



# Anesthesia and Analgesia

for the Veterinary Practitioner: Canine and Feline







Surgical day pre

The preanesthetic proces:

Drug updates

poting

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for the Veterinary Practitioner: Canine and Feline



Book 4

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# Preface

- Individual state practice act requirements and DEA regulations must be met or exceeded in all instances.
- Review Medical Quality Standards. Meet or exceed all Clinical Essentials.

# State regulations

- At all times, every medical team must comply with individual state practice acts.
- It is each doctor's responsibility to know and understand the requirements of his/her specific state, as well as Banfield policies and procedures.
- The doctor must ensure compliance with state regulations regarding:
  - Handling and administration of controlled substances
  - Intubation of pets
  - Anesthetic monitoring
  - Drug administration documentation
  - Which hospital associates can legally perform dental prophylaxis and all other medical procedures
  - Off-label usage of medications

This publication may contain information that is not within the current FDA-approved labeling for several products for companion animals.

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Dr. Beal received her DVM degree from Oregon State University in 2005. After a short time working with non-human primates, she transitioned to a small animal practice in Salem, Oregon. During her 12 years in practice she accepted the role of Medical Director and helped grow the general practice to eventually include a 24 hour emergency and urgent care facility. Dr. Beal earned her master's in veterinary forensic toxicology in 2021 and currently works at Banfield as a Program Manager on the Veterinary Affairs team promoting medical quality and standards.

### Kathryn Boyle, MS, DVM Senior Manager Veterinary Enterprise Support Banfield Pet Hospital

Dr. Boyle received her Master's degree in Biomedical Science in 2003 and Doctorate of Veterinary Medicine in 2008 from the University of Missouri. In school, she was recognized for her passion for dentistry and clinical pathology. Dr. Boyle worked in a private small animal practice in Springfield, Oregon, after graduation before relocating to St. Louis in 2010, where she joined Banfield as an associate veterinarian. While practicing as a veterinarian, she honed her dentistry skills and developed an interest in promoting the human animal bond through safe patient handling. She was promoted to Chief of Staff in 2012. This role allowed her to drive highquality medical standards and lead multiple teams throughout her market. In 2017, she joined the Veterinary Affairs team, where she supports practice-wide project development and implementation to improve hospital efficiency and effectiveness and allow Banfield associates to provide safe, quality medicine. She has received certifications from Fear Free, Low Stress Handling, Cat Friendly, and Human Animal Bond, which allows her to share her learning and passion for safe, patient-centered care with the rest of the practice.

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# Clinical essentials

**Clinical Essentials** are standards of practice that constitute the minimum acceptable level of care. Practice below this level of care is below expectations. Failure to provide at least this level of care, or clearly document sound reasons for not providing this care, can result in disciplinary consequences.

# General

Veterinarians or trained associates under the direct supervision of a veterinarian perform anesthetic procedures<sup>2</sup>

Sedate or an esthetize brachycephalic pets with brachycephalic-specific protocols and monitoring ^6, ^{10, \, 15, \, 23}

Offer referral of critical or unstable pets when appropriate and in the best interests of the  ${\rm pet}^{1,3}$ 

Current CPR recommendations are reviewed, and CPR training is provided to all associates at least annually. A CPR team is available during normal hours of operation<sup>2, 25</sup>

Do not administer vaccines to an anesthetized patient unless there is a significant pet or associate safety concern to vaccinating a fully conscious  $pet^{33}$ 

Document all perianesthetic physical examination findings, changes in physical status and anesthetic procedure complications in the medical record<sup>7</sup>

Administer all anesthetic medications "to effect" and do not exceed maximum drug dosages<sup>6,7,27</sup>

Utilize multimodal analgesic therapy. Identify and address presurgical, immediate and postoperative pain<sup>7</sup>

Administer IV fluids with every general anesthetic event lasting >10 minutes unless patients are hypervolemic<sup>2</sup>

Place an endotracheal tube with every general anesthetic event and/or procedure in which loss of protective airway reflexes occurs<sup>2</sup>

Assisted ventilation is available for every anesthetic procedure<sup>2, 21</sup>

# Equipment

Utilize the Anesthesia Machine Checklist for every general anesthetic event and have the checklist verified by the attending veterinarian prior to induction<sup>6,17,21</sup>

A crash cart containing emergency drugs and equipment is readily available, in a designated place, portable, clearly labeled and appropriately stocked<sup>2, 25</sup>

Thoroughly clean, disinfect, dry and store personal anesthesia equipment in a manner that prevents contamination prior to each use<sup>7,21</sup>

Anesthetic machines and equipment are tested and maintained on a regular basis and a permanent log of maintenance is kept. Anesthetic events are postponed until all equipment is fully functional<sup>7, 16, 26</sup>

