Effects of Combined Treatment with Branched-Chain Amino Acids, Citric Acid, L-Carnitine, Coenzyme Q10, Zinc, and Various Vitamins in Tumor-Bearing Mice

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A functional dietary supplement (FDS) containing Coenzyme Q10, branched-chain amino acids and L-carnitine was administered to tumor-bearing mice, investigating its effects on tumor and muscle tissues. Experiment (A): B16 melanoma cells were implanted subcutaneously into the right side of the abdomen of 8- to 9-week-old C57BL/6J mice. The mice were divided into two groups: a FDS group that received oral administration of FDS (n=10), and a control group that received oral administration of glucose (n=10). The moribund condition was used as the endpoint, and median survival time was determined. Experiment (B): On day 21 after tumor implantation, tumors, soleus muscle, gastrocnemius muscle, and suprahyoid muscles were collected. Tumor and muscle weight and other aspects were evaluated in each group: FDS group (n=15) and control group (n=10). The median survival time was comparable (21 d in the FDS group vs. 18 d in the control group, p=0.30). However, cumulative food intake was significantly higher in the FDS group than the control group (p=0.011). Metastasis of melanoma to the lung was observed in the control group but not in the FDS group (p=0.043). The weight of the suprahyoid muscles was significantly higher in the FDS group than in the control group (p=0.0045). The weight of the tumor was significantly lower in the FDS group than in the control group (p=0.013). The results possibly suggest oral administration of FDS in tumor-bearing mice enhances the maintenance of suprahyoid muscles, resulting in an extended feeding period and suppression of tumor growth and metastasis.

Key words cancer therapy; cachexia; metabolism; branched-chain amino acid; functional maintenance

Cancer cachexia is a syndrome that affects many patients with advanced cancer. It is characterized by refractory malnutrition and has significant impact on prognosis and QOL. Managing cachexia is therefore a high priority when providing palliative care. Cachexia is a complex metabolic syndrome, with loss of muscle mass being a primary symptom. Development of cachexia involves many factors and accompanies progressive functional impairment, but the pathogenesis remains unknown.1,2) Recent studies suggest that metabolic and neuroendocrinological abnormalities in cancer cachexia are caused by inflammatory cytokines secreted by the tumor and host cells and other tumor-derived substances. It is essential for cancer patients to maintain good nutritional status for better prognosis and QOL. As such, managing cancer cachexia is a major goal for the future of cancer therapy and palliative care. Since its foundation in 2003, our laboratory has implemented a unique nutritional support system to manage cancer cachexia.3–5) With the nutritional support, patients with cachexia have shown reduced symptoms and improved bodily function through control of metabolic dynamics. We have developed a functional dietary supplement (FDS) for improving symptoms and function, containing Coenzyme Q10 (CoQ10), branched-chain amino acids (BCAA), L-carnitine and citric acid and reported the effectiveness of its use in clinical settings.6,7) In this study, we have experimentally evaluated the effects of FDS on tumors and muscle tissues to corroborate the clinical findings.

METHODS

Mice C57BL/6J mice (7-week-old males) were purchased from Japan SLC Inc. (Shizuoka, Japan). After 1 to 2 weeks of acclimatization, a tumor-bearing model was established by first removing hair from the right side of the abdomen and then subcutaneously implanting B16 melanoma cells (B16F6). These mice were maintained under laboratory animal management at the Nanakuri Laboratory of Animal Models for Human Diseases, Fujita Health University. The mice were kept alone or in pairs in plastic cages with wood chip bedding and were fed normal diet MF (Oriental Yeast Co., Ltd., Tokyo, Japan) in an animal facility controlled at a temperature of 23±5°C with humidity at 60±5% with a 12-h light/dark cycle.

