For Use in Cats Only

Simbadol
(buprenorphine injection)
1.8 mg/mL

For subcutaneous use in cats
Opioid Analgesic

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

HUMAN SAFETY WARNING
Abuse Potential
SIMBADOL contains buprenorphine (1.8 mg/mL), an opioid agonist and Schedule III controlled substance with an abuse potential similar to other Schedule III opioids. Buprenorphine has certain opioid properties that in humans may lead to dependence of the morphine type. Abuse of buprenorphine may lead to physical dependence or psychological dependence. The risk of abuse by humans should be considered when storing, administering, and disposing of SIMBADOL. Persons at increased risk for opioid abuse include those with a personal or family history of substance abuse (including drug or alcohol abuse or addiction) or mental illness (suicidal depression).

Life-Threatening Respiratory Depression
Respiratory depression, including fatal cases, may occur with abuse of SIMBADOL.

Additive CNS Depressant Effects
SIMBADOL has additive CNS depressant effects when used with alcohol, other opioids, or illicit drugs that cause central nervous system depression.

Accidental Exposure
Because of the potential for adverse reactions associated with accidental injection, SIMBADOL should only be administered by veterinarians or veterinary technicians who are trained in the handling of potent opioids.

See Human Safety for detailed information.

DESCRIPTION:
SIMBADOL is a clear, colorless to slightly yellow, sterile, injectable solution intended for subcutaneous administration for use in cats. Each milliliter of SIMBADOL contains 1.8 mg buprenorphine (equivalent to 1.94 mg buprenorphine hydrochloride), 50 mg anhydrous dextrose, 1.8 mg methylparaben, 0.2 mg propylparaben, 0.2 mg sodium acetate trihydrate, 0.5 mg glacial acetic acid, 100.0 mg anhydrous ethanol, water for injection, and hydrochloric acid and/or sodium hydroxide to adjust pH.

Buprenorphine belongs to the opioid class of drugs and is a narcotic under the Controlled Substances Act due to its chemical derivation from thebaine. Buprenorphine hydrochloride is a weakly acidic, white or off-white crystalline powder with limited solubility in water. Chemically, it is 17-(cyclopropylmethyl)-e-(1,1-dimethyltyethylenyl)-4,5-epoxy-18,19-dihydro-3-hydroxy-6-methoxy-7-methyl-14-ethenomorphinan-7-methanol, hydrochloride [5c, 7α(S)]. Buprenorphine hydrochloride has the molecular formula of C26H31NO5·HCl, the molecular weight of 504.09, and the following structural formula:

INDICATION:
SIMBADOL is indicated for the control of postoperative pain associated with surgical procedures in cats.

DOSE AND ADMINISTRATION:
The dosage of SIMBADOL is 0.24 mg/kg (0.11 mg/lb) administered subcutaneously once daily, for up to 3 days. Administer the first dose approximately 1 hour prior to surgery. Do not discontinue SIMBADOL for administration at home by the pet owner (see Human Safety).

CONTRAINDICATIONS:
SIMBADOL is contraindicated in cats with known hypersensitivity to buprenorphine hydrochloride or any of the components of SIMBADOL, or known intolerance to opioids.

WARNINGS:
For subcutaneous (SQ) injectable use in cats.

Human Safety:

Adult Human User Safety while Handling SIMBADOL in the Hospital:
Mucous membrane or eye contact during administration:
Direct contact of SIMBADOL with the eyes, oral or other mucous membranes could result in absorption of buprenorphine and the potential for adverse reactions. If accidental eye, oral or other mucous membrane contact is made during administration, flush the area with water and contact a physician.

Skin contact during administration:
If human skin is accidentally exposed to SIMBADOL, wash the exposed areas with soap and water and contact a physician. Accidental exposure could result in absorption of buprenorphine and the potential for adverse reactions.

Drug Abuse, Addiction, and Diversion of Opioids:
Controlled Substance:
SIMBADOL contains buprenorphine, a mu opioid partial agonist and Schedule III controlled substance with an abuse potential similar to other Schedule III opioids. SIMBADOL can be abused and is subject to misuse, abuse, addiction, and criminal diversion. SIMBADOL should be handled appropriately to minimize the risk of diversion, including restriction of access, the use of accounting procedures, and proper disposal methods, as appropriate to the clinical setting and as required by law.

