

RILEY'S SEDATION IS QUICK
AND SMOOTH AND HER
RECOVERY WILL BE TOO,
CREATING EXTRA TIME AND A
MORE PEACEFUL ENVIRONMENT
FOR YOU AND YOUR TEAM



Introducing Zenalpha® (medetomidine and vatinoxan hydrochlorides injection)

An innovative alpha-2 combination drug that improves
the sedation experience from start to finish

Zennovation in Sedation

For use in dogs only.
Not intended for use in cats.

 **Zenalpha®**
(medetomidine and vatinoxan
hydrochlorides injection)

WHY THE HESITATION?

Concerns about alpha-2 adrenoceptor agonists

- Sedation is an important tool in patient comfort and care, but it tends to be underutilized
- While alpha-2 adrenoceptor agonists are commonly used in small animal sedation and anesthesia, the binding of these agonists to peripheral receptors has undesirable effects
- Concerns about side effects and patient safety have prevented their full adoption into veterinary practice^{1,2}



Procedures that may require sedation

- Radiographic examination
- Diagnostic imaging
- Ear examination and treatment
- Eye examination and treatment
- Dermatological examination and procedures
- Dental examination
- Dental biopsy
- Minor tooth extraction
- Nail trimming
- Fine needle aspiration/superficial biopsy
- Anal sac treatment
- Minor surgery to remove dermal masses
- Seroma or abscess drainage
- Intravenous catheter placement
- Coat grooming
- Venous blood draw

Due to the pronounced cardiovascular effects of alpha-2 adrenoceptor agonists, only clinically healthy dogs (American Society of Anesthesiologists [ASA] classes I and II) should be administered Zenalpha.



Introducing Zenalpa® (medetomidine and vatinoxan hydrochlorides injection)

- Zenalpa is an innovative alpha-2 agonist-antagonist that minimizes the stress surrounding sedation, while maximizing procedural efficiency
- Zenalpa is effective in a broad range of procedures
- Zenalpa is indicated for use as a sedative and analgesic in dogs to facilitate clinical examinations, clinical procedures, and minor surgical procedures

ZENNOVATION IN SEDATION

Unique mechanism of action that helps minimize cardiovascular effects

- Zenalpa contains 0.5 mg/mL medetomidine hydrochloride and 10 mg/mL vatinoxan hydrochloride
 - Medetomidine is an alpha-2 **agonist** responsible for sedative and analgesic properties
 - Vatinoxan is an alpha-2 **antagonist**, acting in the periphery without crossing the blood-brain barrier



Selectively binding to peripheral alpha-2 adrenoceptors, vatinoxan has limited penetration across the blood-brain barrier into the central nervous system.²

 Vatinoxan  Medetomidine

As with all drugs, side effects may occur. Tachycardia may occur in some dogs after recovery from sedation. The following adverse reactions have been reported: diarrhea, muscle tremors, and colitis.

