PARTICULARS TO APPEAR ON THE OUTER PACKAGE - Cardboard Box: 10 ml, 50 ml, 100 ml or 250 ml vials.

1. NAME OF THE VETERINARY MEDICINAL PRODUCT
Recocam 20 mg/ml solution for injection
Meloxicam

2. STATEMENT OF ACTIVE SUBSTANCES
Meloxicam 20 mg/ml

3. PACKAGE SIZE
1 x 10 ml
1 x 50 ml
1 x 100 ml
1 x 250 ml

4. TARGET SPECIES
Cattle, pigs and horses.

5. INDICATIONS
Read the package leaflet before use.

6. ROUTES OF ADMINISTRATION
Cattle: SC, IV

Pigs: IM

Horses: IV

Read the package leaflet before use.
Do not broach the stopper more than 50 times.

7. WITHDRAWAL PERIODS
Withdrawal period:
Cattle: meat and offal: 15 days; milk: 5 days

Pigs: meat and offal: 5 days

Horses: meat and offal: 5 days.
Not authorised for use in horses producing milk for human consumption.

8. EXPIRY DATE
Exp. {mm/yyyy}

Shelf-life after first opening the container: 28 days
Once broached, use by ….

9. SPECIAL STORAGE PRECAUTIONS

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
Bimeda Animal Health Limited

14. MARKETING AUTHORISATION NUMBERS
Vm 50146/5000

15. BATCH NUMBER
Lot {number}

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE - Glass vial label for 50 ml, 100 ml and 250 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT
Recocam 20 mg/ml solution for injection
Meloxicam

2. STATEMENT OF ACTIVE SUBSTANCES
Meloxicam 20 mg/ml

3. TARGET SPECIES

Cattle, pigs and horses.

4. ROUTES OF ADMINISTRATION
Cattle: SC, IV
Pigs: IM
Horses: IV

Read the package leaflet before use.
Do not broach more than 50 times.

5. WITHDRAWAL PERIODS
Withdrawal periods:
Cattle: meat and offal: 15 days; milk: 5 days

Pigs: meat and offal: 5 days

Horses: meat and offal: 5 days.
Not authorised for use in horses producing milk for human consumption.

6. EXPIRY DATE
Exp. {mm/yyyy}

Shelf-life after first opening the container: 28 days
Once broached, use by ….

7. SPECIAL STORAGE PRECAUTIONS

8. NAME OF THE MARKETING AUTHORISATION HOLDER
Bimeda Animal Health Limited

9. BATCH NUMBER
Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS
- Glass vial label for 10 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT
Recocam 20 mg/ml solution for injection

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES
Meloxicam 20 mg/ml

3. BATCH NUMBER
Lot {number}

4. EXPIRY DATE
Exp. {mm/yyyy}

Shelf-life after first opening the container: 28 days
Once broached, use by ….

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product
Recocam 20 mg/ml solution for injection for cattle, pigs and horses.

2. Composition
One ml contains:

Active substances:
Meloxicam 20 mg

Excipients:
Ethanol 99.9% 150 mg

Clear yellow solution.

3. Target species
Cattle, pigs and horses.

4. Indications for use
Cattle
For use in acute respiratory infection with appropriate antibiotic therapy to reduce clinical signs in cattle.
For use in diarrhoea in combination with oral re-hydration therapy to reduce clinical signs in calves of over one week of age and young, non-lactating cattle.
For adjunctive therapy in the treatment of acute mastitis, in combination with antibiotic therapy.
For the relief of post-operative pain following dehorning in calves.

Pigs
For use in non-infectious locomotor disorders to reduce the symptoms of lameness and inflammation.
For adjunctive therapy in the treatment of puerperal septicaemia and toxaemia (mastitis-metritis-agalactia syndrome) with appropriate antibiotic therapy.

Horses
For use in the alleviation of inflammation and relief of pain in both acute and chronic musculo-skeletal disorders.
For the relief of pain associated with equine colic.

5. Contraindications
Do not use in horses less than 6 weeks of age.

Do not use in pregnant or lactating mares.
Do not use in horses producing milk for human consumption.

Do not use in animals suffering from impaired hepatic, cardiac or renal function and haemorrhagic disorders, or where there is evidence of ulcerogenic gastrointestinal lesions.

