

**PARTICULARS TO APPEAR ON THE OUTER PACKAGE {Outer carton/125 ml  
and 336 ml}**

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

Novaquin 15 mg/ml oral suspension

**2. STATEMENT OF ACTIVE SUBSTANCES**

Each ml contains:

**Active substance:**

Meloxicam 15 mg

**3. PACKAGE SIZE**

125 ml

336 ml

**4. TARGET SPECIES**

Horses.



**5. INDICATIONS**

**6. ROUTES OF ADMINISTRATION**

Oral use.

Shake vigorously at least 20 times before use.

**7. WITHDRAWAL PERIODS**

Withdrawal period:

Meat and offal: 3 days.

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

**8. EXPIRY DATE**

Exp. {mm/yyyy}

Once opened use within: 5 months.

**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**

Le Vet Beheer B.V.

**14. MARKETING AUTHORISATION NUMBERS**

Vm 41821/5000

**15. BATCH NUMBER**

Lot {number}

**PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE {LABEL/HDPE  
Bottle 125 ml or 336 ml}**

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

Novaquin 15 mg/ml oral suspension for horses

**2. STATEMENT OF ACTIVE SUBSTANCES**

Each ml contains:

**Active substance:**

Meloxicam 15 mg

**3. TARGET SPECIES**

Horses.



**4. ROUTES OF ADMINISTRATION**

Read the package leaflet before use.

Oral use

Shake vigorously at least 20 times before use.

**5. WITHDRAWAL PERIODS**

Withdrawal periods:

Meat and offal: 3 days.

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

**6. EXPIRY DATE**

Exp. {mm/yyyy}

Once opened use within: 5 months.

**7. SPECIAL STORAGE PRECAUTIONS**

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

Le Vet Beheer B.V.

**9. BATCH NUMBER**

Lot {number}

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

**PACKAGE LEAFLET**

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

Novaquin 15 mg/ml oral suspension for horses

**2. Composition**

Each ml contains:

**Active substance**

Meloxicam 15 mg.

**Excipients**

Sodium benzoate 1.75 mg

Yellowish-green viscous oral suspension.

**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.

**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 warnings:

None.

Special precautions for safe use in the target species:

Avoid use in any dehydrated, hypovolemic or hypotensive animal, as there is a potential risk of renal toxicity.

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 for teratogenic, foetotoxic, or maternotoxic effects. However, no data have been generated in horses. Do not use the veterinary medicinal product in pregnant or lactating mares. See section "Contraindications".

Interaction with other medicinal products and other forms of interaction:

Do not administer concurrently with glucocorticoids, other non-steroidal anti-inflammatory drugs or with anticoagulant agents.

Overdose:

In case of overdose symptomatic treatment should be initiated.

Special restrictions for use and special conditions for use:

Not applicable.

Major incompatibilities:

None known.

## 7. Adverse events

Horses:

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Diarrhoea <sup>a</sup> , Abdominal pain, Colitis Appetite loss, Lethargy Urticaria <sup>a,b</sup> , Anaphylactoid reaction <sup>c</sup>
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<sup>a</sup> Adverse reactions typically associated with NSAIDs. The clinical signs are reversible.

<sup>b</sup> Slight

<sup>c</sup> May be serious (including fatal) and should be treated symptomatically.

If adverse reactions occur, treatment should be discontinued 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 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](mailto:adverse.events@vmd.gov.uk)

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

For oral use.

### Dosage

Oral suspension to be administered at a dosage of 0.6 mg/kg body weight, once daily, up to 14 days.

### Method and route of administration

Shake vigorously at least 20 times before use. To be administered either mixed with a small quantity of food, prior to feeding, or directly into the mouth.

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

Avoid introduction of contamination during use.

The suspension should be given using the measuring syringe provided in the package. The syringe fits onto the bottle and has a kg-body weight scale.

After administration of the veterinary medicinal product, close the bottle by replacing the cap, wash the measuring syringe with warm water and let it dry.

## **10. Withdrawal periods**

Meat and offal: 3 days.

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

Shelf life after first opening the immediate packaging: 5 months.

## **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 41821/5000

Cardboard box containing one bottle of 125 ml.

Cardboard box containing one bottle of 336 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](http://www.gov.uk).

### **16. Contact details**

#### Marketing authorisation holder:

Le Vet Beheer B.V.  
Wilgenweg 7  
3421 TV Oudewater  
The Netherlands

#### Manufacturer responsible for batch release:

Produlab Pharma B.V.  
Forellenweg 16  
4941 SJ Raamsdonksveer  
The Netherlands

#### Local representatives and contact details to report suspected adverse reactions:

Dechra Veterinary Products Limited  
Sansaw Business Park  
Hadnall  
Shrewsbury  
Shropshire  
SY4 4AS  
United Kingdom  
Tel: +44 (0)1939 211200

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

**17. Other information**

POM-V

Veterinary medicinal product subject to prescription

*Gavin Hall*

Approved: 06 August 2025