 **Zenalpa®**
(medetomidine and vatinoxan
hydrochlorides injection)

AN IMPROVED SEDATION MONITORING EXPERIENCE

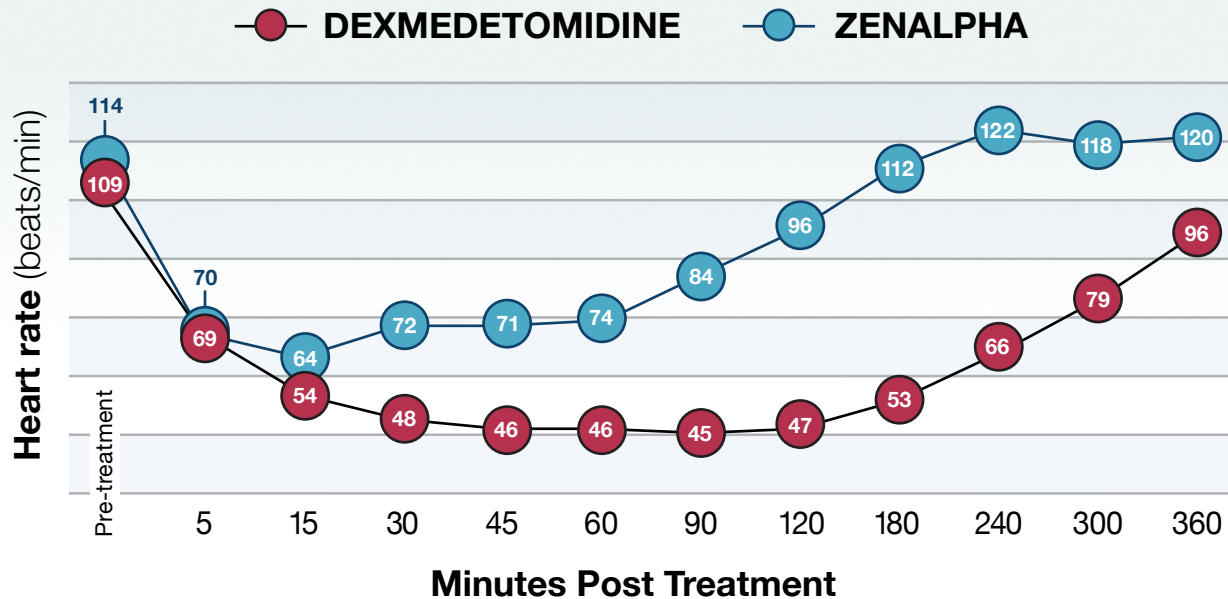
Field effectiveness study results (N = 223)

- Compared to the control group,* Zenalpa® (medetomidine and vatinoxan hydrochlorides injection):
 - Minimizes unwanted side effects while preserving reliable sedation and analgesia
 - Has a shorter time to onset of sedation and a shorter duration of sedation with quicker recovery²
 - Keeps the patient's heart rate and blood pressure closer to normal
- If needed, Zenalpa can be reversed with atipamezole hydrochloride



HEART RATE CLOSER TO NORMAL WITH ZENALPHA VS DEXMEDETOMIDINE

Group Mean Heart Rate (beats per minute)



Normal 60-140 beats/min

*Control = dexmedetomidine

Dogs should be monitored frequently during sedation for changes in heart rate, blood pressure, respiratory rate and body temperature.

Get your patients up and out the door quickly

- Zenalpha takes effect quickly, within **5-15 minutes*** in most dogs, and patients recover within approximately **45 minutes***
- Compared to the control group†:
 - Zenalpha has a shorter time to onset and a shorter duration of sedation in most dogs
 - Patients recover more rapidly and can return home sooner
 - There is minimal vomiting with Zenalpha
 - Zenalpha resulted in fewer severe cardiovascular side effects

INCREASES PROCEDURAL EFFICIENCY

TIME TO ONSET
5-15 Minutes



in most cases*

DURATION OF SEDATION
~45 Minutes



in most cases*



*Field study means were 14 minutes for time to onset of sedation and 38 minutes for duration of sedation.

†Control = dexmedetomidine

Do not use Zenalpha in dogs with cardiac disease, respiratory disorders, shock, severe debilitation, that have hypoglycemia or are at risk of developing hypoglycemia, or are stressed due to extreme heat, cold or fatigue.

 **Zenalpha®**
(medetomidine and vatinoxan hydrochlorides injection)

LESS STRESS FOR VETERINARY TEAMS

Gain more time back in your day



Zenalpha® (medetomidine and vatinoxan hydrochlorides injection) increases procedural efficiency due to its rapid onset and short duration of action



Due to the quick recovery time, the use of reversal agents may not be needed, which can save time and decrease costs



Reducing time to complete procedures and sending dogs home sooner can give veterinary teams time back in their day



DOSAGE AND ADMINISTRATION:

The dose is based on body surface area (BSA). Calculate the dose using 1 mg medetomidine/m² BSA or use the dosing table below. Note that the mg/kg dosage decreases as body weight increases.

