

POSTER PRESENTATION

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# Thermogenic and hemodynamic effects of ingesting a pre-workout supplement with and without synephrine

YP Jung, C Goodenough, M Cho, A O'Connor, R Dalton, K Levers, E Galvan, N Barringer, F Ayadi, J Carter, M Koozechian, S Simbo, A Reyes, B Sanchez, A Coletta, C Rasmussen, R Kreider\*

From The Eleventh International Society of Sports Nutrition (ISSN) Conference and Expo  
Clearwater Beach, FL, USA. 20-21 June 2014

## Background

A number of nutritional strategies have been developed to optimize nutrient delivery prior to exercise. As a result, a number of pre-workout supplements have been developed to increase energy availability, promote vasodilation, and/or positively affect exercise capacity. The purpose of this study was to examine the acute effects of ingesting a pre-workout dietary supplement with and without synephrine on energy metabolism and cardiovascular hemodynamics.

## Methods

In a double-blind, crossover, randomized and placebo-controlled manner; 25 apparently healthy and recreationally active men and women ( $21.76 \pm 3.00$  yr,  $15.24 \pm 5.26\%$  fat,  $25.09 \pm 3.03$  kg/m<sup>2</sup>) volunteered to participate in this study and had resting blood pressure (BP), heart rate (HR), 12-lead electrocardiographs (ECG), and resting energy expenditure (REE) measured for 10 minutes. Participants then ingested in a randomized and counterbalanced manner a dextrose flavored placebo (P); a pre-workout supplement (PWS) containing 3.0 g beta alanine, 2 g creatine nitrate, 2 g arginine AKG, 300 mg of N-acetyl tyrosine, 270 mg caffeine, 15 mg of Mucuna pruriens; or, the PWS with 20 mg of synephrine (PWS+S). Metabolic changes were measured continuously while BP, HR, and ECG's were obtained every 10 minutes during the REE test. Participants repeated the experiment after a one week washout period with the alternate supplements in a randomized and counterbalanced manner. Data were analyzed by repeated measure MANOVA and are presented as means  $\pm$  SD or SEM from baseline. Consent to publish the results was obtained from all participants.

## Results

MANOVA analysis revealed a significant overall Wilks' Lambda time ( $p < 0.001$ ) and time x group interactions ( $p < 0.001$ ) for oxygen uptake (VO<sub>2</sub>), carbon dioxide production (VCO<sub>2</sub>), minute ventilation (Ve), respiratory exchange ratio (RER), and REE values. MANOVA Greenhouse-Geisser univariate analysis revealed significant interactions among groups in VCO<sub>2</sub> ( $p = 0.003$ ) and RER ( $p < 0.001$ ) with a trend toward significance in REE ( $p = 0.098$ ). Delta analysis revealed significant differences among groups in mean change in VO<sub>2</sub> (P:  $3.8 \pm 5.2$ ; PWS:  $15.4 \pm 5.2$ ; PWS+S:  $23.5 \pm 5.2$  ml/min;  $p = 0.03$ ), VCO<sub>2</sub> (P:  $12.5 \pm 5.1$ ; PWS:  $31.8 \pm 5.1$ ; PWS+S:  $37.7 \pm 5.1$  ml/min;  $p = 0.002$ ), RER (P:  $0.033 \pm 0.009$ ; PWS:  $0.071 \pm 0.009$ ; PWS+S:  $0.071 \pm 0.009$ ;  $p = 0.005$ ), and REE (P:  $0.034 \pm 0.025$ ; PWS:  $0.095 \pm 0.025$ ; PWS+S:  $0.132 \pm 0.025$  kcal/min;  $p = 0.02$ ) with significant differences observed among the P group and both supplemented groups. PWS-S ingestion promoted a more prominent increase in VO<sub>2</sub>, VCO<sub>2</sub>, and REE during the initial 5-10 minutes after ingestion with differences minimizing thereafter. Area under the curve (AUC) analysis of changes from baseline revealed that PWS+S and PWS supplementation resulted in significantly greater AUC values than P in VO<sub>2</sub> (PWS+S:  $1,034 \pm 584$ ; PWS:  $802 \pm 434$ ; P:  $684 \pm 376$ ;  $p = 0.01$ ); VCO<sub>2</sub> (PWS+S:  $1,372 \pm 604$ ; PWS:  $1,151 \pm 604$ ; P:  $634 \pm 262$ ;  $p < 0.01$ ); and RER (PWS+S:  $2.79 \pm 0.89$ ; PWS:  $2.44 \pm 0.98$ ; P:  $1.46 \pm 0.66$ ;  $p < 0.01$ ). There were no significant interaction effects for HR ( $p = 0.77$ ), SBP ( $p = 0.35$ ), or DBP ( $p = 0.65$ ) and there was no evidence of an increase in ECG assessed arrhythmias during the REE assessment.

\* Correspondence: rkreider@hlkn.tamu.edu  
Texas A&M University, College Station, Texas, USA

## Conclusion

Ingesting a PWS containing beta alanine, creatine nitrate, arginine AKG, N-Acetyl Tyrosine, caffeine, and Mucuna pruriens increased resting VO<sub>2</sub>, VCO<sub>2</sub>, RER, and tended to increase REE values in comparison to a placebo. Addition of 20 mg of synephrine to the PWS resulted in a greater increase in the metabolic response during the first 5-10 minutes after ingestion but differences were not as apparent thereafter and AUC values were not significantly different between the PWS and PWS+S groups. PWS and PWS+S ingestion did not result in a significantly different HR or BP responses during the REE test in comparison to P responses. Results indicate that ingestion of these pre-workout supplements promoted modest thermogenic response and that addition of 20 mg of synephrine to the PWC provided limited additional benefit.

## Acknowledgement

Supported by Woodbolt International, Inc. (Bryan, TX)

Published: 1 December 2014

doi:10.1186/1550-2783-11-S1-P35

**Cite this article as:** Jung et al.: Thermogenic and hemodynamic effects of ingesting a pre-workout supplement with and without synephrine. *Journal of the International Society of Sports Nutrition* 2014 **11**(Suppl 1):P35.

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