TORBUGESIC- butorphanol tartrate solution
Zoetis Inc.

Torbugesic®
(butorphanol tartrate injection)

NADA 135-780, Approved by FDA
Torbugesic®
(butorphanol tartrate injection)

CAUTION
Federal (USA) law restricts this drug to use by or on the order of a licensed veterinarian.

DESCRIPTION
TORBUGESIC (butorphanol tartrate injection) is a totally synthetic, centrally acting, narcotic agonist-antagonist analgesic with potent antitussive activity. It is a member of the phenanthrene series. The chemical name is Morphinan-3, 14-diol, 17-(cyclobutylmethyl)-, (-)-, (S- (R*, R*))-2,3-dihydroxybutanedioate (1:1) (salt). It is a white, crystalline, water soluble substance having a molecular weight of 477.55; its molecular formula is C21H29NO2•C4H6O6.

Chemical Structure

Each mL of TORBUGESIC contains 10 mg butorphanol base (as butorphanol tartrate), 3.3 mg citric acid, USP, 6.4 mg sodium citrate, 4.7 mg sodium chloride, and 0.1 mg benzethonium chloride, q.s. with water for injection.

CLINICAL PHARMACOLOGY

Comparative Pharmacology
In animals, butorphanol has been demonstrated to be 4 to 30 times more potent than morphine and pentazocine (Talwin®-V) respectively.¹
In humans, butorphanol has been shown to have 5 to 7 times the analgesic activity of morphine and 20 times that of pentazocine.\textsuperscript{2,3}

Butorphanol has 15 to 20 times the oral antitussive activity of codeine or dextromethorphan in dogs and guinea pigs.\textsuperscript{4}

As an antagonist, butorphanol is approximately equivalent to nalorphine and 30 times more potent than pentazocine.\textsuperscript{1}

Cardiopulmonary depressant effects are minimal after treatment with butorphanol as demonstrated in dogs\textsuperscript{5}, humans\textsuperscript{6,7} and horses.\textsuperscript{8}

Unlike classical narcotic agonist analgesics which are associated with decreases in blood pressure, reduction in heart rate and concomitant release of histamine, butorphanol does not cause histamine release.\textsuperscript{1}

Furthermore, the cardiopulmonary effects of butorphanol are not distinctly dosage-related but rather reach a ceiling effect beyond which further dosage increases result in relatively lesser effects.

Reproduction: Studies performed in mice and rabbits revealed no evidence of impaired fertility or harm to the fetus due to butorphanol tartrate. In the female rat, parenteral administration was associated with increased nervousness and decreased care for the newborn, resulting in a decreased survival rate of the newborn. This nervousness was seen only in the rat species.

**Equine Pharmacology**

Following intravenous injection in horses, butorphanol is largely eliminated from the blood within 3 to 4 hours. The drug is extensively metabolized in the liver and excreted in the urine. In ponies, butorphanol given intramuscularly at a dosage of 0.22 mg/kg, was shown to alleviate experimentally induced visceral pain for about 4 hours.\textsuperscript{9}

In horses, intravenous dosages of butorphanol ranging from 0.05 to 0.4 mg/kg were shown to be effective in alleviating visceral and superficial pain for at least 4 hours, as illustrated in the following figure:

<table>
<thead>
<tr>
<th>Analgesic Effects of Butorphanol Given at Various Dosages in Horses with Abdominal Pain</th>
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<tbody>
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</table>
A definite dosage-response relationship was detected in that butorphanol dosage of 0.1 mg/kg was more effective than 0.05 mg/kg but not different from 0.2 mg/kg in alleviating deep abdominal pain.

**Acute Equine Studies**

Rapid intravenous administration of butorphanol at a dosage of 2 mg/kg (20 times the recommended dosage) to a previously unmedicated horse resulted in a brief episode of inability to stand, muscle fasciculation, a convulsive seizure of 6 seconds duration and recovery within three minutes. The same dosage administered after 10 successive daily 1 mg/kg dosages of butorphanol resulted only in transient sedative effects. During the 10-day course of administration at 1 mg/kg (10 times the recommended use level) in two horses, the only detectable drug effects were transient behavioral changes typical of narcotic agonist activity. These included muscle fasciculation about the head and neck, dysphoria, lateral nystagmus, ataxia and salivation.

