

**Capsaicin-induced bladder hyperactivity**

**Model's advantages:**

- Capsaicin is a selective excitotoxin of C-fiber primary afferent neurons.
- Acts through the stimulation of a vanilloid receptor (VR1).
- Release of tachykinins and other mediators at both the peripheral and spinal cord level.
- Useful for quick investigation of the effect of drugs known to act on C-fiber afferents.

**Species:** rat, guinea pig

**Pathophysiological features:**

- Cystometrogram displays:
  - decrease in the intercontraction interval.
  - decrease in the pressure threshold for eliciting contractions.
- Increase C-Fos positive cells in the spinal cord (L6 level).

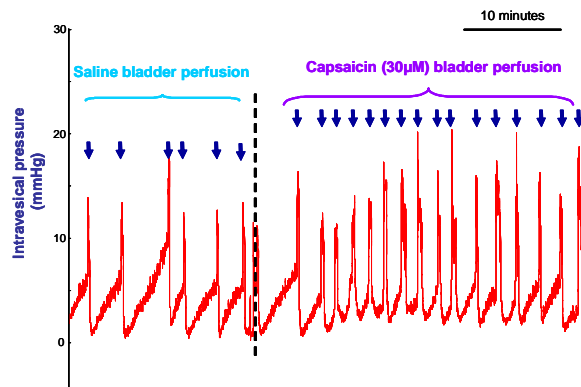


Figure 1: Representative cystometrograms showing the effect of capsaicin-induced bladder hyperactivity in anesthetized female rat. Arrows indicate the voiding contractions. (Pelvipharm, internal data)

**Summarized methodology:**

The bladder is perfused with continuous capsaicin (30 µM) at a rate of 50 µl/min while intravesical and blood pressure are monitored concomitantly.

**Related Pelvipharm bibliography:**

Caremel, R. et al. *Eur Urol* (2010) : 58(4):616-25

**NB:** Pelvipharm will gladly study the feasibility to fit this experimental model in order to meet its client's needs.

**Links to applicable experimental skills:**

- **Administration routes / regimen**
- **Plasma / urine / tissue collection**
- **In vivo experiments – anesthetized animals**
  - \* Urodynamic evaluation (anesthetized)
  - \* Bladder blood flow
  - \* Neural firing recording
- **Biochemistry (Plasma / Urine / Tissue)**
  - \* Spectrophotometric assays
  - \* Protein expression and activity
- **Histology**
  - \* Histomorphology
  - \* Histomorphometry
  - \* Oxydative fuorescence
- **Immunohistology / Confocal microscopy**
  - \* Protein expression – immunohistochemistry / immunofluorescence
  - \* Confocal microscopy