

Top 5 Tips for Sedation & Anesthesia in Fractious Dogs

Katherine Bennett, DVM
Christine Egger, DVM, MVSc, CVA, CVH, DACVAA
University of Tennessee



▲ **FIGURE 1** A long extension set directly connected to the catheter, which is placed in the lateral saphenous vein. An injection port is accessible (out of frame).

Aggression represents over 50% of behavior-related problems in dogs,¹ and fractious animals pose an inherent risk to veterinary staff. Behavior management is the ideal long-term solution for aggressive or fractious animals; however, some surgical or diagnostic procedures require relatively immediate attention and preclude most recommended behavior modifications. Precautions should be taken to ensure both patient and team safety when sedating or anesthetizing these patients.

Following are the authors' tips for safe handling of a sedated or anesthetized fractious dog presented for diagnostic or surgical procedures.

1 Owner Communication
Communication with the pet owner ahead of the scheduled appointment is critical. Discussion should include current medications, patient behavior at home, and whether the owner is

comfortable medicating the patient at home. Owner involvement can help facilitate a team-based approach to safe and effective patient sedation.² In addition, a thorough risk assessment should be explained to the owner, as many sedative medications can have adverse effects on patients with underlying diseases, particularly cardiovascular disease. Patients with underlying systemic disease may require dose alterations and/or alternative drug protocols to account for comorbidities.

TOP 5 TIPS FOR SEDATION & ANESTHESIA IN FRACTIOUS DOGS

1. Owner Communication
2. Preappointment Preparation
3. Sedation Administration
4. Patient Handling While Hospitalized
5. Recovery & Discharge

2 Preappointment Preparation

At-home administration of one or more sedatives (eg, trazodone, clonidine, dexmedetomidine, acepromazine, alprazolam; **Table 1**) the day before and the day of the scheduled visit allows for multimodal anxiolysis and can facilitate delivery of additional sedatives in the clinical setting. Caution should be taken when prescribing multiple serotonin-altering medications, as serotonin syndrome is a potentially lethal side effect (see **Serotonin Syndrome**, next page).³ Combining different medications or introducing new serotonin-altering medications to a dog's treatment protocol can have deleterious effects; owners should be informed that, although uncommon, disinhibition of behavioral tendencies⁴ and/or development of aggression⁵ can occur at home. If adverse behavioral effects or signs of serotonin syndrome do not occur, the dose can be gradually increased over 1 to 2 days until the desired dose is reached or the desired effect is achieved.⁶ Alternatively, medications that do not alter serotonin levels (eg, α_2 agonists, benzodiazepines, gabapentin) can be used.

Common at-home administration protocols include administering oral trazodone, gabapentin, and alprazolam the day before the appointment and on the morning of the scheduled appointment or administering oral acepromazine, gabapentin, and alprazolam, potentially coadministered with maropitant (2 mg/kg PO q24h) to decrease the risk for vomiting after later administration of injectable sedatives (especially those that contain a pure μ opioid).⁷

3 Sedation Administration

Many patients may become more stressed in the hospital waiting area, making it more difficult for sedative medications to reach full efficacy. The owner should be advised to place a muzzle and/or Elizabethan collar on the patient before or just after arrival, if possible. If available, other parts of the hospital (eg, parking lot, grassy relief area, barn) can be used as an environmental distraction for the patient during handling, waiting, and/or sedative administra-

tion.⁸ Because dogs use multiple cues (eg, visual, auditory, olfactory) to influence their behavior and/or reactions to their environment,⁹ soft and calm voices and limited personnel involvement are recommended. Pheromone sprays can help reduce anxiety but have not been shown to consistently reduce aggression in dogs.¹⁰

White coat syndrome (ie, the increase in a patient's sympathetic response to stress due to the appearance of medical personnel in white coats or similar clothing) has been well documented in human medicine.¹¹⁻¹³ To reduce the perceived threat of medical personnel, staff members who interact with the patient should avoid wearing white coats or similar hospital clothing while initially handling the patient (ie, from arrival to administration of injectable sedation). Typically, a coat or other outerwear is recommended to be worn over hospital clothing.^{11,12}

