



Inulin propionate ester increases satiety and decreases appetite but does not affect gastric emptying in healthy humans

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The short chain fatty acid (SCFA) propionate stimulates free fatty acid receptors 2 and 3 (FFAR2, FFAR3) found on enteroendocrine L-cells and increases the release of the anorectic gut hormones glucagon like-peptide-1 (GLP-1) and peptide YY (PYY) *in vitro*⁽¹⁾. Infusion of GLP-1 and PYY significantly delays gastric emptying in humans^(2,3). Furthermore, *in vivo* data shows that intestinal infusions of SCFA significantly inhibits gastrointestinal motility via an increase in PYY release from the gastrointestinal tract⁽⁴⁾. Dietary supplementation with an inulin propionate ester (propionate bound to inulin through an ester linkage) significantly increases the post-prandial release of GLP-1 and PYY, whilst significantly decreasing food intake (unpublished findings). It is unknown whether the mode of action of inulin propionate ester on appetite and food intake is through a central action of PYY and GLP-1 or via a direct effect on gastric emptying.

We hypothesized that dietary supplementation with inulin propionate ester would lead to a reduction in appetite by delaying gastric emptying through the release of GLP-1 and PYY.

Fourteen healthy men and women (Mean \pm SEM age 31 \pm 2 years; body mass index (BMI) 24.0 \pm 0.9 kg/m²) were recruited and completed two study visits in a randomized, double-blind, crossover manner. On the study days, subjects attended having fasted overnight and were given 10 g inulin propionate ester or 10 g inulin control in a standard breakfast (641 kcal; 113 g CHO, 15 g fat, 16 g protein). At 300 min volunteers were served a standard lunch (354 kcal; 47 g CHO, 12 g fat, 12 g protein) together with 100 mg ¹³C-octanoic acid. Breath CO₂ was collected serially for 480 min after the standard lunch by exhaling alveolar breath samples. ¹³CO₂ enrichment was determined by isotope ratio mass spectrometry (IRMS). Subjective hunger, satiety, and nausea were monitored with the use of 100 mm visual analog scales (VAS). This study was completed in accordance with the Declaration of Helsinki.

Ingestion of 10 g of inulin propionate ester significantly increased subjective 'fullness' (Δ AUC_{0–540min} 24 296 \pm 2554 min \times mm inulin control vs. 27 573 \pm 1898 min \times mm propionate ester; $p < 0.05$), and decreased 'desire to eat' (Δ AUC_{0–540min} 20 044 \pm 2873 min \times mm inulin control vs. 17 404 \pm 2881 min \times mm propionate ester; $p < 0.05$) throughout the study period.

The time to 50% AUC excretion of ¹³C in breath (T_{1/2}), a proxy for gastric emptying rate, was not significantly different between acute propionate ester and control trials (T_{1/2} 180 min [95% CI, 163 to 198] control vs. 185 min [95% CI, 168 to 204] control, $P = 0.51$). These data demonstrate that acute ingestion of 10 g of inulin propionate ester increases satiety and reduces appetite in humans but this is not through an effect on gastric emptying, and it is most likely to be a centrally mediated action of GLP-1 and PYY.

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