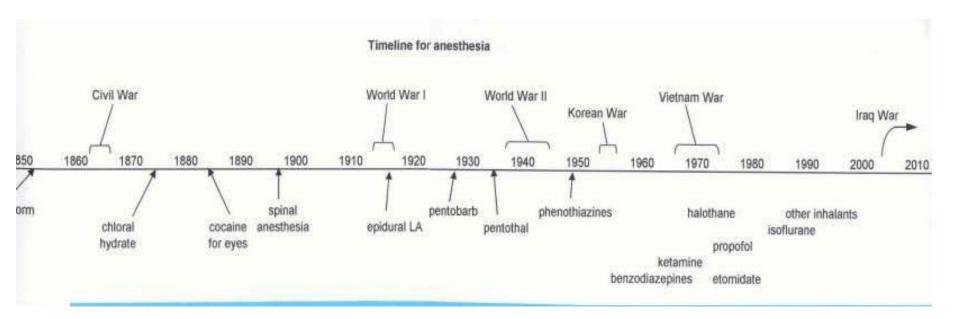
Pre-anasthesia

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History of Anesthesia



Classification of Anesthetic Agents

Route of administration

- Inhalant
- Injectable
- Oral
- Topical
- Time of administration
 - Preanesthetic
 - Induction
 - Maintenance

Principal effect

- Local vs. general
- Sedatives and tranquilizers vs. analgesics
- Neuromuscular blockers
- Anticholinergic agents
- Reversal agents

Chemistry

- Pharmacokinetics
- Pharmacodynamics
- Drug distribution
- Target tissues and stimulation
 - CNS—depression or stimulation

Preanesthetic Medications

- Calm or sedate excited animal
- Minimize adverse drug effects
- Reduce dose of concurrent drugs
- Smoother anesthetic induction and recovery
- Analgesia
- Muscle relaxation

- Route of administration affects onset of action and duration of effects
 - SC—slowest onset, longest duration
 - IM—faster onset, shorter duration
 - IV—fastest onset, shortest duration

Preanesthetic Anticholinergics

- Parasympatholytic drugs—block acetylcholine
 - Prevent and treat bradycardia
 - Decrease salivary secretions

Atropine and glycopyrrolate (dogs and cats)

- ✤IV, IM, SC, or IT
- Atropine—faster onset, shorter peak, shorter duration
- Glycopyrrolate—slower onset, longer peak, longer duration

Anticholinergic Effects

- CNS—limited effect
- Cardiovascular—prevent bradycardia
- Secretions—decrease
- Eye—mydriasis and corneal drying
- Bronchodilation

Anticholinergic Adverse Effects

- Cardiac arrhythmia
 - Contraindicated in animals with elevated heart rates or cardiac diseases
- Temporary bradycardia—atropine
- Thickened respiratory and salivary secretions
 - May lead to airway blockage—cats and ruminants
- Intestinal peristalsis inhibition
 - May lead to colic (horses) or bloat (ruminants)

Tranquilizers and Sedatives

- Phenothiazines
- Benzodiazepines

Alpha₂-adrenoceptor agonists

Alpha₂-antagonists

Phenothiazines—Acepromazine Maleate

Also known as acepromazine or "ace"

- Preanesthetic sedation
- Decrease dose of general anesthetic
- Ease induction and recovery
- May be used with opioids for minor procedures

Approved for horses, dogs, and cats

- Administered IV or IM
- No reversal agent
- Metabolized by liver
- Will slowly cross the placenta

Effects of Acepromazine

CNS

- Calming, reluctance to move, decreased interest in surroundings
- Sedation less pronounced in cats
- Not an analgesic

Cardiovascular System

- Peripheral vasodilation that leads to hypotension, increased heart rate, and hypothermia
- Protects against arrhythmias and decreases cardiac output