TABLE 1

PERIOPERATIVE ANXIOLYTIC & SEDATIVE DOSAGES IN DOGS¹⁵⁻¹⁷

Drug (Drug Category)	Dosage*
Acepromazine (phenothiazine)	0.5-2 mg/kg PO q8h
Alprazolam (benzodiazepine)	0.02-0.04 mg/kg PO q6h
Clonidine (α_2 agonist)	0.01-0.05 mg/kg PO q12h
Dexmedetomidine gel (α_2 agonist)	Refer to product insert
Diazepam (benzodiazepine)	1-2 mg/kg PO q8h
Gabapentin (anticonvulsant, neuropathic pain analgesic)	5-10 mg/kg PO q8-12h
Trazodone** (serotonin antagonist and reuptake inhibitor)	2-10 mg/kg PO q8-12h

*Some dosages are anecdotal based on those used in the authors' facility.

**Indicates commonly prescribed medications that, when combined with other serotonin-altering drugs, may place the patient at risk for serotonin syndrome. Careful and controlled introduction of medication combinations can help mitigate risks for serotonin syndrome development.

SEROTONIN SYNDROME

Serotonin syndrome, defined as a group of clinical signs associated with administration of serotonin-altering medications (eg, selective serotonin reuptake inhibitors, monoamine oxidase inhibitors, antidepressants), although rare in veterinary medicine, can occur when multiple serotonin-altering medications are coadministered.¹⁸ Clinical signs of serotonin syndrome include altered mental status, agitation, nervousness, myoclonus, hyperreflexia, tremors, diarrhea, incoordination, increased heart rate and blood pressure, and hyperthermia.¹⁹ If a patient is already receiving medications for behavior alteration or other reasons, slow introduction of additional medications at lower doses is recommended. Any signs of agitation, restlessness, or myoclonus may suggest serotonin syndrome, and cessation of any additional serotonin-altering medications is recommended.

TABLE 2

SEDATIVE DOSAGES IN DOGS¹⁵⁻¹⁷

Drug	Dosage*	Duration of Full Effect**
Acepromazine	0.01-0.03 mg/kg IM	6-8 hours
Alfaxalone	1-3 mg/kg IM	15-20 minutes
Butorphanol	0.1-0.4 mg/kg IM	30-60 minutes
Dexmedetomidine	1-10 µg/kg IM (not to exceed 10 µg/kg)	30-60 minutes
Hydromorphone	0.05-0.1 mg/kg IM	4-6 hours
Ketamine	3-10 mg/kg IM	30-60 minutes
Midazolam	0.1-0.5 mg/kg IM	20-40 minutes
Tiletamine/zolazepam	1-4 mg/kg IM	30-60 minutes

*Some dosages are anecdotal based on those used in the authors' facility.

**Most drugs have a dose-dependent duration of effect (ie, higher doses usually prolong the effect); however, higher doses can also increase the frequency of adverse events.

Administering sedation via an intramuscular injection (*Table 2*) is preferable and can be done while the patient is walking on a leash, provided the person handling the patient and the person administering the drugs are both experienced enough for a rapid pelvic limb injection and subsequent patient reaction. These drugs are typically used in combination to provide deep sedation and/or general anesthesia. Combining different drug classes (*Table 3*) allows for a dose reduction in all agents, thereby potentially limiting negative adverse effects.

Other handling techniques involve using a half-wall or chain link fence as a barrier between the patient and the injector/handler. An ideal sedative protocol, as recommended in human medicine, is rapid-acting with minimal side effects, although, without physical examination, adverse effects are difficult to predict in fractious patients.¹³ Of note, most anesthetic drugs are associated with some degree of risk¹⁴; this risk is increased in patients that are unable to be assessed for pre-existing comorbidities (eg, heart disease). Reversible drugs (eg, α_2 agonists, opioids) are preferable, as their adverse effects can be mitigated with reversal agents if necessary.