# **Preanesthetic**

Assign and document an ASA status to each pet undergoing general anesthesia and address status appropriately as part of the preanesthetic evaluation. Document discussion of increased risks of anesthesia for pets with an ASA status of ≥III with owner and postpone, cancel or refer anesthetic procedures when indicated<sup>3, 7, 8, 10, 15, 24</sup>

Obtain and review clinical pathology data prior to general anesthesia. Verify, document and address all clinically significant abnormalities prior to premedication, communicate to the team, and discuss all abnormalities with the client. Dismissal of abnormal results is not permitted<sup>6, 7, 18, 24</sup>

Perform a thorough physical exam prior to any anesthetic event, including an accurate weight. Verify, document and address all clinically significant abnormalities prior to premedication, communicate to the team, and discuss with the client. Dismissal of abnormal findings is not permitted<sup>6, 7, 24</sup>

Review the medical history of each pet and pre-emptively identify patientspecific factors that may influence anesthesia (e.g., signalment, adverse drug reactions). Adjust protocols appropriately<sup>21,31</sup> Perform a physical examination (including all cardiovascular parameters) post-premedication and pre-induction for every general anesthetic event<sup>7, 24</sup>

Institute pre-emptive warming measures for those pets below 100°F or for those at greater risk for developing hypothermia (e.g., poor body condition)<sup>21, 31</sup>

Address and resolve physical examination abnormalities that may negatively impact anesthesia (e.g., dehydration, obesity) prior to anesthesia when possible, especially with elective procedures<sup>6</sup>

Provide a minimum of 3 minutes of pre-oxygenation by face mask (flow-by is acceptable if mask is not tolerated), prior to and during the induction process to all pets who will tolerate the procedure<sup>7, 18, 34</sup>

Keep all pets that have been administered preanesthetic medication under visual observation at all times<sup>6</sup>

## Induction & intubation

Coat endotracheal tube cuffs with sterile, water soluble lubricant<sup>7, 22</sup>

Fill endotracheal tube cuffs to the amount required to provide a complete seal and deflate prior to removal (unless otherwise directed by veterinarian)<sup>21</sup>

Keep endotracheal tubes in place until protective, vigorous laryngeal reflexes return without applying noxious stimuli<sup>7</sup>

If patient repositioning is necessary, disconnect intubated pets from the breathing circuit prior to movement and reconnect after attaining proper positioning<sup>7</sup>

# **Monitoring & recovery**

Assign and document at least one hospital associate with the sole responsibility of dedicated, continuous patient monitoring and dedicated recovery to every immobilization or general anesthetic procedure. If there is not a trained, dedicated associate, the procedure must be rescheduled<sup>3, 6, 7, 10, 21</sup>

Continuously measure temperature, heart and respiratory rates, blood pressure, ECG, SpO2, and end-tidal CO2. Document at a minimum of every 5 minutes (or more frequently as clinically indicated) for every general anesthetic event from the time of induction until full recovery<sup>7,21,24</sup> Identify, verify, communicate to the anesthesia team, and address abnormal patient monitoring parameters and trends. Presumptions of malfunctioning equipment and dismissal of abnormal parameters are not permitted. Document findings and treatments on anesthesia form<sup>6,7,21,31</sup>

Abort, as able, elective anesthetic procedures in cases of worsening or refractory patient physical parameters (e.g., hypotension, hypothermia)<sup>6,24</sup>

Keep all patients recovering from an anesthetic procedure under visual observation at all times until full recovery<sup>6, 24</sup>

A final postanesthetic evaluation of each patient is performed by the veterinarian prior to discharge from hospital<sup>6,7</sup>

### IV catheter & multi-dose vial usage

Aseptically place a sterile IV catheter and T-port for every patient receiving IV fluids and for every anesthetic event<sup>12,14,20</sup>

Mark fluid bags with date, time and all additives when initially accessed or when administration sets are attached (fluid bags are spiked), using the appropriate label<sup>13</sup>

Use aseptic technique when placing or accessing patient IV lines, multi-use vials and fluid bags<sup>12, 14, 20</sup>

- Wear new gloves when placing new ports or extensions to IV catheters or fluid bags
- Use a sterile needle and syringe
- Use alcohol to clean ports on IV lines, fluid bags and multi-use vials, allowing to dry before inserting needle
- Discard needle and syringe after use

Change extension sets between each patient undergoing general anesthesia. Use a new, sterile extension set for each patient receiving IV fluids<sup>12, 13</sup>

Discard ALL used fluid bags and administration sets at the end of the day<sup>12, 32</sup>

Discard fluid bags and administration sets upon discontinuation of fluid therapy and replace with new in ANY of the following<sup>11, 14, 20</sup>

- If backflow of blood into any portion of the fluid line is noted
- After fluids have been used on a pet with a known infection
- If any supplemental therapeutics have been injected into the bag or administration line
- If fluid bags and administration sets are used to deliver a fluid which may promote microbial growth

Clamp administration sets closed in between procedures (within day of use window) and place new, sterile needle with cap in place over end of administration set. Hang administration set when not in use so as to not contact patients, tables, or other materials<sup>13, 14, 28</sup>

**Best practices** are standards of practice that meet or exceed an expected level of care and encompass a scale of care from "desirable" to "aspirational."