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

For the treatment of diarrhoea in cattle, do not use in animals of less than one week of age.

6. Special warnings

Treatment of calves with the veterinary medicinal product 20 minutes before dehorning reduces postoperative pain. The veterinary medicinal product alone will not provide adequate pain relief during the dehorning procedure. To obtain adequate pain relief during surgery co-medication with an appropriate analgesic is needed.

Special precautions for safe use in the target species:
If adverse reactions occur, treatment should be discontinued and the advice of a veterinarian should be sought.
Avoid use in very severely dehydrated, hypovolaemic or hypotensive animals which require parenteral rehydration, as there may be a potential risk of renal toxicity.
In case of inadequate relief of pain when used in the treatment of equine colic, careful re-evaluation of the diagnosis should be made as this could indicate the need for surgical intervention.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
Accidental self-injection may give rise to pain.
People with known hypersensitivity to Non-Steroidal Anti-Inflammatory Drugs (NSAIDs) should avoid contact with the veterinary medicinal product.

Pregnancy and lactation:
Cattle and pigs**:** Can be used during pregnancy and lactation.
Horses: Do not use in pregnant or lactating mares.

Interaction with other medicinal products and other forms of interaction:
Do not administer concurrently with glucocorticosteroids, other non-steroidal anti-inflammatory drugs or with anti-coagulant agents.

Overdose:
In the case of overdose, symptomatic treatment should be initiated.

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):	Anaphylactoid reaction[1], Injection site swelling[2]

[1]May be serious (including fatal), should be treated symptomatically.
[2]Slight, transient, following subcutaneous administration.

Pigs:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Anaphylactoid reaction[1]

[1]May be serious (including fatal), should be treated symptomatically.

Horses:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Anaphylactoid reaction[1], Injection site swelling[2]

[1]May be serious (including fatal), should be treated symptomatically.
[2]Transient. Resolves without intervention.

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 or the local representative of 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

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

Cattle
Single subcutaneous or intravenous injection at a dose of 0.5 mg meloxicam/kg body weight (i.e. 2.5 ml/100 kg body weight) in combination with antibiotic therapy or with oral re-hydration therapy, as appropriate.

Pigs
Single intramuscular injection at a dose of 0.4 mg meloxicam/kg body weight (i.e. 2.0 ml/100 kg body weight) in combination with antibiotic therapy, as appropriate. If required, a second administration of meloxicam can be given after 24 hours.

Horses
Single intravenous injection at a dose of 0.6 mg meloxicam/kg body weight (i.e. 3.0 ml/100 kg body weight).
For use in the alleviation of inflammation and the relief of pain in both acute and chronic musculo–skeletal disorders, oral suspensions of meloxicam may be used for continuation of treatment at a dosage of 0.6 mg meloxicam/kg body weight, 24 hours after administration of the injection.

9. Advice on correct administration

Avoid introduction of contamination during use.
Do not broach the stopper more than 50 times.

10. Withdrawal periods

Cattle:
Meat and offal: 15 days
Milk: 5 days

Pigs:
Meat and offal: 5 days

Horses:
Meat and offal: 5 days.
Not authorised for use in horses producing milk for human consumption.

11. Special storage precautions

Keep out of the sight and reach of children.

This veterinary medicinal product does not require any special storage conditions.

Do not use this veterinary medicinal product after the expiry date which is stated on the carton and vial after Exp.
The expiry date refers to the last day of that month.
Shelf life after first opening the immediate packaging: 28 days.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater or household waste.
Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

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.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Vm 50146/5000

Pack size of 1 Type I clear glass vial with Teflon coated bromobutyl rubber stoppers and sealed with an aluminium flip-off tear-off seal. Vials contain 10 ml, 50 ml, 100 ml or 250 ml.
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.

16. Contact details

Marketing authorisation holder and manufacturer responsible for batch release:
Bimeda Animal Health Limited
2/3/4 Airton Close
Tallaght
Dublin 24
Ireland

Local representatives and contact details to report suspected adverse reactions:
Cross Vetpharm Group UK Limited (Trading as Bimeda)
Unit 2, Bryn Cefni Industrial Park
Llangefni, LL77 7XA
United Kingdom
Tel: 01248 725 400

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder.

17. Other information

POM-V

Gavin Hall

Approved: 17 April 2026