Some patients may become sedate enough to lose airway protection. Supplies for intubation and appropriate ventilation should always be available for patients that show signs of requiring a protected airway or ventilatory support (eg, cyanosis, shallow breathing, regurgitation).

4 Patient Handling While Hospitalized

Fractious patients may require additional precautions for handling and drug administration while hospitalized. Standard monitoring procedures are recommended with the patient sedated or anesthetized. Hospitalization of fractious animals typically requires planning.

Placement of an IV catheter in a pelvic limb can be advantageous, as it provides more room between the patient's head and the injection site. If pelvic limb catheter placement is not feasible, additional

placement of long extension sets attached to the IV catheter (**Figure 1**, page 36) can facilitate semi-remote drug administration and provides an additional level of safety for the patient and staff.

An Elizabethan collar and/or basket muzzle can be used to provide additional safety for aggressive patients, and allowing patients to wear a harness with an attached leash while in a cage can be helpful when removing them from the confined space (**Figure 2**). Floor-level cages or runs are preferred, as they prevent the need for the handler to lift the patient out of the cage and onto the floor or into a carrier. Muzzles with connections suitable for oxygen delivery are also helpful for providing flow-by oxygen to aggressive patients.



▲ **FIGURE 2** To ensure patient and staff safety, an Elizabethan collar and a harness are used on the patient, with the leash attached to the harness and placed toward the cage door.

5 Recovery & Discharge

For outpatient procedures (eg, outpatient surgery, diagnostic procedures) requiring sedatives/anesthetic drugs, a basket muzzle

TABLE 3

SEDATIVE COMBINATIONS & DOSAGE RECOMMENDATIONS IN DOGS¹⁵⁻¹⁷

Drug Combination*	Dosage**	Effect
Combination 1		
Butorphanol	0.4 mg/kg IM	High level of sedation with mild analgesia
Dexmedetomidine	5 µg/kg IM	
Tiletamine/zolazepam	3 mg/kg IM	
Combination 2		
Hydromorphone†	0.1 mg/kg IM	Higher degree of analgesia with good sedation
Dexmedetomidine	5 µg/kg IM	
Ketamine	2 mg/kg IM	
Combination 3		
Butorphanol	0.4 mg/kg IM	Dissociative anesthetics or α ₂ agonists are not recommended in patients with questionable cardiac disease or significant comorbidities
Alfaxalone	2 mg/kg IM	
Midazolam	0.5 mg/kg IM	

*Opioids can be substituted within their drug class (eg, butorphanol substituted for hydromorphone) if goals for pain management require a different opioid.

Doses can be adjusted based on recommended dosing ranges (Table 2**). Some dosages are anecdotal based on those used in the authors' facility.

†Any opioid can be substituted for hydromorphone based on availability.

STEP-BY-STEP CATHETER REMOVAL VIDEO

To view a video showing step-by-step removal of an intravenous catheter at the end of an anesthetic procedure in a fractious animal, scan the QR code below or view this article online at cliniciansbrief.com/sedation-tips



Using QR codes from your mobile device is easy and quick!

Simply focus your phone's camera on the QR code as if taking a picture (but don't click!). A notification banner will pop up at the top of your screen; tap the banner to view the linked content.

can be modified so that the endotracheal tube can be removed through the muzzle, which allows the muzzle to be placed on the patient prior to extubation and be in place at the end of the procedure (**Figure 3**). This facilitates safety in the recovery period while still allowing the patient to be closely monitored.

Intravenous catheters can be removed just before discharge. With all tape removed and a bandage left over the catheter, the extension line, which is attached to the catheter hub, can be pulled, thus removing the catheter while keeping the bandage in place for hemostasis (see **Step-by-Step Catheter Removal Video**). Sedatives can be administered intravenously just before catheter removal at the time of discharge and can facilitate a smooth transition from the hospital to the transportation vehicle. The owner should be made aware of the expected nature and duration of the sedation protocol.



▲ **FIGURE 3** Basket muzzle modified to facilitate extubation (A). Placement of the pilot balloon and endotracheal tube ties through the end of the muzzle is necessary to avoid difficulty extubating the patient (B).

