carprofen 50 mg/ml

**3. BATCH NUMBER**

Lot {number}

**4. EXPIRY DATE**

Exp. {mm/yyyy}

Once broached, use within 28 days.

Once broached, use by:

## **PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:**

### **PACKAGE LEAFLET**

#### **1. Name of the veterinary medicinal product**

RIMADYL Cattle 50 mg/ml solution for injection

#### **2. Composition**

Each ml contains:

##### **Active substance:**

Carprofen 50 mg

##### **Excipients:**

Ethanol (96%) 0.1 ml

Benzyl Alcohol 10 mg

Clear, pale straw yellow solution.

#### **3. Target species**

Cattle.

#### **4. Indications for use**

The veterinary medicinal product is indicated as an adjunct to antimicrobial therapy to reduce clinical signs in acute infectious respiratory disease and acute mastitis in cattle.

#### **5. Contraindications**

Do not use in animals suffering from cardiac, hepatic or renal impairment.

Do not use in animals suffering from gastro-intestinal ulceration or bleeding.

Do not use where there is evidence of a blood dyscrasia.

Do not use in cases of hypersensitivity to the active substance or to the excipients.

#### **6. Special warnings**

##### Special precautions for safe use in the target species:

Avoid use in any dehydrated, hypovolaemic or hypotensive animal, as there is a potential risk of increased renal toxicity. Concurrent administration of potentially nephrotoxic drugs should be avoided.

Do not exceed the stated dose or the duration of treatment.

Do not administer other NSAID's concurrently or within 24 hours of each other.

As NSAID therapy can be accompanied by gastro-intestinal or renal impairment, adjunctive fluid therapy should be considered especially in the case of acute mastitis treatment.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Carprofen, in common with other NSAIDs, has been shown to exhibit photosensitising potential in laboratory studies. Avoid skin contact with the product. Should this occur, wash the affected areas immediately.

Pregnancy:

The safety of the veterinary medicinal product has not been established during pregnancy.

Use only according to the benefit-risk assessment by the responsible veterinarian.

Interaction with other medicinal products and other forms of interaction:

In common with other NSAIDs, carprofen should not be administered simultaneously with another veterinary medicinal product of the NSAID or glucocorticoid class.

NSAID's are highly bound to plasma proteins and may compete with other highly bound drugs, such that concomitant administration may result in toxic effects.

However during clinical studies in cattle four different antibiotic classes were used, macrolides, tetracyclines, cephalosporins and potentiated penicillins without known interactions.

Overdose:

In clinical studies, no adverse signs were reported after intravenous and subcutaneous administration of up to 5 times the recommended dose.

There is no specific antidote for carprofen overdosage but general supportive therapy, as applied to clinical overdosage with NSAID's, should be applied.

Major incompatibilities:

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

**7. Adverse events**

Cattle:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):
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Injection site reaction <sup>1</sup>
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<sup>1</sup> Transient

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder using the contact details at the end of this leaflet, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>

e-mail: [adverse.events@vmd.gov.uk](mailto:adverse.events@vmd.gov.uk)

## **8. Dosage for each species, routes and method of administration**

Subcutaneous (s.c) or intravenous use (i.v.)

The recommended dose is 1.4 mg carprofen / kg body weight (1 ml/35 kg) in combination with antibiotic therapy as appropriate.

Single subcutaneous or intravenous injection.

## **9. Advice on correct administration**

When treating groups of animals, use a draw-off needle to avoid excessive broaching of the stopper. The maximum number of broachings should be limited to 20.

## **10. Withdrawal periods**

Meat and offal: 21 days.

Milk: Zero days.

## **11. Special storage precautions**

Do not store above 30°C.

Keep the container in the outer carton in order to protect from light.

Once broached, use within 28 days.

Do not use this veterinary medicinal product after the expiry date which is stated on the label after Exp. The expiry date refers to the last day of that month.

When the container is broached (opened) for the first time, using the in-use shelf-life which is specified on this package leaflet, the date on which any product remaining in the container should be discarded should be worked out. This discard date should be written in the space provided on the label.

## **12. Special precautions for disposal**

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

### **13. Classification of veterinary medicinal products**

Veterinary medicinal product subject to prescription.

### **14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES**

Vm 42058/5152

Available in a cardboard box containing one multidose amber glass (Type I) vial of either 50 ml, 100 ml or 250 ml capped with bromobutyl rubber stopper retained by an aluminium crimped seal.

Not all pack sizes may be marketed.

### **15. PID LINK (Do not print heading)**

*[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]*

Find more product information by searching for the 'Product Information Database' on [www.gov.uk](http://www.gov.uk).

### **16. Contact details**

Marketing authorisation holder and contact details to report suspected adverse events:

Zoetis UK Limited  
1st Floor, Birchwood Building  
Springfield Drive  
Leatherhead  
Surrey  
KT22 7LP  
Tel: +44 (0) 345 300 8034

Manufacturer responsible for batch release:

Bela-Pharm GmbH & Co. KG  
Lohner Str. 19  
49377 Vechta  
Germany

### **17. Other information**

POM-V

*Gavin Hall*  
Approved: 16 December 2025