

LABELLING

PARTICULARS TO APPEAR ON THE OUTER PACKAGE {CARDBOARD BOX (20 sachets or 100 sachets)}.

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Inflacam 330 mg granules

2. STATEMENT OF ACTIVE SUBSTANCES

Each sachet contains:

Meloxicam 330 mg

3. PACKAGE SIZE

20 sachets
100 sachets

4. TARGET SPECIES

Horses

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

In feed use.

7. WITHDRAWAL PERIODS

Withdrawal period:

Meat and offal: 3 days.

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

8. EXPIRY DATE

Exp. {mm/yyyy}

Once incorporated into muesli feed, use immediately.

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

Chanelle Pharmaceuticals Manufacturing Ltd

14. MARKETING AUTHORISATION NUMBERS

Vm 08749/5016

15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS {SACHET}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Inflacam 

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each sachet contains:

Meloxicam 330 mg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Inflacam 330 mg granules for horses

2. Composition

Each sachet contains:

Active substance

Meloxicam 330 mg

Pale yellow granules.

3. Target species

Horses.

4. Indications for use

Alleviation of inflammation and relief of pain in both acute and chronic musculo-skeletal disorders in horses weighing between 500 and 600 kg.

5. Contraindications

Do not use in pregnant or lactating mares.

Do not use in horses suffering from gastrointestinal disorders such as irritation and haemorrhage, impaired hepatic, cardiac or renal function and haemorrhagic disorders. Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

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

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 renal toxicity.

In order to minimise the risk of intolerance, the product should be mixed into muesli feed.

This product is only for use in horses weighing between 500 and 600 kg.

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

People with known hypersensitivity to non-steroidal anti-inflammatory drugs (NSAIDs) should avoid contact with the veterinary medicinal product.

In case of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.

Pregnancy and lactation:

Laboratory studies in cattle have not provided any evidence of teratogenic, foetotoxic, or maternotoxic effects. However, no data have been generated in horses. Therefore, do not use in pregnant or lactating mares.

Interaction with other medicinal products and other forms of interaction:

Do not administer concurrently with glucocorticoids, other NSAIDs or with anti-coagulant agents.

Overdose:

In 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

Horses:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Appetite loss, Lethargy Diarrhoea ¹ , Abdominal pain, Colitis. Urticaria ^{1,2} , Anaphylactoid reactions ³ .
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¹Reversible

²Slight

³May be serious (including fatal). If such reaction occurs, it should be treated symptomatically.

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 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

In-feed use.

To be administered mixed with food at a dose of 0.6 mg/kg body weight, once daily, up to 14 days. The product should be added to 250 g of muesli feed, prior to feeding.

Each sachet contains one dose for a horse weighing between 500 kg and 600 kg and the dose must not be divided into smaller doses.

9. Advice on correct administration

To ensure a correct dosage, body weight should be determined as accurately as possible.

The use of suitably calibrated measuring equipment is recommended.

Avoid introduction of contamination during use.

10. Withdrawal periods

Meat and offal: 3 days.

Not authorised for use in animals 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 sachet after Exp. The expiry date refers to the last day of that month.

Shelf life after incorporation into muesli feed: use immediately.

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 08749/5016.

Pack sizes:

Cardboard box with 20 or 100 sachets.

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:

Chanelle Pharmaceuticals Manufacturing Ltd
Loughrea
Co. Galway
Ireland
H62 FH90
Telephone: +353 (0)91 841788

Local representatives and contact details to report suspected adverse reactions:

Virbac Ltd
Suffolk
IP30 9UP
United Kingdom
Tel: + 44 (0) 1359 243243

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: 11 December 2025