Opioids Overview and General Clinical Effects

Analgesia: MOR are potent analgesics

Behavioral effects: Sedation, euphoria, dysphoria *Species and dose dependent

Sedation: dogs, rabbits, mild in ruminants

Euphoric: cats and horses

Dysphoria: possible in any species *Seem to be less if the animal is in pain when the opioid is administered

Anesthetic sparing effects

Emesis: Tend to promote emesis at the chemoreceptor trigger zone (CRTZ) but inhibit emesis at the vomiting center

Cardiovascular effects: Minimal as long as HR is maintained, can produce vagally mediated reduction in HR but this is treatable with an anticholinergic. Cats and horses may have an increased HR

Histamine release: may decrease systemic vascular resistance *meperidine should not be given IV, morphine can be given slow IV **Respiratory system**: respiratory depression, high doses of mu agonists can produce apnea, panting is common in dogs due to its effect on thermoregulation (they feel warmer than they actually are)

Thermoregulation: Hypothermia in dogs, hyperthermia in cats

Pupil size: Miosis (constriction) in dogs and rabbits, mydriasis (dilation) in cats and rodents

Urinary: MOP agonists can produce urinary retention by increasing urethral sphincter tone, relaxation of bladder smooth muscle and

increased ADH secretion

GIT effects: Delayed gastric emptying, relaxation of lower esophageal sphincter, altered motility

Opioid Drugs	Receptor	Clinical Effects	Duration	Administration	Additional Info
Morphine	Mu agonists	Analgesia	4 hours	Morphine must be	Vomiting is common in
Hydromorphone		Euphoria, dysphoria,		given slowly IV, can	dogs
Oxymorphone		sedation	*Fentanyl has a	cause histamine	
Fentanyl		Respiratory depression	30-60 min duration	release	Hydromorphone may
		Change in pupil size			produce less sedation in
		Reduced GI motility			dogs and hyperthermia
		Urine retention			in cats
		Change in HR			
		Hypothermia			

					Oxymorphone is less likely to produce dysphoria in cats
Agonist/Antagonist -Butorphanol	Kappa agonist Mu receptor antagonist	Analgesia mediated via kappa receptors but has a ceiling effect due to MOR antagonism	Rapid onset after IV administration and has a duration of 30-60 min Up to 90 min after IM administration	IV and IM	Commonly used in large animals
Partial Agonist -Buprenorphine	Mu receptor agonist	Produces all the effects of MOR agonists but to a lesser degree	6 hours	IV – 15 min until action IM – 20-30 min until action	
	Kappa	Analgesia Dysphoria, sedation Diuresis Change in pupil size			
	Delta	Analgesia Respiratory depression Urine retention Dependence *human concern			
	Opioid- receptor-like receptor (ORL-1 NOP)				
Antagonists Naloxone	Used to reverse MOR agonists	Not an effective reversal for buprenorphine due to its high receptor affinity	30 min so re- administration may be necessary	IV- 1-2 min until action	