Table 1. IM dose volume based on body weight

The onset of sedative/analgesic effects in nervous, excited or agitated dogs could be slowed, or the depth and duration of effects could be diminished or nonexistent. Therefore, allow the dog to rest quietly for 10-15 minutes after injection.

Dog body weight		Dose volume
lbs	kg	mL
4.4 to 7	2 to 3	0.3
7.1 to 9	3.1 to 4	0.4
9.1 to 11	4.1 to 5	0.6
11.1 to 22	5.1 to 10	0.8
22.1 to 29	10.1 to 13	1.0
29.1 to 33	13.1 to 15	1.2
33.1 to 44	15.1 to 20	1.4
44.1 to 55	20.1 to 25	1.6
55.1 to 66	25.1 to 30	1.8
66.1 to 73	30.1 to 33	2.0

Dog body weight		Dose volume
lbs	kg	mL
73.1 to 81	33.1 to 37	2.2
81.1 to 99	37.1 to 45	2.4
99.1 to 110	45.1 to 50	2.6
110.1 to 121	50.1 to 55	2.8
121.1 to 132	55.1 to 60	3.0
132.1 to 143	60.1 to 65	3.2
143.1 to 154	65.1 to 70	3.4
154.1 to 176	70.1 to 80	3.6
> 176	> 80	3.8

Zenalpha should not be administered in the presence of pre-existing hypotension, hypoxia or bradycardia.

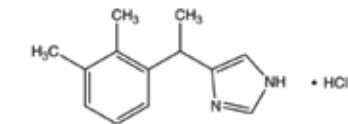
 **Zenalpha®**
(medetomidine and vatinoxan
hydrochlorides injection)

Zenapha®
(medetomidine and vatinoxan hydrochlorides injection)
0.5 mg/mL and 10 mg/mL
For intramuscular injection in dogs only
Sedative, Analgesic

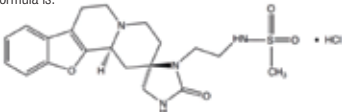
CAUTION:
Federal law restricts this drug to use by or on the order of a licensed veterinarian.

DESCRIPTION:
Zenapha is a combination of medetomidine and vatinoxan hydrochlorides. Medetomidine is a racemic mixture containing the active enantiomer, dexmedetomidine, an alpha₂-adrenoceptor agonist with sedative and analgesic properties. Vatinoxan is a peripherally selective alpha₂-adrenoceptor antagonist, which partially counteracts the cardiovascular depressive effects of dexmedetomidine at peripheral alpha₂-adrenoceptors, while preserving the centrally mediated sedative and analgesic effects of dexmedetomidine.

Medetomidine hydrochloride has the chemical name (S,R)-4-[1-(2,3-dimethylphenylethyl)-1H-imidazole] hydrochloride. It is a white, crystalline, water soluble substance having a molecular weight of 237 g/mol. The molecular formula is C₁₇H₁₉N₂·HCl and the structural formula is:



Vatinoxan hydrochloride has the chemical name N-(2-((2R,12bS)-2'-oxo-1,3,4,6,7,12b-hexahydrospiro[benzofuro[2,3-a]quinolizine-2,4'-imidazolidine]-3'-yl) ethyl) methane sulfonamide hydrochloride. It is a white to pale yellow, crystalline substance, sparingly soluble in water and having a molecular weight of 455 g/mol. The molecular formula is C₂₀H₂₄N₆O₄·HCl and the structural formula is:



Each mL of Zenapha contains 0.5 mg medetomidine hydrochloride, 10 mg vatinoxan hydrochloride, 32.5 mg mannitol (USP), 4.16 mg citric acid monohydrate (USP), 1.8 mg methylparaben (NF), 0.2 mg propylparaben (NF).