Respiratory system

Worsens depressive effect of other drugs

Gastrointestinal system

Antiemetic

Prevents histamine release and decreases allergic response

Adverse Effects of Acepromazine

* CNS

- Reduced seizure threshold
- May produce aggression or excitement

Cardiovascular system

Hypotension—dose dependent

Penile prolapse

Seen in horses and other large animals

May lead to permanent injury

Decreased PCV

Possibly due to splenic engorgement

Use of Acepromazine

- Dose and needle placement
- Increased potency and duration
 - Geriatrics, neonates, debilitated animals
- Breed considerations
 - Australian shepherds
 - Giant breeds, Boxers, Greyhounds
 - Terriers and cats

Benzodiazepines

Tranquilizers—controlled substances

- Diazepam
- Zolazepam
- Midazolam
- Rapid onset of action
- Short duration of action

Effects of Benzodiazepines

CNS

- Calming and antianxiety only in old or ill patients
- Not an effective sedative or analgesic
- Anticonvulsant—use with animals having seizures

Cardiovascular and respiratory systems
 Minimal effect with a high margin of safety

- Skeletal muscle relaxation
- Potentiate general anesthetics
- Appetite stimulation (cats and ruminants)

Adverse Effects of Benzodiazepines

CNS

- Disorientation and excitement—young, healthy dogs
- Dysphoria and aggression—cats
- Muscle fasciculations—horses
- Ataxia and recumbency—any large animal
- Diazepam must be given by IV slowly
 Oral diazepam in cats can cause liver failure

Use of Benzodiazepines

Diazepam

- Not water soluble
- Don't mix with water-soluble drugs
- Don't store in plastic
- Commonly used with ketamine to induce anesthesia in small animals and horses
- Administer IV slowly

Midazolam

- Water soluble
- Can be administered IM or SC
- Excellent sedative for swine, ferrets, rabbits, and birds
- Used in combination with ketamine to induce anesthesia in dogs, small mammals, and birds

Zolazepam

Available only as a component of Telazol[®]

- A powdered product
- Reconstituted with sterile water

Alpha₂-Adrenoceptor Agonists

- * Also written alpha₂-agonists or α_2 -agonists
- Noncontrolled agents
- Sedation, analgesia, and muscle relaxation
- Large and small animals—IM or IV
- Administered prior to minor procedures
- Readily reversed with alpha₂-antagonist

Alpha₂-Agonists

- Xylazine (Rompun, Anased)
- Detomidine (Dormosedan)
- Romifidine (Sedivet)
- Dexmedetomidine (Dexdomitor)

- Stimulates alpha₂ receptors of the sympathetic nervous system (SNS)
 Decrease release of norepinephrine
 No "fight-or-flight" response
- Sedation, analgesia, bradycardia, hypotension, and hypothermia
- Metabolized in liver; excreted in urine
- Rapid sedation; 1-2 hour duration

Effects of Alpha₂-Agonists

CNS

Dose-dependent sedation

Analgesia—short-acting

Cardiovascular system—early phase

- Dose-dependent vasoconstriction and hypertension
- Bradycardia
- Cardiac arrhythmias

Cardiovascular system—late phase Decreased cardiac output

Hypotension and further bradycardia

Respiratory system

- Dose-dependent depression
- Other effects
 - Muscle relaxation
 - Increased effect of other anesthetic agents
 - Vomiting—immediate response (dogs and cats)
 - Hyperglycemia—transient

Hypothermia

Adverse Effects of Alpha₂-Agonists

* CNS

- Change in behavior—varies with species
- Cardiovascular system
 - Bradycardia, hypotension, decreased output

Respiratory system

- Depression—varies from animal to animal
- More severe if given with other drugs

Increased urination

Gastrointestinal effects

Bloat—dogs, cattle, and horses

Salivation and regurgitation—cattle

Premature parturition—cattle (last trimester)

Sweating—horses

Absorbed through skin abrasions and mucous membranes

Wash off immediately

Use of Alpha₂-Agonists

- Use with caution; monitor patients closely
- Avoid use in geriatric, diabetic, pregnant, pediatric, or ill patients
- Administer anticholinergics 10-20 minutes prior

Alpha₂-Agonist—Xylazine

- 2% solution (small animals)
- 10% solution (horses)
- Use 1/10 horse dose in cattle
- Used mostly in large animals