Tumor Cells B16 melanoma cells (B16F6) have been used in tumor-bearing models and cancer cachexia models in a number of studies.8) Because melanoma cells typically form subcutaneous tumors, the size and shape of the tumor can be easily evaluated. The highly malignant cell line B16F6 can also metastasize to the lung.9) B16 melanoma cells were grown in RPMI-1640 (Wako Pure Chemical Industries, Ltd., Osaka, Japan) with 10% fetal bovine serum (Gibco, Life Technologies, Carlsbad, CA, U.S.A.) in humidified atmosphere containing 5% CO2 at 37°C. The medium was replaced every 2 to 3 d, and the subcultures were obtained by harvesting using 0.25% trypsin/ethylenediaminetetraacetic acid (EDTA) (Sigma-Aldrich, St. Louis, MO, U.S.A.).

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For subcutaneous implantation, $2 \times 10^6$ B16 melanoma cells (B16F6) in 0.2 mL of the medium were prepared for each mouse and implanted into the right side of the abdomen.  

**Reagents** The FDS sample used in this study was Inner Power®, purchased from Otsuka Pharmaceutical Co., Ltd. Each 125-g package of FDS contained 139 kcal of energy and the following ingredients: 1.7 g of proteins, 0 g of fat, 13.5 g of sugars, 33 g of carbohydrate, 2 g of fiber, 90 mg of sodium, 3 mg of zinc, 0.3 mg of copper, 1000 mg of citric acid, 2500 mg of BCAA (917 mg of leucine, 833 mg of isoleucine, 750 mg of valine), 30 mg of CoQ10, 50 mg of l-carnitine, 1 mg of vitamin B1, 3 mg of vitamin B2, 0.9 mg of vitamin B6, 1.5 µg of vitamin B12, and 10 mg of vitamin E. The contents of FDS were mixed in a blender into a gel consistency and administered orally. The的规定 of radioimmunoprecipitation assay (RIPA) buffer (Pierce Biotechnology, Rockford, IL, U.S.A.) containing 1% protease inhibitor cocktail (Nacalai Tesque, Inc., Kyoto, Japan). The homogenate was cooling centrifuge at $80^\circ$C for protein and mRNA measurements to be conducted at a later time.

Protein contents were determined as follows: A portion of the frozen gastrocnemius muscle stored at $-80^\circ$C was excised and weighed. The muscle tissue was homogenized in 10 volumes of radioimmunoprecipitation assay (RIPA) buffer (Pierce Biotechnology, Rockford, IL, U.S.A.) containing 1% protease inhibitor cocktail (Nacalai Tesque, Inc., Kyoto, Japan). The homogenate was cooling centrifuge at $80^\circ$C for 10 min, and the supernatant was used for protein assay using BCA Protein Assay Kit (Pierce Biotechnology).

MuRF-1, mTOR, and VEGF mRNA levels were determined as follows: Total RNA was extracted from the tumor tissue and gastrocnemius muscle stored at $-80^\circ$C using High Pure RNA Tissue Kit (Roche Diagnostics GmbH, Mannheim, Germany). cDNA was synthesized using Transcriptor Universal cDNA Master Kit (Roche Diagnostics GmbH) and quantified by real-time PCR using TaqMan probes. Relative mRNA expression levels were calculated by normalizing to an internal control, eukaryotic translation elongation factor 2 (28EF-2) mRNA.

**Experimental Procedures**

**Experiment (A)**

Male C57BL/6j mice (7 weeks old) were allowed ad libitum access to food and water and acclimatized for 1 to 2 weeks. At 8 to 9 weeks of age, B16 melanoma (B16F6) cells ($2 \times 10^6$ cells in 0.2 mL) were subcutaneously implanted into the right side of the abdomen (Fig. 1). The mice were divided into two groups ($n=10$ each). FDS [0.1 mL (0.1 kcal)/10 g body weight (BW)/d] was orally administered in the FDS group, and glucose [0.1 mL (0.1 kcal)/10 g BW/d] was orally administered in the control group. Food intake and general health status of the mice were recorded daily. The moribund condition was used as the experimental endpoint.

