

PARTICULARS TO APPEAR ON THE OUTER PACKAGE BOX WITH VIAL(S) OF 50 ML, 100 ML AND 250 ML

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Labiprofen 150 mg/ml solution for injection for cattle, pigs and horses

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Each ml contains:

Active substance:

Ketoprofen 150 mg

Excipient:

Benzyl alcohol (E1519) 10 mg

3. PACKAGE SIZE

1 x 50 ml

1 x 100 ml

1 x 250 ml

12 x 50 ml

10 x 100 ml

10 x 250 ml

4. TARGET SPECIES

Cattle, pigs and horses.



5. INDICATION(S)

6. ROUTES OF ADMINISTRATION

Intramuscular or intravenous use.

7. WITHDRAWAL PERIODS

Withdrawal period:

Cattle:

Meat and offal: 2 days.

Milk: zero hours.

Horses:

Meat and offal: 1 day.

Milk: Not authorised for use in animals producing milk for human consumption.

Pig:

Meat and offal: 3 days.

8. EXPIRY DATE

Exp {month/year}
Once opened use within 28 days.
Use by:

9. SPECIAL STORAGE PRECAUTIONS

Keep the vial 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

Labiana Life Sciences, S.A.

14. MARKETING AUTHORISATION NUMBERS

Vm 32112/5000

15. BATCH NUMBER

Lot {number}

16. SPECIAL WARNING(S), IF NECESSARY

17. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: Read package leaflet

18. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

POM-V To be supplied only on veterinary prescription.

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE 100 ml and 250 ml Vial

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Labiprofen 150 mg/ml solution for injection for cattle, pigs and horses

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Each ml contains:

Active substance:

Ketoprofen 150 mg

Excipient:

Benzyl alcohol (E1519) 10 mg

3. TARGET SPECIES

Cattle, pigs and horses.



4. ROUTES OF ADMINISTRATION

Intramuscular or intravenous use.

5. WITHDRAWAL PERIODS

Withdrawal period:

Cattle:

Meat and offal: 2 days.

Milk: zero hours.

Horses:

Meat and offal: 1 day.

Milk: Not authorised for use in animals producing milk for human consumption.

Pig:

Meat and offal: 3 days.

6. EXPIRY DATE

Exp {month/year}

Once opened use within 28 days.

Use by:

7. SPECIAL STORAGE PRECAUTIONS

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

8. NAME OF THE MARKETING AUTHORISATION HOLDER

Labiana Life Sciences, S.A.

9. BATCH NUMBER

Lot {number}

10. PACKAGE SIZE

100 ml

250 ml

11. INDICATION(S)

12. SPECIAL WARNING(S), IF NECESSARY

13. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: Read package leaflet.

14. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE

POM-V For animal treatment only. To be supplied only on veterinary prescription.

15. MARKETING AUTHORISATION NUMBER(S)

Vm 32112/5000

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS 50 ML VIAL

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Labiprofen 150 mg/ml solution for injection



2. QUALITATIVE AND QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCE(S)

Each ml contains:

Ketoprofen 150 mg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp {month/year}

Once opened use within 28 days.

Use by:

5. CONTENTS BY WEIGHT, BY VOLUME OR BY NUMBER OF DOSES

50 ml

6. ROUTE(S) OF ADMINISTRATION

Intramuscular or intravenous use.

7. WITHDRAWAL PERIOD

Cattle:

Meat and offal: 2 days.

Milk: zero hours.

Horses:

Meat and offal: 1 day.

Milk: Not authorised for use in animals producing milk for human consumption.

Pig:

Meat and offal: 3 days.

8. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

PACKAGE LEAFLET:

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Labiprofen 150 mg/ml solution for injection for cattle, pigs and horses.

2. COMPOSITION

Each ml contains:

Active substance:

Ketoprofen 150 mg

Excipient:

Benzyl alcohol (E1519) 10 mg

A clear colourless to yellowish solution.

3. TARGET SPECIES

Cattle, pigs, and horses.

4. INDICATIONS FOR USE

Cattle:

- Reduction of inflammation and pain associated with post-partum, musculoskeletal disorders and lameness.
- Reduction of fever associated with bovine respiratory disease in combination with antimicrobial therapy where appropriate.
- Reduction of inflammation, fever and pain in acute clinical mastitis in combination with antimicrobial therapy where appropriate.

