

## Introduction

Obesity in domestic cats is commonly diagnosed and a growing concern in veterinary medicine because it predisposes individuals to health complications like diabetes mellitus. Feline diabetes mellitus is similar to human type II diabetes with pancreatic  $\beta$ -cell failure and insulin resistance made worse by obesity. Previous studies showed that, in overweight neutered-male cats, estrogen replacement therapy results in decreased food intake and possibly downregulates insulin production as observed in a glucose tolerance test.

## Hypotheses

- 1) Estrogen replacement will reduce the post-prandial plasma insulin to glucose ratio helping cats maintain euglycemia with less need for insulin
- 2) Bisphenol-A, an environmentally ubiquitous weak estrogen mimic, will have similar effects as estrogen (decrease food intake, reduce post-prandial insulin:glucose ratio)

## Methods

### Animals

- Six male-neutered cats (4-7 years)
- 4.4-8.6kg, variably overweight (body fat= 27-47%)

**Study design** (Latin-square, 3 treatments)

- Treatments: Oral 17β-estradiol (E2, 1.0 µg/kg/day), bisphenol-A (BPA, 50 µg/kg/day), or vehicle (ethanol, 1.0 µL/kg/day)
- Plasma collected at post-prandial intervals of 0,4,8,12,16 hours



# Effects of estrogen on post-prandial glucose and insulin in cats at risk for diabetes mellitus Lisa M. Mori<sup>1</sup>, Robert C. Backus<sup>2</sup>

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## **Conclusions/on-going research**

- Dose of E2 given appears physiologic and appropriate for replacement therapy to decrease ad libitum food intake toward pre-neuter amounts.
- Oral BPA is absorbed and accumulated, but not eliminated within 24 hrs.
- Post-prandial glucose and insulin concentration and insulin: glucose ratio are not negatively impacted by E2 replacement and allowable limits of BPA contamination of pet foods.
- Unconjugated (free) BPA analysis to be done.

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