**Experiment (B)**

Tumor-bearing mice (8 to 9 weeks of age) were prepared as in Experiment A. The mice were divided into FDS and control groups ($n=15$ each), and their health status was recorded daily. We designed Experiment (A) and observed the condition of the tumor-bearing mice to select a day as an endpoint. Based on the results, the endpoint was set as day 21, which was the median survival time. On day 21, whole blood was collected by cardiac puncture under isoflurane anesthesia. Subsequently, the tumor was removed, and the soleus and gastrocnemius muscles of both hind limbs, and suprahryoid muscles were collected. After weighing, the tissue samples were immediately frozen in liquid nitrogen and stored at $-80^\circ$C for protein and mRNA measurements to be conducted at a later time.

**Statistical Analysis** Statistical analysis was performed using Pro 11.2.1 statistical analysis software (SAS Institute, Inc., NC, U.S.A.). To evaluate incidence of lung metastasis, difference between the FDS and control groups was determined using Fisher’s exact test. For other measurements, difference between the two groups was determined using the Wilcoxon rank-sum test. All contentative variables were expressed as median and interquartile range (IQR). Survival rate was estimated using the Kaplan–Meier method with Log-rank test. The change in food intake was evaluated using two-way repeated measure ANOVA by SPSS 21 software (SPSS Inc., Chicago, IL, U.S.A.). In all analyses, $p<0.05$ was considered statistically significant.

**Ethical Considerations** The care and use of the animals was according to the Regulations for the Management of Laboratory Animals at Fujita Health University, which is accredited by the Japanese Association of Laboratory Animal Facilities of Public and Private Universities (JALAP). The experimental protocols were approved by the Institutional Animal Care and Use Committee of Fujita Health University (Permit No. S0141).

**RESULTS**

**Experiment (A)**

**Survival Rate:** The median survival time (MST) was 21 d (IQR: 15–27 d) in the FDS group, and 18 d (IQR: 8–25 d) in the control group ($p=0.30$) (Fig. 2).

**Cumulative Food Intake:** Food intake was analyzed for the first 9 d when all 10 animals were alive in both FDS and control groups. The change in food intake had significant difference between the groups (from 5.62 to 25.42 mg in the FDS group and from 5.44 to 22.01 mg in the control group).
Survival rate was measured. The median survival time (MST) was 21 d (IQR: 15–27 d) in the FDS group, and 18 d (IQR: 8–25 d) in the control group.

**Fig. 2. Survival Rate in Tumor-Bearing Mice**

The number of survival time after implanting the B16 melanoma cell to mice was measured. The median survival time (MST) was 21 d (IQR: 15–27 d) in the FDS group, and 18 d (IQR: 8–25 d) in the control group.

Food intake was analyzed for the first 9 d when all 10 animals were alive in both FDS and control groups. The change in food intake had significant difference between the groups (*p=0.011).

**Fig. 3. Cumulative Food Intake**

Food intake was analyzed for the first 9 d when all 10 animals were alive in both FDS and control groups. The change in food intake had significant difference between the groups (*p=0.011).

**Lung Metastasis:** We performed dissection of the lungs. Lung metastasis was not present macroscopically in the 10 mice in the FDS group, while lung metastasis was present macroscopically in 4 out of 10 mice in the control group (*p=0.043).

**Experiment (B)**

**Loss of Body Weight:** Changes in body weight from before tumor implantation to day 21 (after tumor removal) were comparable between the FDS groups [2.1% (IQR: −0.72–8.83)] and control groups [2.4% (IQR: −0.89–0.90)] (*p=0.9).