Conclusion

Careful planning, communication, and preparation can facilitate a safe and productive appointment for fractious patients that need to be sedated or anesthetized. Multi-

modal pharmacologic restraint, along with modified approaches to drug administration and patient handling, can mitigate most of the issues encountered with aggressive patients in the hospital setting. ■

References

1. Fatjó J, Amat M, Mariotti VM, Luis Ruiz de la Torre J, Manteca X. Analysis of 1040 cases of canine aggression in a referral practice in Spain. *J Vet Behav Clin App Res.* 2007;2(5):158-165.
2. Sueda KL, Malamed R. Canine aggression toward people: a guide for practitioners. *Vet Clin North Am Small Anim Pract.* 2014;44(3):599-628.
3. Boyer EW, Shannon M. The serotonin syndrome. *N Engl J Med.* 2005; 352(11):1112-1120.
4. Gruen ME, Sherman BL. Use of trazodone as an adjunctive agent in the treatment of canine anxiety disorders: 56 cases (1995-2007). *J Am Vet Med Assoc.* 2008;233(12):1902-1907.
5. Gilbert-Gregory SE, Stull JW, Rice MR, Herron ME. Effects of trazodone on behavioral signs of stress in hospitalized dogs. *J Am Vet Med Assoc.* 2016;249(11):1281-1291.
6. Thomas DE, Lee JA, Hovda LR. Retrospective evaluation of toxicosis from selective serotonin reuptake inhibitor antidepressants: 313 dogs (2005-2010). *J Vet Emerg Crit Care (San Antonio).* 2012;22(6):674-681.
7. Hay Kraus BL. Efficacy of maropitant in preventing vomiting in dogs premedicated with hydromorphone. *Vet Anaesth Analg.* 2013;40(1):28-34.
8. Hsu Y, Sun L. Factors associated with aggressive responses in pet dogs. *Appl Anim Behav Sci.* 2010;123(3-4):108-123.
9. Luescher AU, Reisner IR. Canine aggression toward familiar people: a new look at an old problem. *Vet Clin North Am Small Anim Pract.* 2008;38(5):1107-1130, vii.
10. Mills DS, Ramos D, Estelles MG, Hargrave C. A triple blind placebo-controlled investigation into the assessment of the effect of Dog Appeasing Pheromone (DAP) on anxiety related behaviour of problem dogs in the veterinary clinic. *Appl Anim Behav Sci.* 2006;98(1):114-126.
11. Pickering TG, James GD, Boddie C, Harshfield GA, Blank S, Laragh JH. How common is white coat hypertension? *JAMA.* 1988;259(2):225-228.
12. Wang XX, Shuai W, Peng Q, et al. White coat effect in hypertensive patients: the role of hospital environment or physician presence. *J Am Soc Hypertens.* 2017;11(8):498-502.
13. Moore G, Pfaff JA. Assessment and emergency management of the acutely agitated or violent adult. UpToDate. <https://www.uptodate.com/contents/assessment-and-emergency-management-of-the-acutely-agitated-or-violent-adult>. Updated October 2, 2017. Accessed September 12, 2018.
14. Brodbelt DC, Flaherty D, Pettifer GR. Anesthetic risk and informed consent. In: Grimm KA, Lamont LA, Tranquilli WJ, Greene SA, Robertson SA, eds. *Veterinary Anesthesia and Analgesia*. 5th ed. Ames, IA: John Wiley & Sons; 2015:11-22.
15. Crowell-Davis SL, Seibert LM, Sung W, Parthasarathy V, Curtis TM. Use of clomipramine, alprazolam, and behavior modification for treatment of storm phobia in dogs. *J Am Vet Med Assoc.* 2003;222(6):744-748.
16. Grimm KA, Lamont LA, Tranquilli WJ, Greene SA, Robertson SA, eds. *Veterinary Anesthesia and Analgesia: The Fifth Edition of Lumb and Jones*. Ames, IA: John Wiley & Sons; 2015.
17. Plumb DC. *Plumb's Veterinary Drug Handbook*. 9th ed. Hoboken, NJ: Wiley-Blackwell; 2018.
18. Haberzettl R, Bert B, Fink H, Fox MA. Animal models of the serotonin syndrome: a systematic review. *Behav Brain Res.* 2013;256:328-345.
19. Crowell-Davis SL, Poggiagliolmi S. Understanding behavior—serotonin syndrome. *Compend Contin Educ Vet.* 2008;30(9):490-493.