# General

Designate dedicated anesthetic induction and recovery areas7,21,26

Review anesthetic human safety hazards annually with all hospital associates<sup>2</sup>

Use a new fluid bag and fluid administration set for each pet, regardless of route of fluid administration. Identify each bag with pet name, in addition to date and time<sup>12, 13, 14, 20, 28, 32</sup>

# Equipment

Utilize esophageal instrumentation to provide further means of patient monitoring<sup>7</sup>

# **Preanesthetic**

ldentify, discuss, and address genetic conditions that may impact anesthesia<sup>7</sup>

Utilize the Preanesthetic Timeout Checklist for every general anesthetic procedure<sup>7</sup>

# **Monitoring & recovery**

Train all hospital associates in the appropriate use of pain scales and recognizing pain in pets. Bring concerns of patient analgesia to the attending veterinarian's attention. Review pain recognition annually<sup>5, 16</sup>

Utilize advanced analgesic therapies (soaker catheters, spinal blocks, etc.) appropriately to contribute to pet safety and comfort<sup>16</sup>

Encourage and pursue additional training in advanced anesthetic administration and monitoring for hospital associates<sup>7</sup>

Utilize and follow an anesthetic recovery form with all general anesthetic procedures<sup>3,6</sup>

# Surgical day preparation

A successful anesthesia day begins with the pet owner prior to arrival at the hospital. Clearly communicated recommendations and instructions can lead to a safer and more relaxed visit for our patients.

There are many opportunities to help reduce fear, anxiety and stress in our patients. Options include preparation at home prior to arrival at the hospital and utilization of low-stress techniques during their stay. This begins with educating the client on what they can and should do to make a better experience for their pet. One part of this is ensuring the veterinarian has recommended pre-visit anxiolytics. The client should also have received the medication and have been educated on how to administer the medication appropriately. Utilizing pheromone therapy is also recommended and can be applied in the carrier, in the car or as a collar depending on the patient.

# Home administration of medications

The use of home-administered sedatives/anxiolytics is an often-underutilized opportunity to decrease fear, anxiety and stress for pets, clients, and hospital associates. Many fractious pets are actually displaying a fear response. Decreasing anxiety by providing medication at home is one step to improving their visit. Per AAHA Anesthesia Guidelines, it is strongly recommended that anxiolytics be administered to every fearful, aggressive or stressed patient and should be considered for patients that develop any level of fear, anxiety or stress during their visit. Depending on the procedure being performed, consider the addition of a pre-visit pain medication (opioid), for any pets that may have painful conditions affecting their behavior(arthritis).

# Feline anxiolytics

Drug	Dose	Route
Gabapentin	50–100 mg/cat 150 mg/large cat	2–3 hrs prior to travel
Buprenorphine	0.03 mg/kg oral/transmucosal	60–90 min prior to travel
Buprenorphine + Injectable Acepromazine	0.03 mg/kg oral/transmucosal + 0.05–0.1 mg/kg oral/transmucosal	1.5–2 hrs prior to travel

# Helpful tips:

- Reduce Gabapentin to 50mg/cat in older or petite cats
- For highly anxious cats, begin Gabapentin the night before. Give another dose in early morning
- These medications are not to be used as the pre-anesthetic protocol prior to their anesthetic procedure. Additional medication is needed prior to IV catheter placement and to ensure smooth transition to inhaled anesthesia
- Do not rely on oral gabapentin with patients of unknown health status, acute illness, or respiratory compromise. Unknown health conditions may exacerbate these problems. Instead, rely on lowstress handling techniques and pheromone use

Drug	Dose	Route
Trazodone	5–15 mg/kg PO	1 hour prior to travel
Gabapentin + Injectable Acepromazine	20-40 mg/kg PO + 0.025-0.05 mg/kg oral transmucosal	2–3 hours before travel + 30 min. prior to travel
Gabapentin + Trazadone	20-40 mg/kg PO + 5-15 mg/kg PO	2–3 hours before travel + 1 hour prior to travel

# **Canine anxiolytics**

# **Helpful tips:**

- Paradoxical excitation is uncommon with trazodone, trial doses are still recommended prior to day of visit.
- Gabapentin can be given the night before and repeated in the morning with the acepromazine
- Gabapentin has been noted to not be as effective in medium to large size dogs

# Travel

Pets should be comfortable within their carrier, and it should not be an unfamiliar item utilized only on the day of a veterinary visit. It is best that this is kept within the pet's environment and can be where the pet receives rewards or meals. The carrier should be clean and large enough for the pet to stand up, turn around, and lay down. Consider including an item from home that can also be pre-treated with pheromones.

