Medetomidine Hydrochloride
(medetomidine hydrochloride)

Sterile Injectable Solution
1.0 mg/mL

ANADA 200-610, Approved by FDA
Sedative and Analgesic
For intramuscular and intravenous use in dogs only
Modern Veterinary Therapeutics

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

DESCRIPTION
Medetomidine Hydrochloride (medetomidine hydrochloride) is a synthetic α₂-adrenoreceptor agonist with sedative and analgesic properties. The chemical name is (±)-4-[(2,3-dimethylphenyl)ethyl]-1H-imidazol-1-α hydroxide, up to 5.0 mg hydrochloric acid 0.1N, water for injection, q.s.

CLINICAL PHARMACOLOGY
Medetomidine is a potent non-narcotic α₂-adrenoreceptor agonist which produces sedation and analgesia. These effects are dose dependent in depth and duration. Profound sedation and recumbency, with reduced sensitivity to environmental stimuli (sounds, etc.), are seen with medetomidine.

The pharmacological restraint and pain relief provided by medetomidine facilitates handling and aids in the conduct of diagnostic or therapeutic procedures. It also facilitates minor surgical procedures (with or without local anesthesia) and dental care where intubation is not required. Spontaneous muscle contractions (twitching) can be expected in some dogs during medetomidine. With medetomidine administration, blood pressure is initially increased due to peripheral vasoc constriction and thereafter drops to normal or slightly below normal levels. The initial preoperative response is accompanied by a compensatory marked decrease in heart rate mediated by a vagal baroreceptor mechanism. The bradycardia may be partially prevented by prior (at least 5 minutes before) intravenous administration of an anticholinergic agent (see PRECAUTIONS). A transient change in the conductivity of the cardiac muscle may occur, as evidenced by arrhythmia in some animals. Cardiovascular changes (such as profound bradycardia and second degree heart block) equally affect both heartworm negative and asymptomatic heartworm positive dogs.

Respiratory responses include an initial slowing of respiration within a few seconds to 1–2 minutes after administration, increasing to normal within 120 minutes. An initial decrease in tidal volume is followed by an increase. When medetomidine was given at 3 and 5 times the recommended dose IV, and 5 and 10 times IM, effects were not intensified but were prolonged.

INDICATIONS
Medetomidine Hydrochloride is indicated for use as a sedative and analgesic in dogs over 12 weeks of age to facilitate clinical examinations, clinical procedures, minor surgical procedures not requiring muscle relaxation, and minor dental procedures where intubation is not required. The IV route of administration is more efficacious for dental care.

WARNINGS

Medetomidine hydrochloride can be absorbed and may cause irritation following direct exposure to skin, eyes, or mouth. In case of accidental eye exposure, flush with water for 15 minutes. In case of accidental skin exposure, wash with soap and water. Remove contaminated clothing. If irritation or other adverse reaction occurs (e.g., sedation, hypotension, bradycardia), seek medical attention.

In case of accidental oral exposure or injection, seek medical attention. Precaution should be used while handling and using filled syringes. Use of this drug in patients with cardiovascular disease (e.g., hypertension or ischemic heart disease) should take special precautions to avoid any exposure to this product.

The safety data sheet (SDS) contains more detailed occupational safety information. To report suspected adverse drug events, contact FDA at 1-888-FDA-VETS or online at http://www.fda.gov/AnimalVeterinary/SafetyHealth.

NOTE TO PHYSICIANS
This product contains an α₂-adrenoreceptor agonist.

CONTRA-INDICATIONS
Medetomidine Hydrochloride should not be used in dogs with the following conditions: cardiac disease, respiratory disorders, liver or kidney diseases, dogs in shock, dogs which are severely dehydrated, or dogs which are stressed due to extreme heat, cold or fatigue.

PRECAUTIONS
In extremely nervous or excited dogs, levels of endogenous catecholamines are high due to the animal’s state of agitation. The pharmacological response elicited by α₂-agonists (e.g., medetomidine) in such animals is often reduced in depth and duration. Sedative and analgesic effects ranging from slightly diminished to nonexistent. Highly agitated dogs should therefore be put at ease and allowed to rest quietly prior to receiving Medetomidine Hydrochloride. Allowing dogs to rest quietly for 10 to 15 minutes after injection may improve the response to Medetomidine Hydrochloride. In dogs not responding satisfactorily to treatment with Medetomidine Hydrochloride, repeat dosing is not recommended. Caution should be exercised when handling sedated animals. Handling or any other sudden stimulus may cause a decrease in reaction in an animal that is sedated. Medetomidine Hydrochloride is a potent α₂-agonist which should be used with caution with other adrenergic or sympathomimetic drugs. Additive or synergistic effects are likely, possibly resulting in overdose. Although bradycardia may be partially prevented by prior (at least 5 minutes) intravenous administration of an anticholinergic agent, the administration of anticholinergic agents to treat bradycardia either simultaneously or following sedation with medetomidine could lead to adverse cardiovascular effects.

