





# The Problems - Clinical

• Vomiting



# The Problems - Clinical

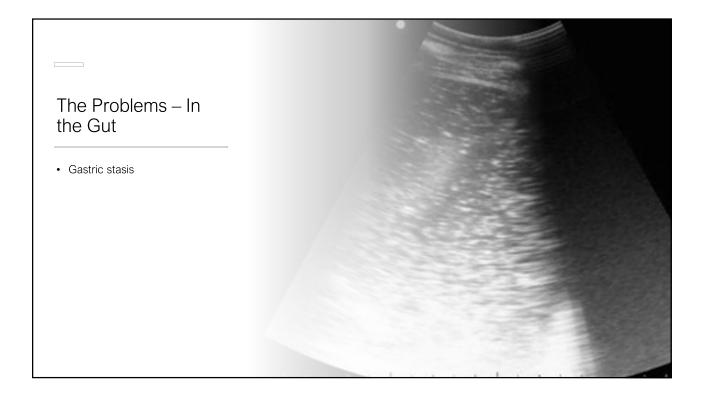
• +/- Oesophagitis

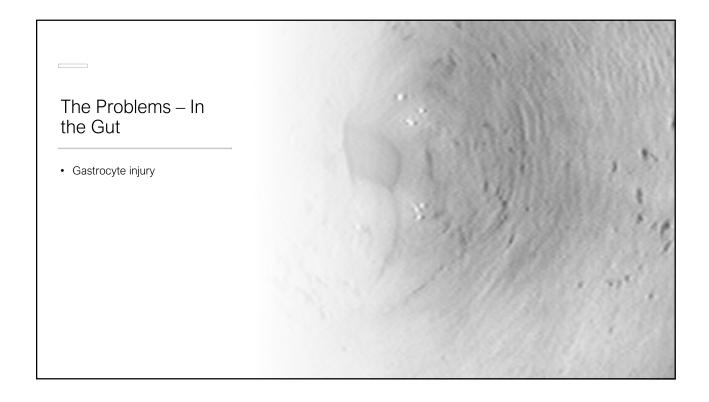


## The Problems - Clinical

• Diarrhoea

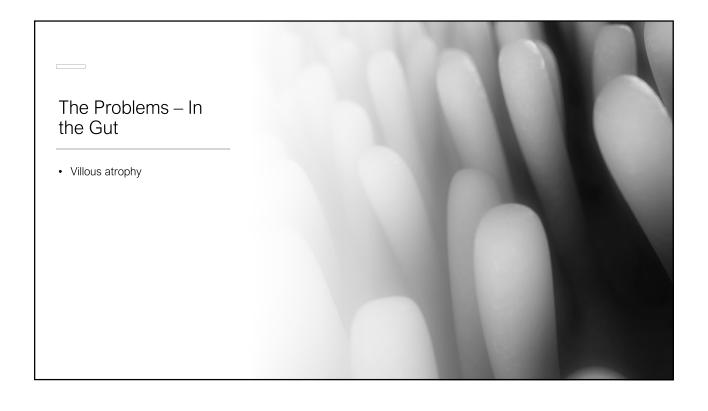




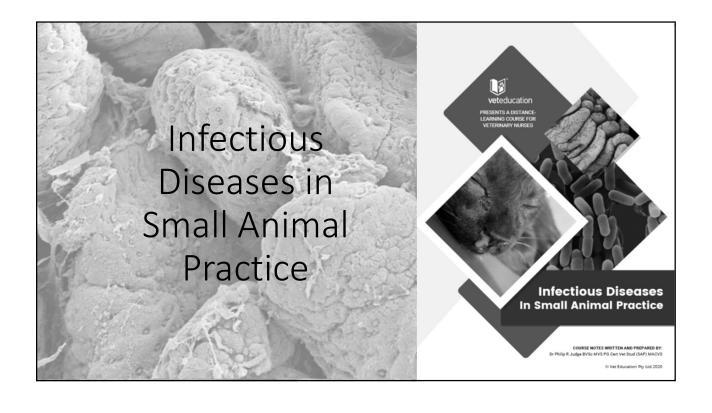


