

HUMAN NUTRITION

Title: Effect of pork ingestion on postprandial mitochondrial protein synthesis and inflammation in healthy weight, overweight, and obese adults - **NPB # 16-012**

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Scientific Abstract:

Context:

Excess fat mass may diminish the anabolic potency of protein-rich food ingestion to stimulate muscle protein sub-fractional synthetic responses. However, the impact of adiposity on mitochondrial protein synthesis rates (MPS) after protein-rich food ingestion has not been thoroughly examined *in vivo* in humans.

Objective:

We compared basal and postprandial MPS and markers of muscle inflammation (Toll-like receptor 4 [TLR4] and myeloid differentiation primary response protein 88 [MyD88] protein content) in young adults with different BMIs.

Methods:

10 normal-weight (NW; BMI 22.7 ± 0.4 kg/m²), 10 overweight (OW; BMI 27.1 ± 0.5 kg/m²), and 10 obese (OB; BMI 35.9 ± 1.3 kg/m²) adults received primed continuous L-[ring-¹³C₆]phenylalanine infusions, blood sampling, and skeletal muscle biopsies before and after the ingestion of 170 g of pork.

Results:

Pork ingestion increased muscle TLR4 and MyD88 protein content in the OB group ($P < 0.05$), but not in the NW or OW groups. Basal MPS were similar between groups ($P > 0.05$). Pork ingestion stimulated MPS ($P < 0.001$) (0-300 min) in the NW (2.5 ± 0.6 -fold above baseline values), OW (1.7 ± 0.3 -fold), and OB groups (2.4 ± 0.5 -fold) with no group differences ($P > 0.05$).

Conclusions:

Protein-dense food ingestion promotes muscle inflammatory signaling only in obese adults. However, the consumption of a dinner-sized amount of protein strongly stimulated a postprandial MPS response irrespective of BMI. Our data suggest that alterations in postprandial mitochondrial protein synthesis are unlikely to contribute to compromised muscle macronutrient metabolism witnessed with obesity.

These research results were submitted in fulfillment of checkoff-funded research projects. This report is published directly as submitted by the project's principal investigator. This report has not been peer-reviewed.

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