Pigs:

- Reduction of pyrexia in cases of respiratory disease and Postpartum Dysgalactia SyndromePDS- (Metritis Mastitis Agalactia syndrome) in sows, in combination with antimicrobial therapy, where appropriate.

Horses:

- Reduction of inflammation and pain associated with osteoarticular and musculoskeletal disorders (lameness, laminitis, osteoarthritis, synovitis, tendinitis, etc.).
- Reduction of postoperative pain and inflammation.
- Reduction of visceral pain associated with colic.

5. CONTRAINDICATIONS

Do not use in cases of gastro-intestinal ulceration or bleeding, in order not to aggravate their situation.

Do not use in cases of cardiac, hepatic, or renal disease.

Do not use in cases of known hypersensitivity to ketoprofen or acetyl-salicylic acid or to any of the excipients.

Do not use in cases of blood dyscrasia, coagulopathy or haemorrhagic diathesis. Do not administer other non-steroidal anti-inflammatory drugs (NSAIDs) concurrently or within 24 hours of each other.

6. SPECIAL WARNING(S)

For Animal Treatment Only

Special precautions for safe use in the target species:

Do not exceed the recommended dose. Do not exceed the recommended treatment period.

The use of ketoprofen is not recommended in foals less than one month of age.

When administering to animals of less than 6 weeks of age, ponies or in aged animals it is necessary to adjust the dose accurately as well as to perform a close clinical follow-up.

Avoid intra-arterial injection.

Avoid use in any dehydrated, hypovolaemic or hypotensive animals as there is a potential risk of increased renal toxicity.

Since gastric ulceration is a common finding in PMWS (Post-weaning Multisystemic Wasting Syndrome), the use of ketoprofen in pigs affected by this pathology is not recommended, in order not to aggravate their situation. In horses, avoid extravascular administration.

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

Hypersensitivity reactions (skin rash, urticaria) could occur. People with known hypersensitivity to the active substance or to any of the excipients should avoid contact with the veterinary medicinal product.

Avoid accidental self-injection. In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the physician.

Avoid contact with the skin, eyes and mucous membranes. In case of accidental skin, eye or mucous membrane contact, wash the affected area thoroughly with clean running water immediately. If irritation persists seek medical advice and show the package leaflet or the label to the physician.

Wash hands after use.

Pregnancy:

Laboratory studies in rats, mice and rabbits, and studies in cattle have not produced any evidence of adverse effects. Can be used during pregnancy in cows.

The safety of the veterinary medicinal product has not been established during pregnancy in sows and mares. Use only according to the benefit-risk assessment by the responsible veterinarian.

Lactation:

Can be used during lactation in cows and sows.

The use is not recommended during lactation in mares.

Interaction with other medicinal products and other forms of interaction:

- Concurrent administration of diuretics or potentially nephrotoxic drugs should be avoided since there is an increase of renal disturbances, including renal failure. This is secondary to the diminished blood flow caused by the inhibition of prostaglandins synthesis.

- Do not administer other non-steroidal anti-inflammatory drugs (NSAIDs), corticosteroids, anticoagulants or diuretics concurrently or within 24 hours of administration of the product since the risk of gastrointestinal ulceration and other adverse reactions may be exacerbated.

- The treatment free period should however take into account the pharmacological properties of the products used previously.

- Ketoprofen is highly bound to plasma proteins and may compete with other highly bound drugs which can lead to toxic effects.

Overdose:

Overdose with non-steroidal anti-inflammatory drugs can lead to gastro-intestinal ulceration, loss of proteins, hepatic and renal impairment.

In tolerance studies performed in pigs, up to 25% of the animals treated at three times the maximum recommended dose (9 mg/kg bw) for three days or at the recommended dose (3 mg/kg bw) for triple the maximum recommended time (9 days) showed erosive and/or ulcerative lesions in both the aglandular (pars oesophagica) and glandular parts of the stomach. Early signs of toxicity include loss of appetite and pasty faeces or diarrhoea.