Special care is recommended when treating very young animals and older animals. Information on the possible reproductive effects of medetomidine is limited; therefore, the drug is not recommended for use in dogs used for breeding purposes or in pregnant dogs.

ADVERSE REACTIONS
As with all α₂-agonists, the potential for isolated cases of hypersensitivity, including paradoxical response (excitation), exists. Incidents of prolonged sedation, bradycardia, cyanosis, vomiting, apnea, death from circulatory failure with severe congestion of lungs, liver, kidney and recrudescence of sedation after initial recovery have been reported.

SIDE EFFECTS
Bradycardia with occasional atrioventricular blocks will occur together with decreased respiratory rates. Body temperature is slightly or moderately decreased. Urination typically occurs during recovery at about 90 to 120 minutes posttreatment. In approximately 10% of treated dogs, occasional episodes of vomiting occur between 15 to 10 minutes posttreatment. An increase in blood glucose concentration is seen due to α₂-adrenoreceptor-mediated inhibition of insulin secretion.

ANIMAL SAFETY
In target animal safety studies, medetomidine was tolerated in dogs at up to 5 times the recommended IV dose and up to 10 times the recommended IM dose. Additive or synergistic effects are likely, possibly resulting in overdose. Although bradycardia may be partially prevented by prior (at least 5 minutes) intravenous administration of an anticholinergic agent, the administration of anticholinergic agents to treat bradycardia either simultaneously or following sedation with medetomidine could lead to adverse cardiovascular effects.

DOSE AND ADMINISTRATION
Medetomidine Hydrochloride should be administered at the rate of 150 mcg IV or 1,000 mcg IM per square meter of body surface. Use the table below to determine the correct dosage on the basis of body weight.

<table>
<thead>
<tr>
<th>Body Weight (lb) (IB EXCLUSION)</th>
<th>Injection Volume (mL)</th>
<th>Body Weight (lb) (IM Administration)</th>
</tr>
</thead>
<tbody>
<tr>
<td>3-4</td>
<td>0.1</td>
<td>4-5</td>
</tr>
<tr>
<td>5-7</td>
<td>0.15</td>
<td>6-7</td>
</tr>
<tr>
<td>8-11</td>
<td>0.2</td>
<td>8-12</td>
</tr>
<tr>
<td>12-15</td>
<td>0.25</td>
<td>9-8</td>
</tr>
<tr>
<td>16-21</td>
<td>0.3</td>
<td>10-14</td>
</tr>
<tr>
<td>22-31</td>
<td>0.4</td>
<td>15-20</td>
</tr>
<tr>
<td>32-43</td>
<td>0.5</td>
<td>21-27</td>
</tr>
<tr>
<td>44-55</td>
<td>0.6</td>
<td>28-35</td>
</tr>
<tr>
<td>56-68</td>
<td>0.7</td>
<td>36-44</td>
</tr>
<tr>
<td>69-82</td>
<td>0.8</td>
<td>45-53</td>
</tr>
<tr>
<td>83-97</td>
<td>0.9</td>
<td>54-63</td>
</tr>
<tr>
<td>98-121</td>
<td>1.0</td>
<td>64-78</td>
</tr>
<tr>
<td>122-156</td>
<td>1.2</td>
<td>79-101</td>
</tr>
<tr>
<td>157-194</td>
<td>1.4</td>
<td>102-126</td>
</tr>
<tr>
<td>195+</td>
<td>1.6</td>
<td>127-165</td>
</tr>
</tbody>
</table>

Following injection of Medetomidine Hydrochloride, the dog should be allowed to rest quietly for 15 minutes.

STORAGE
Store at 20° to 25°C (68° to 77°F). Excursions between 15°-30°C (59°-86°F) are allowed. Protect from freezing. Use within 28 days of the first puncture.

BOND
Medetomidine Hydrochloride is supplied in 10-mL, multidose vials containing 1.0 mg of medetomidine hydrochloride per mL. NDC 015914-005-01 - 10 mL vials

TAKE THE Observer DELAY DIRECTIONS
Manufactured for:
Modern Veterinary Therapeutics, LLC
Miami, Florida 33186 - USA