The hospital team should recognize if a pet arrives showing signs of fear, anxiety, or stress. Signs to look for would be dogs not wanting to walk on their leash or refusing to get on the scale. Cats may be howling or hiding at the back of the carrier. If any of these signs are noted, the anesthesia team should be notified immediately to prevent further escalation.

### Home instructions for chronic medications

For those pets on chronic medications, a discussion should be had regarding when medications should be discontinued prior to anesthesia and when next doses can be given again. Below is a general guideline for some of the more common medications. Each decision is at the veterinarian's discretion.

Medications to continue	Discontinue day of anesthesia	Administer based on recommendation
<ul> <li>Thyroid: supplements or methimazole</li> <li>Behavioral drugs</li> <li>Oral anxiolytics</li> <li>Cardiac medications: pimobendan, furosemide</li> <li>Antibiotics</li> <li>Steroids</li> </ul>	<ul> <li>Antihypertensive medications, especially ACE inhibitors: enalapril, benazepril</li> <li>Anticoagulants: up to 2 weeks prior depending on risk of bleeding</li> </ul>	Insulin: full dose should not be administered to fasted patients

# **Pre-surgical fastings**

The purpose of fasting before surgery is to decrease the risk of regurgitation and vomiting. Fear of complications related to aspiration of gastric contents has led to continued, conservative, preoperative fasting standards. The historic recommendation of "no food overnight" and restricted water does not take into account differences in gastric emptying or the varying times of surgery.

Reflux is influenced by several factors including: the volume and acidity of gastric contents, age, surgical procedure, drugs used for premedication and/or anesthesia and preoperative fasting. Positioning on the surgical table has not shown to influence incidence of regurgitation for those pets without pre-existing conditions.

There is evidence that an increased duration of preoperative fasting is associated with an increased incidence of reflux in dogs. Decreasing the fasting times and providing a small meal, can decrease the reflux episodes.

The following guidelines supported by AAHA, may help decrease regurgitation episodes seen in surgical patients. This is a guideline only and recommendations should be tailored to the needs of each patient and the resources of each hospital.

Patient status	Witl wc (H	nhold ater Irs )	Withhold food (Hrs)				Feed pāté consistency wet food
	0*	6-12	1–2	2-4	4-6	6-12	wettoou
Healthy	<ul> <li>Image: A start of the start of</li></ul>				<ul> <li>Image: A start of the start of</li></ul>		
<8 wks of age or <2 kg	~		No longer than 1-2 hr				In pre-op period
Diabetic	<b>~</b>			~			1/2 meal 2-4 hrs prior
History of or at risk for regurgitation		<ul> <li>Image: A start of the start of</li></ul>				<ul> <li>Image: A start of the start of</li></ul>	Consider feeding 10%-25% of normal amt. 4-6 hr prior to induction
Emergent		ASAP					

\* Ohrs = allow free access to water

# The preanesthetic process

The preanesthetic protocol should be the entire process from the time the patient arrives at the hospital until intubation. There are multiple steps to this process:

- A thorough history, physical exam, ASA, and pain score
- Evaluation of bloodwork, urinalysis, radiographs, BP, ECG, etc.
- Communication with the pet owner to update all normal and abnormal findings, prognosis, and risk
- Preparation of all equipment, fluids, and medications
- Administration of additional premedication/sedation/analgesia
- The beginning of any monitoring as the patient allows
- IV catheter placement
- Preoxygenation for at least 3–5 minutes

# Reviewing bloodwork

Blood analyzers do not take into consideration the guidelines recommended by the International Renal Interest Society (IRIS). To improve patient safety prior to anesthesia, it is recommended to

# IRIS CKD staging guidelines

	Blood creatinine				
Stage	ge Canine		Feli	ne	
	µmol/l	mg/dl	µmol/l	mg/dl	
1	<125	<1.4	<140	<1.6	
2	125-250	1.4-2.8	140-250	1.6-2.8	
3	251-440	2.9-5.0	251-440	2.9-5.0	
4	>440	>5.0	>440	>5.0	

review blood work results compared to any previous values we may have on record. A rising trend in creatinine and/or an elevated creatinine per IRIS guidelines on a fasted patient may indicate underlying early kidney disease and may warrant further investigation prior to anesthesia.

# Premedication and monitoring

# **Route of administration**

IM injections are the preferred route of administration for premedication. They are the least stressful and most reliable route of administration for most patients. Skeletal muscle has a consistent blood supply, and is less likely influenced by factors such as hypotension, hypothermia, and dehydration vs. SC injections.