INDICATION:
Zenapha is indicated for use as a sedative and analgesic in dogs to facilitate clinical examination, clinical procedures and minor surgical procedures.
Zenapha is not intended for use in cats (see Animal Safety Warnings).

DOSAGE AND ADMINISTRATION:
The dose is based on body surface area (BSA). Calculate the dose using 1 mg medetomidine /m² BSA or use the dosing table below. **Note that the mg/kg dosage decreases as body weight increases.**

Table 1. IM dose volume based on body weight

Dog body weight		Dose volume
lbs	kg	mL
4.4 to 7	2 to 3	0.3
7.1 to 9	3.1 to 4	0.4
9.1 to 11	4.1 to 5	0.6
11.1 to 22	5.1 to 10	0.8
22.1 to 29	10.1 to 13	1.0
29.1 to 33	13.1 to 15	1.2
33.1 to 44	15.1 to 20	1.4
44.1 to 55	20.1 to 25	1.6
55.1 to 66	25.1 to 30	1.8
66.1 to 73	30.1 to 33	2.0
73.1 to 81	33.1 to 37	2.2
81.1 to 99	37.1 to 45	2.4
99.1 to 110	45.1 to 50	2.6
110.1 to 121	50.1 to 55	2.8
121.1 to 132	55.1 to 60	3.0
132.1 to 143	60.1 to 65	3.2
143.1 to 154	65.1 to 70	3.4
154.1 to 176	70.1 to 80	3.6
>176	>80	3.8

After administration of Zenapha, the dog should be allowed to rest quietly until evidence of sedation has occurred (5-15 minutes). The average duration of sedation is 38 minutes. As with all alpha₂-adrenoceptor agonists, onset of sedation may be delayed or may be inadequate in some dogs.

Reversal of Zenapha: Administration of IM atipamezole hydrochloride at the approved dose to reverse IM medetomidine hydrochloride results in reversal of the sedative and cardiovascular effects of Zenapha. Reversal of sedation occurs within 5-10 minutes after administration of atipamezole hydrochloride.

CONTRAINDICATIONS:
Do not use Zenapha in dogs with cardiac disease, respiratory disorders, shock, severe debilitation, that have hypoglycemia or are at risk of developing hypoglycemia, or are stressed due to extreme heat, cold or fatigue.

Zenapha is contraindicated in dogs with a known sensitivity to medetomidine or vatinoxan.

WARNINGS:
Human User Safety Warnings

Not for use in humans. Keep this and all medications out of reach of children and pets. Avoid skin, eye or mucosal contact. Use caution while handling and using filled syringes. Absorption of the active ingredients is possible following exposure via the skin, eye or mucosa. In case of accidental eye exposure, flush eyes with water for 15 minutes, remove contact lenses then continue to flush. In case of accidental skin exposure, wash with soap and water and remove contaminated clothing. If symptoms occur, seek the advice of a physician.

In case of accidental oral intake or self-injection, seek medical advice immediately and show the package insert to the physician. DO NOT DRIVE as sedation, loss of consciousness, and changes in blood pressure may occur.

Persons with cardiovascular disease (for example, hypertension or ischemic heart disease) should take special precautions to avoid any exposure to this product.

Pregnant women should exercise special caution to avoid exposure. Uterine contractions and decreased fetal blood pressure may occur after accidental systemic exposure.

Persons with known hypersensitivity to any of the ingredients should avoid contact with Zenapha.

Caution should be exercised when handling sedated animals. Handling or any other sudden stimuli, including noise, may cause a defense reaction in an animal that appears to be heavily sedated.

Note to physician: Zenapha contains medetomidine, an alpha₂-adrenoceptor agonist, in combination with vatinoxan, a peripherally selective alpha₂-adrenoceptor antagonist. Symptoms after absorption or accidental self-injection may include dose-dependent sedation, respiratory depression, bradycardia, tachycardia, and hypotension.