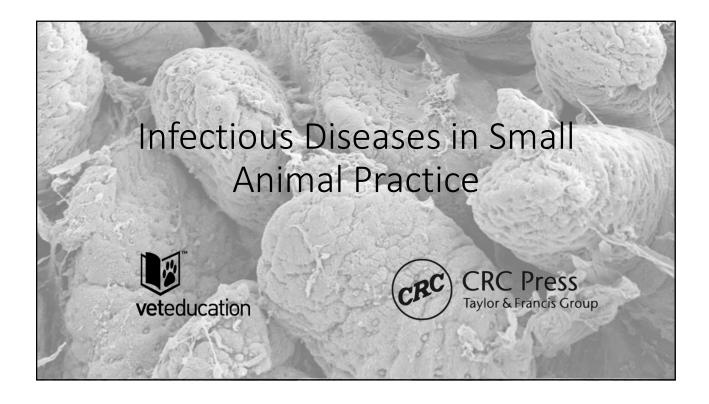




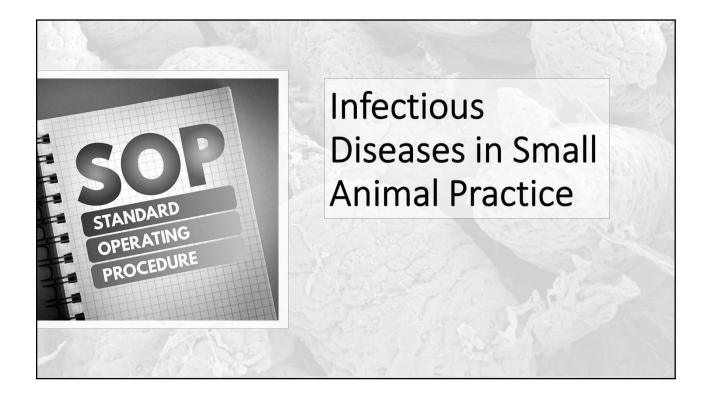




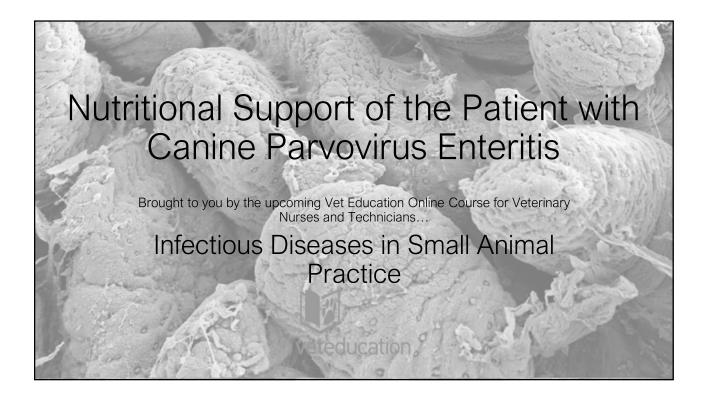


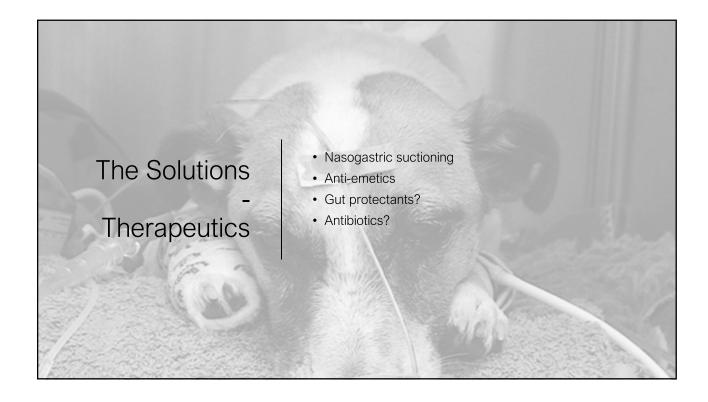


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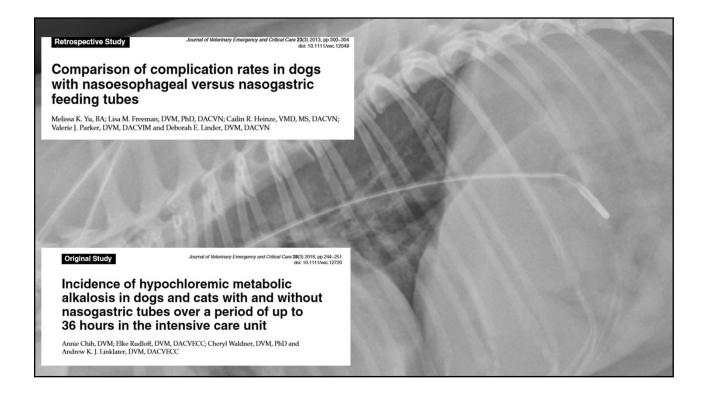


