Introduction

The use of pre-workout supplement (PWS) is common in the active duty population, with 50% to 87% of soldiers reporting regular use.1–3 These supplements often contain long lists of vitamins, amino acids, energy supplements, and herbal formulations in their ingredient list. However, under the Dietary Supplement and Health Education Act of 1994, these products are not considered drugs, and therefore are not required to demonstrate purity, efficacy, or safety before distribution. These products are available without a prescription and without physician knowledge or approval. As a result, the individuals taking PWS may not be aware of the significant health risks associated with them. Furthermore, although physicians are aware of many of the signs and symptoms of adverse events due to PWS, very few of these events are reported.1 This impairs public health efforts to identify and monitor potentially dangerous products currently on the market. Previous case reports of adverse events associated with PWS use in the active duty population have included acute liver damage,4,5 rhabdomyolysis,6 ischemic colitis,7 acute psychosis,8 and ventricular fibrillation.

Case Presentation

A 23-year-old male navy soldier who was in excellent physical condition (last physical fitness test placed him above the 90th percentile), suffered a sudden cardiac arrest while running in formation with his battalion for about 5 minutes, was found to be pulseless; bystanders initiated CPR with the return of spontaneous circulation within 5 minutes, he subsequently developed 3 episodes of ventricular fibrillation (V-FIB) on route to the emergency department which was successfully defibrillated and aggressive resuscitation efforts were applied including active core cooling via an endovascular device, endotracheal intubation, and fluid restoration.
Sudden Unexplained Cardiac Arrest In Apparently Healthy Soldier: Is Pre-Workout Supplement Safe!

smokes half pack per day, occasional marijuana use and social drinking. He has a paternal grandmother who
had a heart attack at her 30. He described the daily exercise regimen composed of weight training and aerobic
activity approximately five times a week of varying time length; prior to each session, he ingested
a caffeine-containing drink (Red Bull) and pre workout supplement called Cellucor C4. On the day of the
cardiac arrest, he mixed the Red Bull and approximately one serving of Cellucor C4 with the engagement of
his exercise routine. Upon arrival at the hospital, physical examination was remarkable for hyperthermia
per protocol, intubated, blood pressure 127/70 mmHg, pulse 66, respiratory rate 17, and saturating at 97%.
Electrocardiogram (ECG) showed normal sinus rhythm, right axis deviation and small positive deflection (‘blip’)
buried in the end of the QRS complex in lead V1, II, III and aVF suggestive of epsilon wave. Initial laboratory
values were remarkable for a troponin I of 0.04 ng/mL, white blood cell count of 18000. Toxicology analyses
were negative. Laboratory evaluation revealed no findings consistent with a hypercoagulable state, nor was
there other evidence of systemic thromboembolic disease.

Transthoracic echocardiogram showed a normal left ventricle function. Coronary angiography was performed
showing normal coronary arteries. A cardiac MRI showed no evidence of myocardial fibrosis or Arrhythmogenic
right ventricular cardiomyopathy (ARVC). Subsequent cardiac genetic testing was done without yield. He was
eventually extubated and transferred to our step down unit. Cardiac electrophysiology (EP) study was performed
without inducible arrhythmias and ultimately underwent an ICD placement for secondary prevention.

Fig1. Electrocardiogram (ECG) showed normal sinus rhythm, right axis deviation and small positive deflection
(‘blip’) buried in the end of the QRS complex in lead V1, II, III and aVF suggestive of epsilon wave (Arrow)

DISCUSSION

Nutrient timing refers to the methodical, timed ingestion of carbohydrate, protein, fat and other dietary
supplements either before, during, or after physical activity [10]. Supplementation during the period immediately
preceding physical activity has become an increasingly popular strategy among competitive and recreational
athletes alike as a means of improving performance [11]. In response to this trend, manufacturers have developed
pre-workout supplements (PWS), which typically combine caffeine with any number of purported ergogenic
substances, such as dimethylamylamine, creatine, arginine, β-alanine, taurine, nitrate and phosphates. As the
number of PWS (unregulated dietary supplements) available on the market grows, each containing their own
“proprietary blend” of active ingredients, it must be determined which, if any, are safe for chronic consumption. This becomes particularly important as concerns have arisen over the concept of proprietary blends, namely the fact that the FDA does not monitor the amounts of ingredients used in these blends or the accuracy of product labeling by manufacturers [11]. While some dietary supplement labels instruct consumers to seek the advice of a health care professional before using the products, the labels usually do not disclose all ingredients or their precise amounts, and evidence to support the purported performance-enhancing benefits is generally lacking. There is limited evidence to support the use of some pre-workout supplement ingredients. For example, in one small placebo-controlled study (n = 12), the use of the energy drink Red Bull (containing caffeine and taurine) 40 minutes before a simulated cycling time trial appeared to provide a meaningful ergogenic benefit; in another small study (n = 12), the use of a similar caffeine-containing product (Redline) by strength-trained athletes was found to improve reaction time, energy, and mental focus relative to placebo use. However, published evidence on the use of the other ingredients listed above is scant, inconclusive, or conflicting. Adverse effects reported in association with pre-workout supplements include gastrointestinal symptoms, cardiac arrhythmia, blood pressure increases, and potential effects on lipids and blood glucose.

Although an expanding body of literature reports adverse cardiovascular outcomes in association with the use of synephrine-containing products, our case report is, to our knowledge, the first in which active duty soldier with no history of identifiable cardiovascular risk factors experienced V-FIB arrest shortly after ingesting a Cellocur C4. While the cause of his SCA remains uncertain, it is plausible that the pre-workout C4 might triggered his V-FIB event as it contains a variety of sympathomimetic and stimulant compounds.

The reported specificity qualified the epsilon wave not only as a major criterion for ARVC diagnosis but also as a hallmark of the disease. It is, however, a relatively rare finding, manifesting itself only in from 1% to 25% of ARVC patients [12]. It is important to acknowledge that the epsilon wave is not specific to ARVC only and has been reported in patients with other conditions such as sarcoidosis [13], myocarditis [14], or acute RV infarction [15]. Moreover, an iatrogenic form of epsilon wave has been described, occurring after catheter ablation in the RV [16].

Our case highlights the potential health hazards associated with athletic-performance-enhancing and weight-reduction supplements that contain sympathomimetic compounds similar to ephedra. In light of these facts, as well as the empiric risk of synergism in the above-mentioned agents, we believe that performance enhancement use was the precipitant of this patient’s cardiac event, and we believe that greater FDA and some government agencies such as US military involvement in the regulation of such supplements is warranted.

CONCLUSION

Although evidence exists to support the performance-enhancement efficacy of some pre-workout ingredients as standalone agents, published data on combination products are scant, inconclusive, or conflicting. The safety of these products may be compromised if users consume larger-than-recommended amounts or use more than one product. Thus, the safety of Cellocur C4 remains an open question and needs more investigation. Regulatory action may be necessary.

REFERENCES


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