Abuse:
Abuse of SIMBADOL poses a hazard of overdose and death. This risk is increased with concurrent abuse of alcohol and other substances including other opioids and benzodiazepines. Buprenorphine has been diverted for non-medical use into illicit channels of distribution. All people handling opioids require careful monitoring for signs of abuse. Drug abuse is the intentional non-therapeutic use of a prescription drug for its rewarding psychological or physiological effects. Abuse of opioids can result in the presence of true addiction.

Storage and Discard:
SIMBADOL is a Class III opioid. Store in a locked, substantially constructed cabinet according to DEA and local controlled substance guidelines. Discard baulched vials after 28 days. Any unused or expired vials must be destroyed by a DEA registered reverse distributor; for further information, contact your local DEA field office or call Zeotis Inc. at 1-888-965-8471.

Information for physician:
SIMBADOL injectable solution is a mu opioid partial agonist (1.8 mg buprenorphine/mL). In the case of an emergency, provide the physician with the package insert. Naloxone may not be effective in reversing respiratory depression produced by buprenorphine. The onset of naloxone effect may be delayed by 30 minutes or more. Doxapram hydrochloride has also been used as a respiratory stimulant.

PRECAUTIONS:
Hyperactivity (opioid excitation) has been observed up to 8 hours after anesthetic recovery (see ADVERSE REACTIONS).

Safety has not been evaluated in moribund cats (i.e., those not expected to live more than 24 hours with or without surgery). Use in such cases should be based on the risk-benefit assessment of the veterinarian.

Use with caution in cats with impaired hepatic function.

The use of SIMBADOL has not been evaluated in breeding, pregnant, or lactating cats, or in cats younger than 4 months of age.

ADVERSE REACTIONS:
In two controlled field studies, a total of 450 male and female cats 4 months to 16 years old, weighing between 2.6 – 20.0 lb were included in the field safety analysis. In one study, cats underwent a soft tissue surgical procedure (soft tissue). In the other study, cats underwent onychectomy, orchietomy and castration, or orchietomy and ovariohysterectomy (orthopedic). The following tables (one table for each study) show the number of cats exhibiting each observation.

Adverse Reactions in the Soft Tissue Field Study

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>SIMBADOL (N = 119)</th>
<th>Control (N = 112)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypotension</td>
<td>33 (33.6%)</td>
<td>29 (26.6%)</td>
</tr>
<tr>
<td>Tachycardia</td>
<td>26 (25.9%)</td>
<td>29 (26.6%)</td>
</tr>
<tr>
<td>Hyperthermia</td>
<td>≥98°F</td>
<td>27 (25.7%)</td>
</tr>
<tr>
<td>Hyperthermia</td>
<td>≥103°F</td>
<td>0</td>
</tr>
<tr>
<td>Hypertension</td>
<td>6 (5.4%)</td>
<td>20 (18.3%)</td>
</tr>
<tr>
<td>Bradycardia</td>
<td>5 (4.6%)</td>
<td>1 (0.9%)</td>
</tr>
<tr>
<td>Tachypnea</td>
<td>4 (3.6%)</td>
<td>2 (1.8%)</td>
</tr>
<tr>
<td>Arrhythmia</td>
<td>1 (0.9%)</td>
<td>0</td>
</tr>
<tr>
<td>Hyperactivity</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Blindness</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Apnea/Death</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

a. Cats may have experienced more than one type or occurrence of an adverse reaction. Cats experiencing the same reaction both during and after surgery are presented in both time periods.

b. During surgery is the time from the administration of the anesthetic induction agent until discontinuation of the surgery.

c. Hypertension is defined as a mean blood pressure of ≥160 mmHg during surgery and ≥30 mmHg after surgery.

d. Tachycardia is defined as a heart rate ≥180 beats per minute during surgery and ≥200 beats per minute after surgery.

e. Hyperthermia is defined as a core temperature ≥103.0°F.