SC injections are unpredictable in their timing of peak effect. If the patient is well hydrated and there is sufficient time available prior to the desired effects, SC injections are a low-stress option.

IV injections are the fastest route to peak effect, and a predictable way to ensure the patient received the entire dose. Increasing anxiety and stress with heavy restraint to achieve an IV injection is not recommended. Any preinduction struggling and excitation can lead to increased doses of induction drugs, predisposing to hypotension and arrhythmias.

 Allow the appropriate amount of time before induction for the medication used. Peak effect occurring after induction can lead to rapid hypotension and bradycardia. Decisions should be made by the veterinarian, based on patient safety, and individualized for each situation.

# Monitoring and preoxygenation

Monitoring should begin as soon as tolerated.

- BP
- SpO2
- Check temperature
- ECG

Oxygen should be provided as tolerated by the patient. Deliver oxygen by face mask, for 3-5 minutes prior to and during the induction process. If there are delays during induction, return mask to the face and continue oxygen delivery to prevent hypoxemia. Delivery of 100% oxygen for at least 3 minutes provides approximately 6 minutes of complete saturation of oxygen with hemoglobin. This is even more critical in patients with airway disease, breathing difficulty, restricted thoracic movement, and in patients with an expected difficult intubation.

# Notes

# Troubleshooting

The following section provides quick reference lists for some of the most common complications seen during anesthesia. These lists are only a guide. Continued patient monitoring, signalment, medications used and procedures being performed all need to be taken into consideration when troubleshooting complications.

# Acute Bradycardia

Check:	Do:
Pulse – matches ECG?	<ul> <li>Adjust ECG</li> </ul>
Pop-off valve	Open valve – provide assisted
Depth of anesthesia	ventilation if needed
<ul> <li>Body temperature</li> </ul>	Decrease inhalant
ECG – AV block?	Provide warming measures
Recent movement – vagal event?	<ul> <li>Anticholinergics (was dexmedetomidine given?)</li> </ul>
■ BP	<ul> <li>Reverse dexmedetomidine if possible</li> </ul>
Assess for hemorrhage	Treat underlying cause if found
Hyperkalemia? Hypercalcemia?	(stop hemorrhage, administer tx, etc)

# Acute Hypotension (MAP < 60 mm Hg)

Check:	Do:					
BP cuff placement and	Reposition and monitor					
body position	Decrease inhalant and					
Depth of anesthesia	increase O2					
HR and Temperature	Provide analgesia					
Fluid status	Provide warming measures					
Check for blood loss	Anticholinergics:					
	• < 60 large canine					
	• < 80 small canine					
	• < 90 feline					
	<ul> <li>Reversal drugs if applicable</li> </ul>					
	Fluid bolus					
	<ul> <li>Vasopressors if above not working</li> </ul>					

# Hypoxemia (SpO2 <95%)

Check:	Do:
<ul> <li>Assess probe placement</li> </ul>	<ul> <li>Moisten gauze on lingual probe</li> </ul>
<ul> <li>Verify intubation – check for one-lung intubation</li> </ul>	<ul><li>Change tank or abort procedure</li><li>If extubated, provide flow-by, mask</li></ul>
<ul> <li>Verify oxygen source and supply</li> </ul>	or nasopharyngeal O2

# Hypothermia (T < 100°F)

Check:	Do:				
<ul> <li>Warming devices on and working</li> </ul>	<ul> <li>Forced warm air, circulating warming water devices</li> </ul>				
<ul> <li>Depth of anesthesia</li> </ul>	<ul> <li>Warm saline lavage in body cavity</li> </ul>				
Length of surgical prep	<ul> <li>Decrease inhalant if possible</li> </ul>				
Length of anesthesia	<ul> <li>Stage procedure if possible</li> </ul>				
Wet from dental – wet blankets/towels?					
Open body cavity					

# Body positioning

# Risk factors during anesthesia, sedation, and surgery related to body position:

Tracheal damage	<ul> <li>Always disconnect from the breathing circuit prior to any repositioning</li> </ul>
Respiratory difficulty	<ul> <li>Patients positioned with their head below their heart can have difficulty ventilating</li> </ul>
Pain	<ul> <li>Avoid overstretching joints.</li> <li>Provide appropriate padding on table edges, pressure points and areas of trauma</li> </ul>
Compromised airways	<ul> <li>Sternal recumbency is vital during the recovery period to prevent respiratory emergencies</li> </ul>
Vagal events	Acute bradycardia from stimulation of the parasympathetic system. Can be caused by handling the neck during moving