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References

1. Gommeren K, van Hoek I, Lefebvre HP, Benckroun G, Smets P, Daminet S. Effect of thyroxine supplementation on glomerular filtration rate in hypothyroid dogs. *J Vet Intern Med.* 2009;23(4):844-849.
2. Panciera DL, Lefebvre HP. Effect of experimental hypothyroidism on glomerular filtration rate and plasma creatinine concentration in dogs. *J Vet Intern Med.* 2009;23(5):1045-1050.
3. Williams TL, Elliott J, Syme HM. Association of iatrogenic hypothyroidism with azotemia and reduced survival time in cats treated for hyperthyroidism. *J Vet Intern Med.* 2010;24(5):1086-1092.
4. Peterson ME, Nichols R, Rishniw M. Serum thyroxine and thyroid stimulating hormone concentration in hyperthyroid cats that develop azotemia after radioiodine therapy. *J Small Anim Pract.* 2017;58(9):519-530.
5. Nykamp SG, Dykes NL, Zarfoss MK, Scarlett JM. Association of the risk of development of hypothyroidism after iodine 131 treatment with the pretreatment pattern of sodium pertechnetate Tc 99m uptake in the thyroid gland in cats with hyperthyroidism: 165 cases (1990-2002). *J Am Vet Med Assoc.* 2005;226(10):1671-1675.
6. Mooney CT. Radioactive iodine therapy for feline hyperthyroidism: efficacy of administration routes. *J Small Anim Pract.* 1994;35(6):289-294.
7. Meric S, Rubin S. Serum thyroxine concentrations following fixed-dose radioactive iodine treatment in hyperthyroid cats: 62 cases (1986-1989). *J Am Vet Med Assoc.* 1990;197(5):621-623.
8. Chun R, Garrett LD, Sargent J, Sherman A, Hoskinson JJ. Predictors of response to radioiodine therapy in hyperthyroid cats. *Vet Radiol Ultrasound.* 2002;43(6):587-591.
9. Lucy JM, Peterson ME, Randolph JF, et al. Efficacy of low-dose (2 millicurie) versus standard-dose (4 millicurie) radioiodine treatment for cats with mild-to-moderate hyperthyroidism. *J Vet Intern Med.* 2017;31(2):326-334.
10. Boag AK, Neiger R, Slater L, Stevens KB, Haller M, Church DB. Changes in the glomerular filtration rate of 27 cats with hyperthyroidism after treatment with radioactive iodine. *Vet Rec.* 2007;161(21):711-715.
11. Peterson M, Melian C, Nichols R. Measurement of serum concentrations of free thyroxine, total thyroxine, and total triiodothyronine in cats with hyperthyroidism and cats with nonthyroidal disease. *J Am Vet Med Assoc.* 2001;218(4):529-536.
12. Peterson M, Gamble D. Effect of nonthyroidal illness on serum thyroxine concentrations in cats: 494 cases (1988). *J Am Vet Med Assoc.* 1990;179(9): 1203-1208.
13. Mooney C, Little C, Macrae A. Effect of illness not associated with the thyroid gland on serum total and free thyroxine concentrations in cats. *J Am Vet Med Assoc.* 1996;208(12):2004-2008.
14. Wakeling J, Moore K, Elliott J, Syme H. Diagnosis of hyperthyroidism in cats with mild chronic kidney disease. *J Small Anim Pract.* 2008;49(6):287-294.
15. Williams TL, Elliott J, Syme HM. Effect on renal function of restoration of euthyroidism in hyperthyroid cats with iatrogenic hypothyroidism. *J Vet Intern Med.* 2014;28(4):1251-1255.