Alpha₂-Agonist—Dexmedetomidine

- Dexdomitor[®]
- Most commonly used in dogs and cats
- Produces sedation and analgesia
- More potent and safer than xylazine
- Antagonist—atipamazole (Antisedan[®])
- Preanesthetic in low doses
- Can be mixed with other drugs

Alpha₂-Agonists—Detomidine and Romifidine

Detomidine

- Used in horses
- Sedation, analgesia, muscle relaxation
- Two times the duration of xylazine
- Standing sedation with butorphanol
- Romifidine
 - Produces less ataxia

Alpha₂-Antagonists

Reverse all effects of alpha₂-agonists

- Seneficial effects—for example, analgesia and sedation
- Detrimental effects—for example, bradycardia
- Wide margin of safety

Effects of overdose

- Neurological—excitement and muscle tremors
- Cardiovascular—hypotension and tachycardia
- Gastrointestinal—salivation and diarrhea

Dose is expressed as a ratio

Agonist to antagonist

10:1 means the dose of the antagonist is 1/10 of the dose of the agonist

Administer slowly by IV

 Reduce dose if more than 30 minutes has elapsed since the agonist was administered

Alpha₂-Antagonist—Tolazoline

- Nonspecific alpha₂-antagonist
- Used in ruminants at a 1:10 dose ratio with xylazine
- Reverses cardiovascular and sedative effects

Alpha₂-Antagonist—Yohimbine

- Used in dogs, cats, horses, and exotic species
- Reverses cardiovascular and sedative effects of xylazine
- Dose ratio is species dependent
 - Dogs and horses—10:1

Cats—2:1

Alpha₂-Antagonist—Atipamezole

- Antisedan[®]
- Specific antagonist for dexmedetomidine
- IM injection (IV in emergencies)
- Use ½ the dose in cats compared to dogs
- Reversal—5-10 minutes after IM injection

Opioids

- Derivatives of opium
- Opiates—naturally derived compounds
- Produce analgesia and sedation
- Anesthetic induction when combined with other drugs
- Classified as agonists, partial agonists, agonist-antagonists, or antagonists

Used of Opioids

Agonists

Morphine, hydromorphone, oxymorphone, fentanyl, and meperidine

Partial agonist

Buprenorphine

Agonist-antagonists

Butorphanol and nalbuphine

Antagonists

Naloxone, etorphine, and carfentenil

Preanesthetic

- Agonists, partial agonists, or agonist-antagonist
- May be used alone or in combination with
 - Tranquilizers
 - Anticholinergics
- Analgesia
 - Prevent and treat postoperative pain
 - Sed with tranquilizer to produce neuroleptanalgesia

Effects of Opioids

CNS

- Effect depends on many factors
- Dogs
 - Causes sedation
 - Narcosis
- Cats, horses, and ruminants
 - Causes CNS stimulation
 - Bizarre behavior patterns or dysphoria

Use lower dose

Analgesia

- Pure agonists are most effective against severe pain
- Used as a premedication for painful surgery

Cardiovascular system

Bradycardia

Respiratory system

Minimal decreased rate and tidal volume

Other Effects of Opioids

- Miosis in dogs
- Mydriasis in cats, horses, and ruminants
- Hypothermia in dogs
- Hyperthermia in cats
- Increased responsiveness to noise
- Sweating in horses
- Decreased urine production with urine retention

Adverse Effects of Opioids

CNS

Anxiety, disorientation, excitement, dysphoria

- Cardiovascular system
 - Pronounced bradycardia
- Respiratory system
 - Decreased respiration and tidal volume
 - Decreased PaO₂ and PaCO₂
 - Dose dependent with some agents
 - Ceiling effect with some agents

Gastrointestinal system

- Salivation and vomiting—small animals
- Initial diarrhea, vomiting, and flatulence
- Pretreat with atropine or acepromazine
- GI stasis follows initial GI stimulation
 - May predispose to colic in horses
 - Avoid administration to any animal with a GI obstruction