

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

folding carton

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Detonervin 10 mg/ml, solution for injection for Horses and Cattle
detomidine hydrochloride

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

1 ml contains detomidine hydrochloride 10 mg (equivalent to 8.36 mg detomidine)

3. PHARMACEUTICAL FORM

Solution for injection

4. PACKAGE SIZE

5 ml
5 x 5 ml
20 ml
5 x 20 ml

5. TARGET SPECIES

Horse and cattle

6. INDICATION(S)

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Intramuscular use, Intravenous use.

8. WITHDRAWAL PERIOD

Meat and offal: 2 days
Milk: 12 hours

9. SPECIAL WARNING(S), IF NECESSARY

Read the package leaflet before use.

10. EXPIRY DATE

EXP:

Shelf-life after first opening: 28 days.

11. SPECIAL STORAGE CONDITIONS

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

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: read package leaflet.

13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, if applicable

For animal treatment only.

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

Keep out of the sight and reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

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

16. MARKETING AUTHORISATION NUMBER

Vm 19994/4015

17. MANUFACTURER’S BATCH NUMBER

Batch:

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

(glass vial)

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Detonervin 10 mg/ml, solution for injection for Horses and Cattle
detomidine hydrochloride



2. QUANTITY OF THE ACTIVE SUBSTANCE(S)

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

5 ml
20 ml

4. ROUTE(S) OF ADMINISTRATION

IM, IV

5. WITHDRAWAL PERIOD

Meat and offal: 2 days
Milk: 12 hours

6. BATCH NUMBER

Batch:

7. EXPIRY DATE

EXP:
Once opened, use by.....

8. THE WORDS "FOR ANIMAL TREATMENT ONLY"

For animal treatment only

PACKAGE LEAFLET

Detonervin 10 mg/ml, solution for injection for Horses and Cattle

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

Marketing authorisation holder:

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

Manufacturer for the batch release:

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

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

Detonervin 10 mg/ml, solution for injection for Horses and Cattle
Detomidine hydrochloride

3. STATEMENT OF THE ACTIVE SUBSTANCE(S) AND OTHER INGREDIENT(S)

Detonervin is a clear and colourless solution for injection containing:

Active substance:

Detomidine hydrochloride 10.0 mg/ml
(equivalent to 8.36 mg detomidine)

Excipient:

Methyl parahydroxybenzoate (E218) 1.0 mg/ml

4. INDICATION(S)

For the sedation and slight analgesia of horses and cattle, to facilitate physical examinations and treatments, such as minor surgical interventions.

Detomidine can be used for:

- Examinations (e.g. endoscopy, rectal and gynaecological examinations, X-rays).
- Minor surgical procedures (e.g. treatment of wounds, dental treatment, tendon treatment, excision of skin tumours, teat treatment).
- Before treatment and medication (e.g. stomach tube, horse shoeing).

For premedication prior to administration of injection- or inhalation anaesthetics.
See section 12 before use.

5. CONTRAINDICATIONS

- Do not use in animals with cardiac abnormalities or respiratory diseases.
- Do not use in animals with liver insufficiency or renal failure.
- Do not use in animals with general health problems (e.g. dehydrated animals).
- Do not use in combination with butorphanol in horses suffering from colic.
- Do not use in the last trimester of pregnancy.

6. ADVERSE REACTIONS

Injection of detomidine may cause the following side effects:

- Bradycardia
- Transient hypo- and/or hypertension
- Respiratory depression, rarely hyperventilation
- Increase in blood glucose
- As with other sedatives, in rare cases paradoxical reactions (excitations) can develop
- Ataxia
- Uterine contractions
- In horses: Cardiac arrhythmia, atrioventricular and sino-atrial block
- In cattle: Inhibition of rumen motility, tympany, paralysis of the tongue.

At doses above 40 µg/kg bodyweight, the following symptoms can also be observed: sweating, pilo-erection and tremor of muscles, transient penis prolaps in stallions and geldings and mild, transient tympany of rumen and increased salivation in cattle.

In very rare cases horses may show mild symptoms of colic following administration of alpha-2 sympathomimetics because substances of this class transiently inhibit the motility of the intestines. Detomidine should be prescribed with caution in horses which present with signs of colic or impaction.

A diuretic effect is usually observed within 45 to 60 minutes after treatment.

If you notice any serious effects or other effects not mentioned in this leaflet, please inform your veterinary surgeon.

7. TARGET SPECIES

Horse, cattle.

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

For intravenous (IV) or intramuscular (IM) use. The product should be injected slowly. Onset of effect is more rapid following intravenous use.

Dosage in μg detomidine hydrochloride /kg	Dosage in ml/100 kg	Level of sedation	Commencement of effect (min)		Duration of effect(hrs)
			horse	cattle	
10-20	0.1-0.2	Light	3-5	5-8	0.5-1
20-40	0.2-0.4	Moderate	3-5	5-8	0.5-1

When prolonged sedation and analgesia is required, doses of 40 to 80 μg detomidine hydrochloride /kg can be used. The duration of effect is up to 3 hours.