# Quick protocol reference guide

Quick protocol reference guide

# Anxiolytics/sedatives

Dose adjustments	el Start with lower dose in larger dogs	avel Noted to not be as effective in medium avel	rel Reduce to 50 mg in ill or geriatric cats Increase to 150 mg in large cats	el	avel	avel	el	avel	avel
Timing	1hr prior to trav	2-3 hrs prior to tr	1 hr prior to trav	1hr prior to trav	2-3hrs prior to tr	2-3 hrs prior to tr	1hr prior to trav	1-2 hrs prior to tr	1-2 hrs prior to tr
Healthy dose	5–15 mg/kg PO	20-40 mg/kg	50-100 mg/cat PO	5–15 mg/kg PO	20-40 mg/kg PO	20-40 mg/kg PO +	0.025–0.05 mg/kg transmucosal	0.03 mg/kg transmucosal	0.03 mg/kg transmucosal + 0.05-0.1 mg/kg transmucosal
Species	Canine	Canine	Feline				Canine	Feline	Feline
Drug	Trazadone		Gabapentin	Trazadone	Gabapentin	Gabapentin	+ Acepromazine [Inj]	Buprenorphine	Buprenorphine + Acepromazine [Inj]

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Drug option	Species	Level of sedation	Health status	Examples
Opioid	Feline	Light	Healthy	Butorphanol 0.2–0.4 mg/kg IM/IV
Opioid	Canine	Light-Moderate	Healthy & Compromised	Butorphanol 0.2–0.4 mg/kg IM/IV Midazolam 0.2 mg/kg IM/IV
Benzodiazepine	Feline	Light-Moderate	Compromised	Butorphanol 0.2–0.4 mg/kg IM/IV Midazolam 0.2 mg/kg IM/IV
	Canine	Moderate	Healthy	Butorphanol 0.4 mg/kg IM Acepromazine 0.01–0.03 mg/kg IM
Opioid Tranquilizer	Feline	Moderate	Healthy	Butorphanol 0.4 mg/kg IM OR Buprenorphine 0.02 mg/kg OTM Acepromazine 0.025-0.1 mg/kg IM
	Canine	Moderate	Healthy	Butorphanol 0.4 mg/kg IM Dexmedetomidine 3-7 <b>mcg</b> /kg IM
Opioid Alpha-2 Agonist	Feline	Moderate	Healthy	Butorphanol 0.4 mg/kg IM OR Buprenorphine 0.02 mg/kg OTM (cat Dexmedetomidine 3-10 <b>mcg</b> /kg IM

Examples	Butorphanol 0.4 mg/kg IM Acepromazine 0.01–0.03 mg/kg IM Dexmedetomidine 3–7 <b>mcg</b> /kg IM	Butorphanol 0.4 mg/kg IM OR Buprenorphine 0.02 mg/kg OTM Acepromazine 0.01-0.03 mg/kg IM Dexmedetomidine 3-10 <b>mcg</b> /kg IM	Dexmedetomidine 7–15 <b>mcg</b> /kg IM Alfaxalone 1–2 mg/kg IM	Butorphanol 0.2-0.4 mg/kg IM	Alfaxalone 1-2 mg/kg IM		DKT 1:1:1
Health status	Healthy	Healthy	Healthy	Compromised	Compromised	Healthy	Healthy
Level of sedation	Moderate	Moderate Heavy Heavy			:	Heavy	
Species	Canine	Feline	Canine	Canine	Feline	Canine	Feline
Drug option		Opiola Tranquilizer Alpha-2 Agonist	Alpha-2 Agonist Alfaxalone	Opioid	Benzoalazepine Alfaxalone	Alpha-2 Agonist	Dissociative Opioid

Premedication drug combinations

# Helpful hints:

- Patient response to handling in the hospital is ultimate determination of dose
- Consider any prior treatment with anxiolytics prior to dosing

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Duration		6–8 hours Slowe	24 hours	96 hours	1–2 hours	[mostly sedation]	1-2 hours	it no CRI
Dose	0.01-0.03 mg/kg IM, IV	0.01-0.02 mg/kg IM, IV 0.02-0.03 mg/kg PO	0.18 mg/kg SC	Size dependent tube			Loading dose: 2–10 <b>mcg</b> /kg	CRI: 2-10 <b>mcg</b> /kg/h
Species	Canine	Feline	Feline	Feline	Canine	Feline	Canine	Feline
ס	Buprenorphine		orphine Icting)	orphine lermal)		2	Inu	D