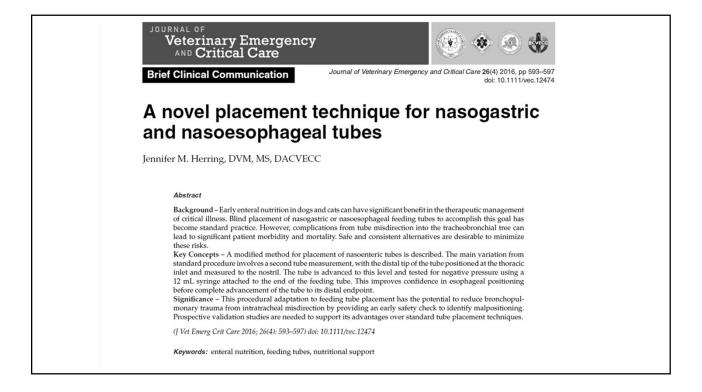














### A novel placement technique for nasogastric and nasoesophageal tubes

Jennifer M. Herring, DVM, MS, DACVECC

- Technique describing placement of nasogastric or naso-esophageal tubes, designed to reduce complications
  - Tracheal or bronchial rupture
  - Pulmonary tears
  - Pneumothorax
  - Pleural effusion
- Involves placing a mark on the tube from nares to thoracic inlet
  - Aspiration of air at this level should prompt tube re-direction
  - · Confirmation with radiography still recommended
  - · Minimises risk to distal pulmonary structures

Original Study

Journal of Veterinary Emergency and Critical Care 12(4) 2002, pp 227-233

#### Capnographic documentation of nasoesophageal and nasogastric feeding tube placement in dogs

Paula A. Johnson, DVM, F. A. Mann, DVM, MS, DACVS, DACVECC, John Dodam, DVM, MS, PhD, DACVA, Keith Branson, DVM, MS, DACVA, Colette Wagner-Mann, DVM, PhD, Mark A. Brady, DVM and Elizabeth Dunphy, DVM

Objective: To evaluate the ability of capnography to document proper placement of nasoesopha-geal (NE) and nasogastric (NG) feeding tubes. This study was conducted in 3 phases. Phase I of this study was designed in order to test the efficacy of capnography to distinguish placement of a feeding tube in the alimentary fract versus the respiratory tract. Phase II was designed in order to document that carbon dioxide (CO) could be measured through a polyvinyl chloride (PVC) feeding tube. Phase III was performed in order to evaluate the technique of continuous monitoring during insertion of the feeding tube into the esophagus and stomach as would be performed during a -divid-al-tube n lacement.

Phase III was performed in order to evaluate the technique of continuous monitoring during insertion of the feeding tube into the esophagus and stomach as would be performed during a clinical-tube placement.

Design: Prospective study.

Setting: Research laboratory.

Animals: 24 adult dogs.

Interventions: In Phase II, sedated dogs were instrumented with an intratracheal catheter and an 8 French feeding tube was placed down the endotracheal tube, then into the esophagus and later advanced into the stomach. In Phase II, dogs were anesthetized and an 8 French feeding tube was placed down the endotracheal tube, then into the esophagus and later advanced into the stomach. In Phase II, sedated dogs were instrumented with an 8 French feeding tube inserted intransally and then advanced to the level of the nasopharynx, distal esophagus and, lastly, the stomach. Fluoroscopy was used in order to determine location of the feeding tube.