The intramuscular administration of the product to cattle, at up to 3 times the recommended dose or for 3 times the recommended duration of the treatment (9 days) did not result in clinical signs of intolerance. However, inflammation as well as necrotic subclinical lesions were detected at the injection site of the treated animals as well as an increase in CPK levels. The histopathological examination showed erosive or ulcerative abomasal lesions related to both dosage regimes.

Horses have been found to tolerate intravenous dosages of ketoprofen up to 5 times the recommended dose for three times the recommended duration (15 days) with no evidence of toxic effects.

If clinical signs of overdose are observed, there is no specific antidote, therefore 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, pigs

Undetermined frequency (cannot be estimated from the available data)	Necrotic muscular lesions ¹ Erosive and ulcerative lesions of the gastrointestinal tract ² Gastric or renal intolerance ³
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Horse

Undetermined frequency (cannot be estimated from the available data)	Necrotic muscular lesions ¹ Erosive and ulcerative lesions of the gastrointestinal tract ² Gastric or renal intolerance ³ Injection site reactions ⁴
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1. After intramuscular injection, subclinical, mild and transient, gradually resolving in the days after completion of treatment. Administration in the neck region minimizes the extension and severity of these lesions.
2. After repeated administrations (due to the mechanism of action of ketoprofen).
3. In certain individuals. Due to the action of inhibition of prostaglandins' synthesis (in common with all NSAIDS).
4. Transient. Observed after one administration of the product at the recommended volume by extravascular route. Disappeared after 5 days.

If side effects occur treatment must be stopped, and the advice of a veterinarian should be sought.

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 its local representative using the contact details at the end of this leaflet, or via your national reporting system for UK: <https://www.gov.uk/report-veterinary-medicine-problem> or IE www.hpra.ie

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

Intramuscular use: cattle, pigs

Intravenous use: cattle, horses

- Cattle:

3 mg ketoprofen/kg body weight, i.e. 1 ml of the veterinary medicinal product per 50 kg body weight/ day, administered via the intravenous or intramuscular route, preferably in the neck region.

The duration of treatment is 1-3 days, and should be established according to the severity and duration of symptoms.

- Pigs:

3 mg of ketoprofen/kg body weight i.e. 1 ml of the veterinary medicinal product per 50 kg body weight/ day, administered via the intramuscular route on a single occasion. Depending on the response observed and based on the benefit-risk analysis by the responsible veterinarian treatment may be repeated at intervals of 24 hours for a maximum of three treatments. Each injection should be given at a different site.

- Horses:

2.2 mg of ketoprofen/kg body weight, i.e. 0.75 ml of the veterinary medicinal product per 50 kg body weight/ day, administered via the intravenous route.

The duration of treatment is 1-5 days, and should be established according to the severity and duration of symptoms. In the case of colic one injection is normally sufficient. A second administration of ketoprofen requires a clinical re-examination.

9. ADVICE ON CORRECT ADMINISTRATION

10. WITHDRAWAL PERIOD(S)

Cattle:

Meat and offal: 2 days.

Milk: zero hours.

Horses:

Meat and offal: 1 day.

Milk: Not authorised for use in animals producing milk for human consumption.

Pig:

Meat and offal: 3 days.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.

Keep the immediate packaging in the outer carton in order to protect from light.

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.

When a immediate packaging 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 immediate packaging should be discarded should be worked out. This discard date should be written in the space provided.

12. SPECIAL PRECAUTIONS FOR DISPOSAL

Medicines should not be disposed of via wastewater.

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required. 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 32112/5000

Package sizes:

Box containing 1 vial of 50 ml

Box containing 1 vial of 100 ml

Box containing 1 vial of 250 ml

Box containing 12 vials of 50 ml

Box containing 10 vials of 100 ml

Box containing 10 vials of 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 and contact details to report suspected adverse events:

Labiana Life Sciences SA, Calle Venus 26, 08228 Terrassa, Barcelona, Spain

Local representative:

Interchem (Ireland) Ltd, Unit 29, Cookstown Industrial Estate, Dublin 24, Phone 01 451 8959 and or E-mail: SADR@interchem.ie

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

17. OTHER INFORMATION

POM-V

To be supplied only on veterinary prescription

Gavin Hall
Approved: 01 April 2026