Animal Safety Warnings
Zenapha should not be administered in the presence of pre-existing hypotension, hypoxia or bradycardia. Due to the pronounced cardiovascular effects of alpha₂-adrenoceptor agonists, only clinically healthy dogs (American Society of Anesthesiologists [ASA] classes I and II) should be administered Zenapha. Dogs should be monitored frequently for cardiovascular function and body temperature during sedation.

Zenapha is not intended for use in cats. The use of Zenapha in cats has been associated with hypotension.

PRECAUTIONS:
Dogs should be monitored frequently during sedation for changes in heart rate, blood pressure, respiratory rate and body temperature. Tachycardia may occur in some dogs after recovery from sedation.

In the event of hypoxia or apnea, supplemental oxygen should be administered. Following administration of Zenapha, a decrease in body temperature may occur and an external heat source may be needed to maintain body temperature. Hypothermia may persist longer than sedation and analgesia.

The analgesic effect of Zenapha will not last longer than the sedative effects. Additional analgesic(s) should be administered as needed (see Effectiveness).

Nervous, excited or agitated dogs with high levels of endogenous catecholamines may exhibit a reduced pharmacological response to Zenapha (ineffectiveness). The onset of sedative/analgesic effects could be slowed, or the depth and duration of effects could be diminished or nonexistent. Therefore, allow the dog to rest quietly for 10 to 15 minutes after injection.

With the alpha₂-adrenoceptor agonist drug class, including Zenapha, the potential for isolated cases of hypersensitivity, including paradoxical response (excitation) exists. Repeat dosing with Zenapha has not been evaluated.

Zenapha has only been evaluated in fasted dogs; therefore, the effects on fed dogs (for example occurrence of vomiting) has not been characterized.

The concurrent use of anticholinergic medications and Zenapha has not been evaluated.

Zenapha may decrease serum glucose in healthy dogs and this effect may persist longer than sedation (see Animal Safety).

The safe use of Zenapha in dogs with hepatic or renal impairment has not been evaluated. The safe use of Zenapha has not been evaluated in dogs younger than 4.5 months old. The safe use of Zenapha has not been evaluated in dogs that are pregnant, lactating, or intended for breeding.

ADVERSE REACTIONS:
In the field study safety evaluation, 110 dogs received Zenapha and 113 dogs received dexmedetomidine (control group) for sedation. Dogs ranged in age from 5 months to 14.5 years, weighed 5.1 to 154 lbs, and represented purebreds and breed mixes.

Table 2. Adverse reactions during the field study

Adverse Reaction	Zenapha (N = 110) n (%)	Dexmedetomidine (N = 113) n (%)
Diarrhea	4 (3.6)	0 (0)
Muscle tremor	2 (1.8)	0 (0)
Signs of colitis	2 (1.8)	0 (0)
Hypothermia*	1 (0.9)	13 (11.5)
Vomiting	1 (0.9)	6 (5.3)
Involuntary defecation	1 (0.9)	0 (0)
Nausea	1 (0.9)	0 (0)
Tachycardia, transient	1 (0.9)	0 (0)
Prolonged sedation	0 (0)	3 (2.7)
Urinary incontinence	0 (0)	2 (1.8)
Retching	0 (0)	1 (0.9)
Apnea	0 (0)	1 (0.9)
Bradycardia	0 (0)	1 (0.9)
Hyperthermia	0 (0)	1 (0.9)

*Hypothermia that necessitated use of an external heat source

Field Study Safety
Decreased body temperature occurred within 30 minutes after administration for some dogs in both treatment groups and lasted up to 2 hours in the Zenapha group and 2-6 hours in the control group. There were 57/110 dogs (51.8%) in the Zenapha group and 77/113 dogs (68.1%) in the control group that had a body temperature ≤99 °F. The respiratory rates decreased in both groups within 5 minutes post dose compared to baseline. The group mean respiratory rate in the Zenapha group decreased earlier and recovered faster than the control group. Thirty-nine dogs out of 110 (35.4%) in the Zenapha group and 33/113 dogs (29.2%) in the control group had respiratory rates < 10 breaths per minute (bpm) during the study. The group mean respiratory rate in the Zenapha group was lowest at 30 minutes post-treatment (14 bpm) and in the control group was lowest at 90 minutes post-treatment (15.5 bpm). Mucous membrane color and capillary refill time were normal for all dogs at all timepoints during the study.