Measurements and main results: Phase I measurements included respiratory rate and CO<sub>2</sub> from the trachea, esophagus, and stomach and pH of gastric fluid sample. Phase II measurements included respiratory and the stomach and pH of gastric fluid sample. Phase II measurements included respiratory rate and CO<sub>2</sub> from the trachea, leveline the distal esophagus, and feeding tube in the stomach. Plase III data collection included respiratory rate and CO<sub>2</sub> so the tube was passed through the usal cavity, nasopharynx, esophagus and stomach. Plase III disamples were collected from 5 of the 9 dogs and had pH values from 1.68 to 42 lin both phases, values for the respiratory rate and CO<sub>2</sub> from the tendous significant difference between the respiratory rate and CO<sub>2</sub> from the tendous significant difference between the respiratory rate and CO<sub>2</sub> and respiratory and so significant difference between the respiratory rate place of the respiratory rate of the respiratory rate of the respiratory rate of the second place of the respiratory rate of the respiratory rate of the respiratory rate of the second pl

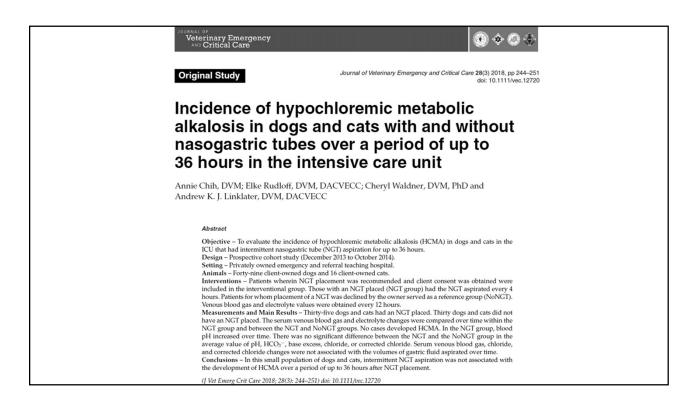
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- Prospective study
  - N = 24 adult dogs
  - 8 Fr nasogastric tube inserted under fluoroscopic guidance
  - Capnography measured at
    - Pharynx
    - Oesophagus
    - Stomach
- Results
  - When tube in oesophagus or stomach:
    - Respiratory rate = 0
    - Capnography = 0



Veterinary Emergency



Original Study

#### Incidence of hypochloremic metabolic alkalosis in dogs and cats with and without nasogastric tubes over a period of up to 36 hours in the intensive care unit

Annie Chih, DVM; Elke Rudloff, DVM, DACVECC; Cheryl Waldner, DVM, PhD and Andrew K. J. Linklater, DVM, DACVECC

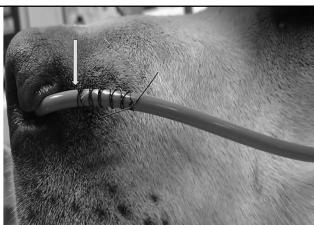
- Prospective study
  - 49 dogs

    - 16 cats
    - Diseases: pancreatitis, enterotomy, GDV, and other conditions
    - 35 with nasogastric tubes placed (23 dogs; 12 cats); 30 without
    - Nasogastric tubes suctioned q 4 hrs, with suctioned fluid discarded
    - Venous blood samples collected at baseline, and every 12 hrs until 36 hrs  $\,$ 

      - Co2
      - BE Sodium
      - Chloride
      - Bicarbonate

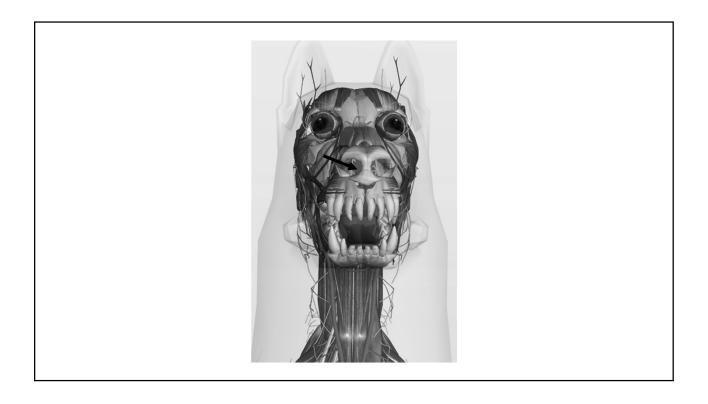
- Results:
  - No patient developed hypochloraemic metabolic alkalosis
  - No significant differences in pH, BE chloride or bicarbonate concentration between groups
  - Short-term nasogastric suctioning was not associated with  $\operatorname{HCMA}$