Adverse Reactions in the Orthopedic Field Study

<table>
<thead>
<tr>
<th>Adverse Reaction</th>
<th>SIMBADOL (N = 115)</th>
<th>Control (N = 114)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tachycardia</td>
<td>29 (25.2%)</td>
<td>39 (35.8%)</td>
</tr>
<tr>
<td>Hypotension</td>
<td>25 (22.1%)</td>
<td>32 (28.3%)</td>
</tr>
<tr>
<td>Hyperthermia</td>
<td>≥103.0°F</td>
<td>0</td>
</tr>
<tr>
<td>Hyperthermia</td>
<td>≥98.0°F</td>
<td>8 (7.0%)</td>
</tr>
<tr>
<td>Hyperactivity</td>
<td>0</td>
<td>16 (14.0%)</td>
</tr>
</tbody>
</table>

Hyperactivity:
0 (0%)
After surgery, 27 out of 93 (29.0%) SIMBADOL cases and 57 out of 102 (55.9%) placebo treatment group (66/93 or 71.0%) compared to the placebo control group (45/102 or 44.1%) was observed. Twenty-seven out of 93 (an active metabolite) via CYP3A4. Buprenorphine and norbuprenorphine subsequently form via the bile into the gastrointestinal tract. Following subcutaneous injection in cats, there is considerable inter-cat variability in plasma concentration and pharmacokinetic parameters. Formulated as an immediate release product, buprenorphine is quickly absorbed following subcutaneous injection (e.g. mydriasis) may occur within minutes after injection. Buprenorphine plasma concentrations, following subcutaneous injection did not appear to correlate to pharmacodynamics measurements (change in the thermal threshold data). In studies with SIMBADOL anesthetic effects of buprenorphine appeared about one hour after injection with a 24 to 28 hour duration of action. Pharmacokinetically, buprenorphine studies have demonstrated a marked time delay between plasma concentrations and the onset and offset of the anesthetic effect which is due to the slow equilibration between drug concentrations in the biophase and the slow association and dissociation of drug binding to the receptor. Following subcutaneous injection in cats, there is considerable inter-cat variability in plasma concentration and pharmacokinetic parameters. Formulated as an immediate release product, buprenorphine is quickly absorbed following subcutaneous injection (e.g. mydriasis) may occur within minutes after injection. Buprenorphine plasma concentrations, following subcutaneous injection did not appear to correlate to pharmacodynamics measurements (change in the thermal threshold data). In studies with SIMBADOL anesthetic effects of buprenorphine appeared about one hour after injection with a 24 to 28 hour duration of action. Pharmacokinetically, buprenorphine studies have demonstrated a marked time delay between plasma concentrations and the onset and offset of the anesthetic effect which is due to the slow equilibration between drug concentrations in the biophase and the slow association and dissociation of drug binding to the receptor. 

**EFFECTIVENESS:**

The effectiveness of SIMBADOL was demonstrated in two randomized, masked, placebo-controlled, multi-site field studies involving client-owned cats of various breeds. In one study (soft tissue), 221 cats underwent a soft tissue surgical procedure. In the other study (orthopedic), 229 cats underwent a soft tissue surgical procedure. In the orthopedic field study, a statistically significant difference in pain intensity was observed in the SIMBADOL group compared to the placebo control group (45102 or 44.1%) was observed. Twenty-seven out of 93 (39.9%) SIMBADOL cases and 57 out of 102 (55.9%) placebo cases were treatment failures. In the orthopedic field study, a statistically significant difference in the proportion of treatment successes in the SIMBADOL treatment group (6693 or 71.0%) compared to the placebo control group (45102 or 44.1%) was observed. Twenty-seven out of 93 (39.9%) SIMBADOL cases and 69 out of 102 (67.6%) placebo cases were treatment failures. For both studies, the majority of the treatment failures required rescue within 4 hours after anesthetic recovery.

Combining both studies (450 cats), sedation was observed in 68 cats in the buprenorphine group and 62 cats in the placebo control group for up to 4 hours after anesthesia recovery. In both studies, during surgery, mean respiratory rates and mean blood pressures were lower in the buprenorphine group compared to the placebo control group. There were a higher number of cats with a higher incidence of increased respiratory rate compared to the placebo group (30 cats [28 incidences] compared to the placebo control group (8 cats, 10 incidences). The results of two field studies demonstrate that SIMBADOL is effective and has an acceptable safety margin for the control of postoperative pain in cats.