CONTACT INFORMATION:
To report suspected adverse reactions, for technical assistance, or to obtain a copy of the Safety Data Sheet (SDS), call Dechra at (866) 933-2472. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or www.fda.gov/reportanimalae.

CLINICAL PHARMACOLOGY:
Medetomidine is a potent non-narcotic alpha₂-adrenoceptor agonist which produces sedation and analgesia. These effects are dose dependent in depth and duration. Medetomidine is a racemic mixture containing the active enantiomer dexmedetomidine. Within the central nervous system, sympathetic neurotransmission is inhibited and the level of consciousness decreases. Respiratory rate and body temperature can also decrease. In the peripheral vasculature, medetomidine stimulates alpha₂-adrenoceptors within vascular smooth muscle which induces vasoconstriction and hypertension which consequently decreases the heart rate and cardiac output. Dexmedetomidine also induces a number of other alpha₂-adrenoceptor mediated effects, which include piloerection, depression of motor and secretory functions of the gastrointestinal tract, diuresis and hyperglycemia.

Vatinoxan is a peripherally selective alpha₂-adrenoceptor antagonist which lacks activity in the central nervous system. By limiting its effects to peripheral organs

systems, vatinoxan will prevent or attenuate the cardiovascular and other effects of dexmedetomidine outside the central nervous system when administered simultaneously with the alpha₂-adrenoceptor agonist. The central effects of dexmedetomidine remain unaltered, although vatinoxan will reduce the duration of sedation and analgesia induced by dexmedetomidine, predominantly by increasing the clearance of the latter via improving the cardiovascular function.

No pharmacokinetic assessment was performed on Zenapha [medetomidine (1 mg/m²) + vatinoxan (20 mg/m²)]. However, after IM administration of a pilot formulation of medetomidine (1 mg/m²) + vatinoxan (30 mg/m²), both medetomidine and vatinoxan were rapidly and highly absorbed from the injection site. Maximal plasma concentration was reached at 12.6 ± 4.7 (mean ± standard deviation) minutes and 17.5 ± 7.4 minutes for dexmedetomidine (the active enantiomer of medetomidine) and vatinoxan, respectively. Vatinoxan increased the volume of distribution and the clearance of dexmedetomidine. Thus, the clearance of dexmedetomidine was increased two-fold when given in combination with vatinoxan. The same phenomena were also observed with intravenous administration.

Medetomidine plasma protein binding is high (85-90%). Medetomidine is mainly oxidized in the liver, a smaller amount undergoes methylation in the kidneys, and excretion is mainly via urine. Vatinoxan plasma protein binding is approximately 70%. Low levels are detectable in the central nervous system. Only a small amount (<5%) of vatinoxan dose has been found to be excreted via the urine.

EFFECTIVENESS:
A prospective, randomized, masked, multi-center study was conducted at six veterinary clinics to evaluate the effectiveness of Zenapha to provide sedation and analgesia for non-painful or mildly painful, non-invasive procedures and examinations in client-owned dogs. Effectiveness was evaluated in 208 of the 223 enrolled dogs (109 in the Zenapha group and 99 in the control group). Dogs ranged in age from 5 months to 14.5 years, weighed 5.1 to 154 lbs (2.3-70 kg) and represented purebreds and breed mixes. Dogs received one IM injection of Zenapha at 1 mg/m² medetomidine and 20 mg/m² vatinoxan, or dexmedetomidine at 0.5 mg/m² (control group) to produce sedation prior to conduct of the examination or procedure. Procedures included: radiographic examination or diagnostic imaging, ear examination and treatment, eye examination and treatment, anal sac treatment, dermatological examination and procedures, dental examination, dental biopsy, dental minor extraction, fine needle aspiration/superficial biopsy, minor surgery to remove dermal masses, drain seroma or abscess, nail trimming, coat grooming, venous blood draw and intravenous catheter placement. Many dogs underwent more than one procedure while sedated.

