## Obesity 1255

## An Exogenous Ketone Ester Modulates Appetite but Not Dietary Intake

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**Objectives:** Previous research suggests exogenous ketone esters (KE) suppress appetite by directly modulating regulatory hormones; however, their impact upon eating behaviors is unknown. The authors aimed to determine if the diminished appetite resulting from KE consumption is accompanied by a reduction in dietary intake.

**Methods:** After informed consent participants (n = 7) were recruited to a randomized cross-over trial. Participants recorded their diet for three consecutive days, starting the day prior to their first study appointment. During this visit, fasted participants were randomized to consume either a KE or matched dextrose placebo (DP) beverage. Blood samples were drawn at regular intervals and analyzed for  $\beta$ hydroxybutyrate (BHB), glucose, leptin and ghrelin. Appetite was self-reported using a visual analogue scale (VAS). One-week later participants were invited to a second visit where the study was repeated using the other beverage. Dietary data was analyzed using MyFood24 and statistical analysis was performed using Microsoft Excel and IBM SPSS (v.26).

Results: BHB increased 30 minutes after consuming the KE  $(0.21 \pm 0.20$  to  $4.21 \pm 0.66$  mmol/L) (P < 0.001) and remained elevated. Blood glucose increased 30 minutes after consuming the DP (4.87  $\pm$  0.42 to 8.11  $\pm$  1.41 mmol/L) (P < 0.001) and promptly returned to baseline. Although there were no changes in leptin levels, those who consumed the KE demonstrated suppressed ghrelin production 120 minutes after baseline (2430.00  $\pm$  323.46 to 1763.14  $\pm$  367.67 pg/mL) (P = 0.026). Furthermore, the VAS also revealed that 120 minutes after baseline participants who consumed the DP reported a greater desire to eat  $(+26.86 \pm 23.55 \text{ mm})$ (P = 0.038) and were less satisfied  $(-30.43 \pm 12.52 \text{ mm})$  (P = 0.003). Despite this, there was no significant differences in the calorie intake of those who consumed the KE compared to the DP on the day before (1941.06  $\pm$  1048.13 vs 1792.86  $\pm$  833.23 kcal), during  $(1594.64 \pm 677.07 \text{ vs} 1536.52 \pm 457.22 \text{ kcal})$ or after (1674.41  $\pm$  801.43 vs 1914.35  $\pm$  804.78 kcal) the study visits.

**Conclusions:** Consuming a KE, despite impacting upon selfreported measures of appetite and associated biomarkers, does not modulate dietary intake. This should be considered when assessing the potential role of KE for appetite management.

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