**ANIMAL SAFETY:**

**Nine-Day Target Animal Safety Study:** In a 9 day safety study, 4 month old healthy cats (4/sec/group) were administered SIMBADOL subcutaneously at 0X (saline), 1X (0.24 mg/kg), 3X (0.72 mg/kg), or 5X (1.2 mg/kg) once daily. All 32 cats survived to study termination. Buprenorphine-related clinical observations included difficulty in handling, lower incidence of urination, abnormal oral dryness, dilated pupils, and decreased pupillary light reflex. The incidence of temperatures ≤103°F was higher in the buprenorphine-treated groups compared to the control group. The highest temperature observed in the buprenorphine-treated group was 103.8°F in a 5X cat. One 1X cat and one 3X cat experienced an episode of hyperactivity, difficulty in handling, slight disorientation, ataxia, dilated pupils (which were responsive to light), and respiratory sinus arrhythmia. One 1X cat (one episode) and one 3X cat (three episodes) were reported with nystagmus. One 1X and one 3X cat were reported with decreased blink response (one episode). Three cats in the 5X group lost body weight (79 g or less) during the study which correlated with decreased food consumption. All other cats gained weight during the study.

The incidence of "moderate responses" (minor vocalization or wincing and quick resolution) and "severe responses" (tried to bite or scratch or had marked vocalization or persistent attention to the injection site) to injection was higher in the buprenorphine-treated groups compared to the control group. Respiratory rate, heart rate, and blood pressure were similar between all groups, including the control group.

Buprenorphine-related clinical pathology findings included an increase in creatine kinase values in the 3X and 5X groups and correlated with subcutaneous inflammation at the injection sites. Histologic lesions included minimal to moderate subcutaneous inflammation at the injection sites, which correlated with the administration of buprenorphine compared to the control group. The incidence of clinical signs similar between buprenorphine-treated groups; however, sites with mild and moderate inflammation were observed in the 5X group compared to the 1X and 3X groups where more sites with minimal inflammation were observed. Mineralization at an injection site was seen in one 1X and one 3X cat. Chronic inflammation in the heart (valve or myocardium) was seen in two 5X cats. Subcutis liver inflammation was seen in one control cat, two 1X cats, three 3X cats, and two 5X cats. Lymphoid hyperplasia of the mediastinal lymph node was seen in one 1X cat, and acute inflammation was seen in the mediastinal lymph node of one 3X cat. Lymphoid hyperplasia of the Peyer’s Patches was seen in two 1X cats and one 5X cat. Lymphoid hyperplasia, lymphocytic infiltrate, or subcutis inflammation of the stomach was seen in four 1X cats, four 3X cats, and three 5X cats. Subcutaneous lymphoid infiltrate of the thyroid glands was seen in two 1X cats, one 3X cat, and four 5X cats. Arterial Blood Pressure Study in Cats: Healthy 8.5 to 29.1 month old cats (4/sec/group) were subcutaneously administered SIMBADOL at 0.24 mg/kg (1X) or meloxicam (control), 1 hour prior to anesthetic induction for a 1 hour exploratory laparotomy. Arterial blood pressure was monitored following anesthetic induction and through laparotomy, with indirect blood pressure monitoring prior to anesthesia and for 8 hours following anesthetic recovery. All 16 animals were clinically healthy for the duration of the study. There were no differences between treatment groups in mean blood pressure during the study. During surgery and postoperatively, heart rate was higher for the buprenorphine group. During surgery, the incidence of heart rates ≤150 beats/minute was higher in the buprenorphine-treated group compared to the control group. Post-operatively, the incidence of heart rates ≤200 beats/minute was higher in the buprenorphine-treated group compared to the control group. During surgery, respiration rate was lower for the buprenorphine group. Post-operatively, body temperature was higher for the buprenorphine group. Four cats in the buprenorphine group had temperatures ≤104°F post-operatively compared to none in the control group. The highest temperature observed in the buprenorphine group was 104.3°F. Electrocardiograms were qualitatively normal in all cats. During surgery, one cat in the buprenorphine group had hemoglobin saturation less than 90% (88% at one time point).

**STORAGE INFORMATION:**

Store at temperatures up to 25°C (77°F). Protect from light and excessive heat (above 40°C or 104°F). Use within 28 days of first puncture.

**HOW SUPPLIED:**

**SIMBADOL (buprenorphine injection) is supplied in a carton containing one 10 mL amber glass vial with a multi-dose cartridge of 1.5 mg/mls, of buprenorphine. NADA 141-434, Approved by FDA**

**REFERENCES:**


Distributed by: Zoetis Inc. Kalamazoo, MI 49007

Product of United Kingdom

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