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ig Species	Canine	rphone Feline		Canine	done Feline	Canine	line	Feline	sal drugs		d Dpioids (partial b			
Dose	0.05-0.2 mg/kg IM, IV, SC	0.02-0.05 mg/kg IM, IV, SC		0.02-0.05 mg/kg IM, IV, SC			0.1–0.5 mg/kg IM, IV, SC		0.5-1.0 mg/kg IM, SC			Reverses	utorphanol and buprenor	
Duration		4 hours			6–12 hours		4-6 hours				phine)			
Notes	Panting, vomiting, diarrhea, bradycardia	Consider pretreatment with maropitant	Less vomiting with IV use	Reduced nausea vs other opioids Cost limits use to cats and	small dogs Reduced inhalant need, monitor for respiratory depression	Sedation, vomiting, diarrhea, bradycardia Hypotension and	bronchoconstriction [histamine release [V]	Consider pretreatment with maropitant		Dose	0.01-0.04 mg/kg IV			

equal volume IM

Alpha-2 agonist

Atipamezole

Notes         Reversible         Very low doses potentiate opioids         Agitated and prolonged recovery         (dose dependent)         Avoid in severe heart, renal or         liver disease         Avoid presurgical, unless chronic use         Avoid with Gl disease, renal disease,         bleeding, or dehydration         Avoid with Gl disease, renal disease,         Avoid presurgical, unless chronic use         Avoid with Gl disease, renal disease,         Avoid with Gl disease, renal disease,         Avoid with Gl disease, renal disease,         bleeding, or dehydration         Avoid with Gl disease, renal disease,         bleeding, or dehydration         Avoid with Gl disease, renal disease,         bleeding, or dehydration         Rooid with Gl disease, renal disease,         bleeding, or dehydration         Rooid with Gl disease, renal disease,         bleeding, or dehydration         HICH RISK	Duration Variable I+-6 hours 24 hours 24 hours	Dose       2-10 mcg/kg IM, IV combined with opioid       2.0.02-1.0 mg/kg IM, SC, PO       0.02-1.0 mg/kg IM, SC, PO       0.02-1.0 mg/kg IM, with opioid       4 mg/kg SC day 1, then 0.1 mg/kg PO       0.3 mg/kg SC day 1, then 0.1 mg/kg PO       0.1 mg/kg SC day 1, then 0.05 mg/kg PO	Species Species Canine Feline Canine Canine Feline Feline	<b>brug</b> Dexmedetomidine Ketamine Carprofen Meloxicam
Wide safety margin (overdosing less dangerous) vs other options	24 hours	2 mg/kg SC	Feline	Robenacoxib
Avoid with GI disease, renal of bleeding, or dehydratic HIGH RISK		OR 0.1 mg/kg SC day 1, then 0.05mg/kg PO	Feline	
Avoid presurgical Avoid with GI disease, renal diseas	24 hours	0.3 mg/kg SC once OR		Meloxicam
Avoid presurgical, unless chronic u Avoid with GI disease, renal diseas bleeding, or dehydration		0.2 mg/kg SC day 1, then 0.1mg/kg PO	Canine	
Avoid presurgical, unless chronic us Avoid with GI disease, renal diseas bleeding, or dehydration	24 hours	H mg/kg SC	Canine	Carprofen
Avoid in severe heart, renal or liver disease		Combined with opioid	Feline	
Very low doses potentiate opioids Agitated and prolonged recovery [dose denendent]		0.02-1.0 mg/kg IM, SC, PO	Canine	2
Avold in Udesarean sections, hear disease, acute illness, or severe inju	variable	iv combined with opioid	Feline	Jexmedetomidine
Reversible		2-10 mcg/kg IM,	Canine	
Notes	Duration	Dose	Species	Drug
			6	ther analgesic:

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algesics	Species Dose	<b>Canine</b> 10 mg/kg PO 8-	Canine 1-2 mg/kg PO Feline	Canine 2-4 mg/kg	1-2 mg/kg/d 3-4 mg/kg/d	Canine 5 mg/kg
	Duration	8-12 hours x 7 days	8 hours	12–24 hours (dose dependent)	24 hours 24 hours x 7 days max	24 hours
	Notes	Toxic to cats Used for mild to moderate pain Do not use in patients with liver disease	Moderate to severe pain and cough suppressant Bad taste, can cause dysphoria Minimal toxicity	Potential for Glulceration, renal failure and idiosyncratic hepatic dysfunction Avoid in bleeding or dehydrated patients	Avoid with GI disease, renal disease, bleeding, or dehydration	Avoid with GI disease, renal disease, bleeding, or dehydration

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Drug	Species	Dose	Duration	Notes
	Canine	0.1 mg/kg PO	24 hours	Avoid with GI disease, renal disease,
Meloxicam	Feline	0.05 mg/kg PO	24 hours x 3 days max	bleeding, or dehydration HIGH RISK in cats
Robenacoxib	Feline	2.5-6 kg: 6 mg tablet PO >6 kg: 2 × 6mg tablet PO	24 hours x 3 days max	Palatable for most cats Avoid in cats with renal or liver disease or coagulopathy Best absorption when given WITHOUT food
	Canine	2–8 mg/kg PO	4-8 hours	Controversial - lack of data for efficacy
Tramadol	Feline	2-4 mg/kg PO	12 hours	in dogs for acute pain Best if used as adjunct to NSAIDs (chronic pain)

# Drug updates

# Alfaxalone

Alfaxalone is a neurosteroid (alters neuronal excitability) derived from progesterone and used as an induction and/or maintenance agent to induce general anesthesia. Similar to propofol, alfaxalone has a rapid onset, lack of accumulation with short and effective duration while also having dose dependent cardiovascular and respiratory depression. It is a class IV-controlled substance with an FDA label for IV administration in the US. There are multiple formulations and use of the multi-dose vial is recommended due to the 28-day shelf-life.