Repeated administration of butorphanol at 1 mg/kg (10 times the recommended dose) every four hours for 48 hours caused constipation in one of two horses.

**Subacute Equine Studies**

Horses were found to tolerate butorphanol given intravenously at dosages of 0.1, 0.3 and 0.5 mg/kg every 4 hours for 48 hours followed by once daily injections for a total of 21 days. The only detectable drug effects were slight transient ataxia observed occasionally in the high dosage group. No clinical, laboratory, or gross or histopathologic evidence of any butorphanol-related toxicity was encountered in the horses.

**INDICATIONS**
TORBUGESIC (butorphanol tartrate injection) is indicated for the relief of pain associated with colic in adult horses and yearlings. Clinical studies in the horse have shown that TORBUGESIC alleviates abdominal pain associated with torsion, impaction, intussusception, spasmotic and tympanic colic and postpartum pain.

WARNINGS
DO NOT USE IN HORSES INTENDED FOR HUMAN CONSUMPTION. NOT FOR HUMAN USE.

CAUTION
TORBUGESIC, a potent analgesic, should be used with caution with other sedative or analgesic drugs as these are likely to produce additive effects. There are no well-controlled studies using butorphanol in breeding horses, weanlings and foals. Therefore, the drug should not be used in these groups.

ADVERSE REACTIONS
In clinical trials in horses, the most commonly observed side effect was slight ataxia which lasted 3 to 10 minutes.
Marked ataxia was reported in 1.5% of the 327 horses treated. Mild sedation was reported in 9% of the horses.

DOSAGE
The recommended dosage in the horse is 0.1 mg of butorphanol per kilogram of body weight (0.05 mg/lb) by intravenous injection. This is equivalent to 5 mL of TORBUGESIC for each 1000 lbs body weight.
The dose may be repeated within 3 to 4 hours but treatment should not exceed 48 hours.
Use contents within 4 months of first puncture. Pre-clinical model studies and clinical field trials in horses demonstrate that the analgesic effects of TORBUGESIC are seen within 15 minutes following injection and persist for about 4 hours.

HOW SUPPLIED
50 mL vials TORBUGESIC (butorphanol tartrate injection) Veterinary Injection, 10 mg base activity per mL.
10 mL vials TORBUGESIC (butorphanol tartrate injection) Veterinary Injection, 10 mg base activity per mL.
Store at controlled room temperature 20°-25°C (68°-77°F) with excursions between 15°-30°C (59°-86°F).

REFERENCES


Distributed by:
Zoetis Inc.
Kalamazoo, MI 49007
Made in Spain
Revised: April 2018

**PRINCIPAL DISPLAY PANEL - 10 mL Bottle Label**

Torbugesic®
(butorphanol tartrate injection)
contains **10 mg** butorphanol base per mL as butorphanol tartrate

**CAUTION:** Federal (USA) law restricts this drug to use by or on the order of a licensed veterinarian.

NADA 135-780, Approved by FDA
10 mL
40022903
Zoetis
PRINCIPAL DISPLAY PANEL - 50 mL Carton Label

Torbugesic®
(butorphanol tartrate injection)
contains 10 mg butorphanol base per mL as butorphanol tartrate

CAUTION: Federal (USA) law restricts this drug to use by or on the order of a licensed veterinarian.
NADA 135-780, Approved by FDA
50 mL
Distributed by:
Zoetis Inc.
Kalamazoo, MI 49007
Made in Spain
40022907
TORBUGESIC
(butorphanol tartrate injection)

**Product Information**

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**Active Ingredient/Active Moiety**

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**Inactive Ingredients**

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### Marketing Information

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**Labeler** - Zoetis Inc. (828851555)

Revised: 10/2018