Effectiveness of Zenapha was based on: 1) the ability to complete the planned examination or procedure while the dog was sedated, and 2) demonstration of less severe cardiovascular adverse effects (determined by heart rate) compared to the control group. Respiratory rate, body temperature, analgesia, and adverse reactions were evaluated as secondary variables for the safety assessment (see Adverse Reactions). Dogs were evaluated at 5, 15, 30, 60, and 90 minutes and 2, 3, 4, 5 and 6 hours post-treatment. The success rate for ability to complete the procedure was 94.5% (103/109) in the Zenapha group and 90.9% (90/99) in the control group. The Zenapha group mean heart rate remained within the normal range (60-140 bpm), while the control group mean heart rate was below normal from 5-180 minutes post-treatment. The mean heart rate in the Zenapha group was lowest at 15 minutes (64 bpm) and the mean heart rate in the control group was lowest at 90 minutes (45 bpm). There were 9 dogs (8.3%) in the Zenapha group and 54 dogs (52.4%) in the control group that had heart rates <40 beats per minute (bpm) during sedation. Zenapha met the criteria for success because it was not inferior to the control product for sedation and completion of the procedure and provided a decrease in the occurrence of bradycardia.

Dogs in the Zenapha group typically had a shorter time to onset of sedation and shorter duration of sedation compared to the control group. The mean time to onset of sedation was 14 minutes in the Zenapha group and 18 minutes in the control group. The mean duration of sedation in the Zenapha group was 38 minutes (maximum of 90 min.) and in the control group was 90 minutes (maximum 5 hrs. 28 min.). The analgesic effects in the Zenapha and the control groups were similar, and both waned with the loss of sedation.

The results of the field study demonstrated that Zenapha is safe and effective, and provides sedation and analgesia needed to perform common non-painful or mildly painful veterinary examinations and procedures. Dogs treated with Zenapha had a shorter time to onset of sedation, shorter duration of sedation (recovered quicker), and less cardiovascular and respiratory depression and adverse reactions compared to dogs in the control group.

ANIMAL SAFETY:
In a 4-day laboratory study using final market formulation, 32 healthy Beagle dogs (4 dogs/sex/group) aged 4.5 - 6.5 months were administered saline control, 1/20 medetomidine/vatinoxan, 3/60 medetomidine/vatinoxan, or 5/100 medetomidine/vatinoxan mg/m² Body Surface Area (BSA) by intravenous injection. These Zenapha doses correspond to 1, 3, or 5X the recommended intramuscular dose. The administration of Zenapha resulted in sedation, hypotension, hypothermia, and initially sinus bradycardia followed later by sinus tachycardia as the dogs recovered from sedation. Generally, dogs were provided supplemental heat support starting at 1 or 2 hours post-dose and continuing through 4 hours post-dose. Dogs recovered from sedation within 4 hours for the 1X group and 4-8 hours for the 3 and 5X groups. Physiologic variables returned to baseline within 8 hours post-dose in all dose groups. Bloodwork was evaluated 3-4 hours post-dose. Two dogs in the 5X group had decreased blood glucose on post-dose bloodwork. One dog in the 1X group and one dog in the 3X group had decreased potassium on post-dose bloodwork. Clinical observations included mucoid, soft, or watery feces and observations of salivation, trembling, tremors, vocalization, defecation or vomiting after dosing, injected sclera, struggling after dosing, and skin cool to touch. There were no Zenapha-related effects on morbidity, body weight, ophthalmological examinations, mucous membrane color, capillary refill time, gross pathology observations, organ weights, or histopathological findings.