Multiple studies have been performed showing efficacy and safety when given IM. If used IM, do not switch to propofol for induction; continue to use alfaxalone as the induction drug.

### Tips for IV induction with alfaxalone:

- Alfaxalone is similar to propofol and can have similar side effects when given rapidly
- Administer 0.5 mg/kg IV every 30 seconds until the patient relaxes and becomes less responsive to stimuli
- Provide O2 supplementation by face mask as soon as tolerated
- Most patients will need between 1-2mg/kg total dose depending on differences in premedication drugs/doses, health status, and age
- If the patient still has a pronounced cough/swallow reflex at intubation, continue with an additional alfaxalone 0.5 mg/kg to help smooth the transition to inhalant.
- Additional incremental 0.5 mg/kg IV doses of alfaxalone may be administered, if required during the transition period, to inhalant anesthesia.

### Tips for IM sedation with alfaxalone:\*

- Should be combined with an opioid +/- benzodiazepine
- Short procedures are more likely to have rough recoveries (paddling, myoclonus, rolling, flailing), more pronounced when used alone
- Large volume needed for IM sedation will limit use to cats and small dogs
- Should not be painful on injection
- Ideal for stressed/fearful cats with unknown health status

# Maropitant

The anesthesia outcomes that occur most frequently in human anesthesia are pain, nausea, and vomiting. Nausea may not result in vomiting and this clinical sign may be overlooked in our patients. Maropitant (Cerenia) is a selective neurokinin-1 receptor antagonist which blocks the binding of the neurotransmitter Substance P (SP). SP is found in high concentrations in both the chemoreceptor trigger zone and the vomiting center and is the key neurotransmitter involved in vomiting. Cerenia is most commonly utilized in the treatment of motion sickness, nausea and vomiting. Studies show that there is some benefit beyond these common ailments and the application of Cerenia in anesthetic protocols is becoming more common.

### Anesthesia and Surgical applications of maropitant:

- Administration of maropitant prior to routine spays, decreased the amount of inhalant anesthetic needed during visceral stimulation (pain) including stimulation of the ovaries and ovarian ligaments
- Maropitant administration prevents nausea and vomiting commonly seen after use of hydromorphone and morphine
- Maropitant treated dogs are significantly more likely to eat within 3 hours of extubating compared to dogs who received opioids alone
- Smoother recovery from anesthesia

### High risk patients for vomiting and aspiration pneumonia:

- Opioid administration (Hydromorphone, etc.)
- Brachycephalic, especially bulldog breeds
- Gl disease
- Increase intraocular/intracranial pressure

\*extra-label use in United States

History of motion sickness, vomiting, regurgitation

### **Maropitant Use in Cats**

Subcutaneous administration of maropitant in cats is effective in decreasing, but not eliminating, vomiting and signs of nausea prior to morphine or dexmedetomidine premedication. Feline patients exhibit significant aversive behaviors and pain upon SC injection, and therefore oral administration at least 2–3 hours prior to opioid premedication may be a more humane option. Oral maropitant significantly decreases but does not eliminate vomiting and signs of nausea associated with morphine or dexmedetomidine premedication in cats.

# Buprenorphine - Transdermal (Zorbium)

Zorbium is a Schedule III opioid, indicated for the control of post-operative pain associated with surgical procedures in cats. It is a one-time administration, applied in the veterinary hospital, providing analgesia for 4 days. . Zorbium can be applied 1-2 hours prior to surgery as a premedication for surgical pain. Transdermal buprenorphine has similar side effects to other buprenorphine products; monitor for dysphoria, hyperthermia, and constipation.

### **Tips for application:**

- Wear latex or nitrile gloves, protective glasses, and a lab coat to prevent skin, eye, or mucosa contact
- Twist to open the applicator tube, the top will not come off
- Do not clip the fur and do not apply to injured/diseased skin
- Part the fur and apply the tube directly to the skin at the base of the head/neck ONLY, emptying the entire contents
- Allow 30 minutes of dry time
- DO NOT send home with clients to apply
- Should only be administered by veterinarians or certified technicians who have been trained in handling opioids

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