For combination with other product to intensify the sedation or for premedication prior to general anaesthesia, doses of 10 to 30 μg detomidine hydrochloride /kg can be used.

It is recommended to wait 15 minutes after the detomidine administration before starting the planned procedure.

The bodyweight of the animal to be treated should be determined as accurately as possible to avoid overdosing.

9. ADVICE ON CORRECT ADMINISTRATION

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10. WITHDRAWAL PERIOD

Horse, cattle:

Meat and offal: 2 days

Milk: 12 hours

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.

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

Do not use after the expiry date stated on the label and on the carton after EXP.

Shelf-life after first opening the container: 28 days.

Discharge any product remaining in the container at this time.

12. SPECIAL WARNING(S)

Special precautions for use in animals

- As sedation begins, especially horses may start to sway and lower the head rapidly while they remain standing. Cattle and especially young cattle will try to lie down. To prevent injuries the location should therefore be chosen carefully. Especially for horses usual precautionary measures should be taken to prevent human or animal injury. To avoid ruminal bloat and aspiration of feed or saliva,

cattle should be maintained in sternal recumbency during and following treatment and head and neck of recumbent cattle should be lowered.

- Animals suffering from shock or liver or kidney disease should only be treated according to the benefit risk assessment by the responsible veterinarian.
- The product should not be used in animals suffering from cardiac diseases (with pre-existing bradycardia and risk of atrioventricular block), respiratory-, liver- or renal insufficiencies, shock or any other extraordinary stress conditions.
- Detomidine/butorphanol combination should not be used in horses with a history of liver disease or cardiac irregularities.
- - It is recommended that feed should be withheld for at least 12 hours prior to anaesthesia. Water or food should not be offered to treated animals until the drug effect has passed.
- In painful procedures detomidine should be used only in combination with an analgesic or a local anaesthetic.
- While waiting for sedation animals should remain in calm surroundings.
- In case of sustained effect it is necessary to protect the animals from heat or cold.

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

- In the case of accidental oral intake or self-injection seek medical advice immediately and show the package leaflet or the label to the physician. DO NOT DRIVE as sedation and changes in blood pressure may occur.
- Avoid skin, eye or mucosal contact.
- Wash the exposed skin immediately after exposure with large amounts of water.
- Remove contaminated clothes that are in direct contact with skin.
- In case of accidental contact of the product with eyes, rinse abundantly with fresh water. If symptoms occur, seek the advice of a physician.
- If pregnant women handle the product, special caution should be observed not to self-inject as uterine contractions and decreased foetal blood pressure may occur after accidental systemic exposure.

Advice to doctors:

Detomidine is an alpha₂-adrenoreceptor agonist, which after absorption may involve clinical effects including dose-dependent sedation, respiratory depression, bradycardia, hypotension, a dry mouth and hyperglycaemia. Ventricular arrhythmias have also been reported. Respiratory and haemodynamic symptoms should be treated symptomatically.

Use during pregnancy, lactation or lay

Do not use during the last trimester of pregnancy. Use only according to the benefit/risk assessment by the responsible veterinarian during the other months of pregnancy.

Interaction with other medicinal products and other forms of interaction

Concurrent use of other sedatives only after consultation of the warnings and precautions of the product concerned.

Detomidine should not be used in combination with sympathomimetic amines such as adrenaline, dobutamine and ephedrine.

The concurrent use of certain potentiated sulphonamides may cause cardiac arrhythmia with fatal outcome. Do not use in combination with sulphonamides.

Detomidine in combination with other sedatives and anaesthetics should be used carefully because additive/synergistic effects may be possible. Where anaesthesia is induced with a combination of detomidine and ketamine, prior to maintenance with halothane, the effects of halothane may be delayed and care must be taken to avoid overdosage. When detomidine is used as a premedicant prior to general anaesthesia, the product may delay the onset of induction.

Overdose (symptoms, emergency procedures, antidotes), if necessary

In the event of an accidental overdose, cardiac arrhythmias, hypotension, delayed recovery and profound CNS and respiratory depression may occur. Should the effects of detomidine become life-threatening, general measures for circulatory and respiratory stabilization and administration of an alpha2-adrenergic antagonist is recommended.

Incompatibilities

In the absence of compatibility studies, this veterinary medicinal product should not be mixed with other veterinary medicinal products in the same syringe.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

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

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

<to be completed nationally>

15. OTHER INFORMATION

Pack sizes:

1 x 1 glass vial with 5 ml.

5 x 1 glass vials with 5 ml.

1 x 1 glass vial with 20 ml.

5 x 1 glass vials with 20 ml.

Not all pack sizes may be marketed.

Approved: 09 June 2016