STORAGE INFORMATION:
Store below 77°F (25°C).
In use shelf life: 3 months at 77°F (25°C).
Protect from freezing and store in outer carton to protect from light.

HOW SUPPLIED:
Zenapha is supplied in 10 mL multi-dose glass vials. Each mL contains 0.5 mg medetomidine hydrochloride and 10 mg vatinoxan hydrochloride.
NDC 17033-090-05

Approved by FDA under NADA # 141-551



Manufactured for:
Dechra Veterinary Products
7015 College Boulevard, Suite 525
Overland Park, KS 66211 USA

Manufactured in Sweden.
©2021 Dechra Limited
Zenapha is a registered trademark of Dechra Limited;
all rights reserved.
Rev. November 2021



HELPS IMPROVE THE SEDATION EXPERIENCE FROM START TO FINISH



Minimize the stress surrounding sedation. Maximize procedural efficiency.

- Zenalpa® (medetomidine and vatinoxan hydrochlorides injection) is used as a sedative and analgesic in dogs to facilitate clinical examinations, clinical procedures, and minor surgical procedures
- Combination of medetomidine and vatinoxan helps minimize unwanted cardiovascular side effects while preserving reliable sedation and analgesia
- When compared to the control,* Zenalpa has a shorter time to onset (5-15 minutes) and a shorter duration of sedation (~45 minutes) in most dogs
- Increases procedural efficiency and is effective for a broad range of procedures



**FOR MORE INFORMATION,
VISIT WWW.ZENALPHA.US**

 **Zenalpa®**
(medetomidine and vatinoxan
hydrochlorides injection)

IMPORTANT SAFETY INFORMATION

As with all drugs, side effects may occur. For use in dogs only. Not intended for use in cats. Not for use in humans. Avoid skin, eye or mucosal contact. In case of accidental oral intake or self-injection, seek medical advice immediately and show the package insert to the physician. DO NOT DRIVE as sedation, loss of consciousness, and changes in blood pressure may occur. People with cardiovascular disease and pregnant women should exercise special caution to avoid exposure. Uterine contractions and decreased fetal blood pressure may occur after accidental systemic exposure. Do not use Zenalpa in dogs with cardiac disease, respiratory disorders, shock, severe debilitation, that have hypoglycemia or are at risk of developing hypoglycemia, or are stressed due to extreme heat, cold or fatigue. Zenalpa should not be administered in the presence of pre-existing hypotension, hypoxia or bradycardia. Due to the pronounced cardiovascular effects of alpha2-adrenoceptor agonists, only clinically healthy dogs (American Society of Anesthesiologists [ASA] classes I and II) should be administered Zenalpa. Dogs should be monitored frequently during sedation for changes in heart rate, blood pressure, respiratory rate and body temperature. Tachycardia may occur in some dogs after recovery from sedation. The following adverse reactions have been reported: diarrhea, muscle tremors and colitis. Refer to the prescribing information for complete details or visit www.dechra-us.com.

References: 1. Murrell JC, Hellebrekers LJ. Medetomidine and dexmedetomidine: a review of cardiovascular effects and antinociceptive properties in the dog. *Vet Anaesth Analg*. 2005;32(3):117-127.
2. Johnson RA, Kennedy MJ. Dexmedetomidine and atipamezole. *Clinician's Brief*. August 2015:65-67.

*Control = dexmedetomidine

Approved by FDA under ANADA # 141-551.

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

Dechra is a registered trademark of Dechra Pharmaceuticals PLC.

Zenalpa is a registered trademark of Dechra Limited; all rights reserved.

© 2022 Dechra Veterinary Products.

07SD-ZEN2016-0622