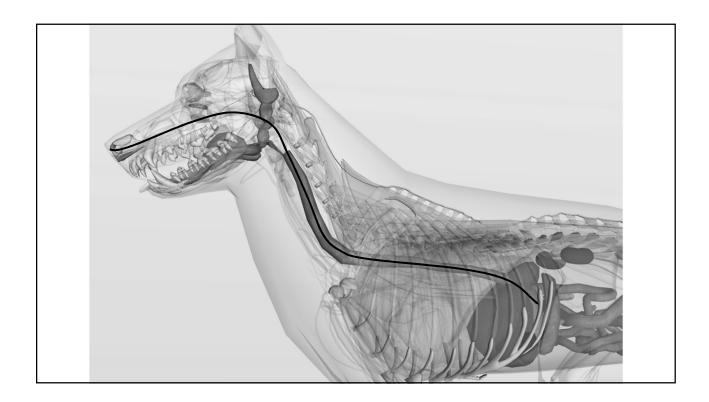


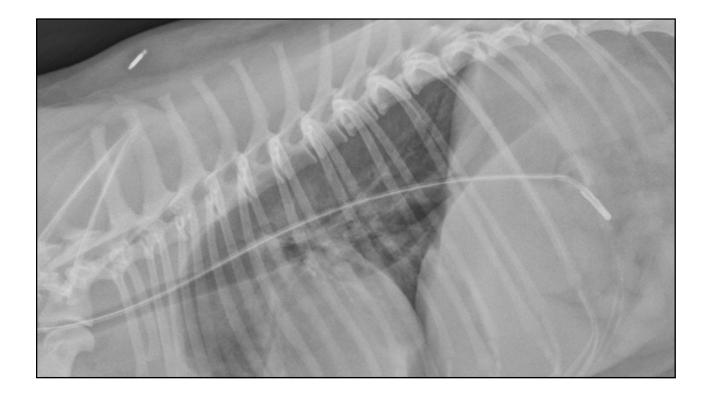


## Nasogastric Suctioning

Suction residual gastric volume q 1-2 hrs.











### **Anti-Emetics**

First-Line Anti-Emetics

- Maropitant
  - 1 mg/kg SC (IV) SQ q 24 hrs.
- Metoclopramide
  - 0.2-0.5 mg/kg slow IV q 6-8 hrs.
  - CRI: 0.4 mg/kg loading dose; then 0.3-0.5 mg/kg/hr
- Ondansetron
  - 0.1-0.2 mg/kg slow IV q 12-24 hrs. Can increase to 0.5 mg/kg

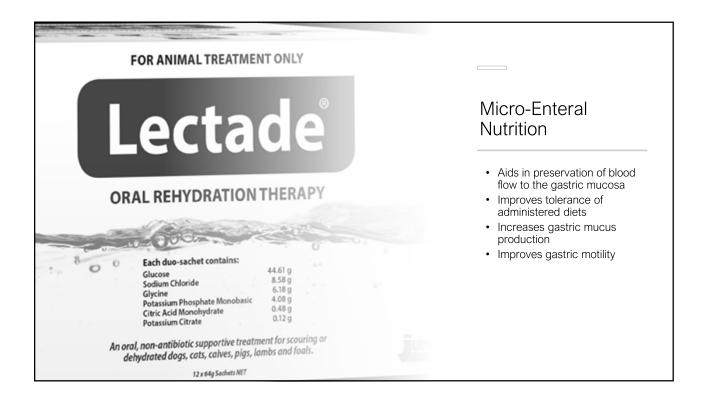


### **Anti-Emetics**

Adjunctive Therapy

- Butorphanol
  - 0.1 mg/kg/hr CRI
- Ranitidine
  - 0.5-2 mg/kg IV q 8-12 hrs.







## Micro-Enteral Nutrition

- Solutions
  - · Vytrate or Lectade
    - Balanced electrolyte and glucose solutions
    - Discard solutions made up after 12 hours
  - Lactated Ringers Solution Spiked with Glucose to make up 2.5% solution
    - Sterile
  - Able to be kept for up to 1 week after being made up
  - Addition of Potassium chloride, Amino Acids, Glycine and Glutamine may improve effectiveness