**Muscle Tissues:** Gastrocnemius and soleus muscles of both hind limbs, and suprahyoid muscles were collected, and the tissue weight was compared between the FDS and control groups. The median weight of suprahyoid muscles was 0.021 g (IQR: 0.017–0.02) in the FDS group and 0.017 g (IQR: 0.014–0.017) in the control group, indicating that the FDS group maintained significantly more muscle mass (*p=0.0045) (Table 1). There was no significant difference in the weight of gastrocnemius and soleus muscles, and protein contents of the gastrocnemius muscle between the two groups. Relative expression levels of mTOR and MuRF1 in left gastrocnemius muscle were determined. The median mTOR expression was 0.96 (IQR: 0.89–1.06) in the FDS group and 0.94 (IQR: 0.88–1.12) in the control group. The median MuRF1 expression was 0.93 (IQR: 0.79–0.98) in the FDS group and 1.01 (IQR: 0.91–1.11) in the control group. Although expression levels of mTOR and MuRF1 were not significantly different between the two groups, the median values were higher in the FDS group, indicating a trend for better muscle growth and less muscle atrophy in the FDS group (Table 2).

**Tumor Weight:** The median tumor weight was significantly lower in the FDS group [4.50 g (IQR: 3.96–5.44)] compared to the control group [6.08 g (IQR: 5.12–6.76)] (*p=0.013) (Fig. 4). When the ratio of the tumor weight to the net body weight (weight after tumor removal) was compared, the median value was significantly lower in the FDS group [23.24–31.05] (IQR: 0.56–12.34) than in the control group [27.1% (IQR: 17.39–24.95)] (*p=0.016) (Fig. 5).

**Relative mTOR and VEGF Expression Levels in the Tumor Tissue:** The median value of relative mTOR expression within the tumor tissue showed no significant difference between the FDS group [0.96 (IQR: 0.89–1.06)] and the control group [0.94 (IQR: 0.88–1.12)] (*p=0.82). The median value of relative VEGF expression didn't show significant difference between the FDS group [0.70 (IQR: 0.59–0.80)] and the control group [0.67 (IQR: 0.64–0.90)] (*p=0.79) (Table 3).
In our ongoing effort, we have developed a dietary supplement for improving symptoms and function, named “CoQ10+BCCA+L-carnitine (FDS),” and reported its effectiveness in clinical cases of cachexia. To our knowledge, this study is the first to evaluate the effects of FDS on tumors and muscle tissues, focusing on mechanistic aspects. The results possibly suggest oral administration of FDS in tumor-bearing mice enhances the maintenance of suprahyoid muscles, resulting in an extended feeding period and suppression of tumor growth and metastasis.

The most important issue in this study, cachexia-associated loss of suprahyoid muscles was significantly reduced in the FDS group. To date, there is no report on the effect of cancer cachexia on muscles involved in mastication, including suprahyoid muscles. Our study revealed that cancer cachexia may affect mastication muscles before affecting hind limb muscles, and that FDS administration could ameliorate the sarcopenia. In cancer cachexia-associated anorexia, ghrelin has drawn much attention because of its role in the regulation of gas- 

Effect of CoQ10 against hepatocellular carcinoma was reported by Fouad et al. CoQ10 significantly reduced peroxidation of lipids and prevented the loss of reduced glutathione (GSH) and superoxide dismutase activity (SOD). It also reduced the increase of tumor necrosis factor-α (TNF-α) and nitric oxide (NO) in the liver of the rats with hepatocellular carcinoma. Histopathology of the liver revealed that dysplastic...
changes were ameliorated by CoQ10.

Antioxidant effect of zinc was reviewed by Prasad. In a human model of zinc deficiency, even mild zinc deficiency caused an increase in interleukin (IL)-1β production in monocytes ex vivo. Wong et al. reported that zinc deficiency caused abnormal activation of immune cells and dysregulation of IL-6. Taken together, these findings suggest that zinc deficiency itself can activate macrophages, induce proinflammatory cytokines, and increase oxidative stress.

There is a concern that one of the BCAA components in FDS, leucine, may facilitate tumor growth. Liu et al. reported that leucine supplementation enhanced pancreatic cancer growth in both lean and overweight mice. On the other hand, Diaz-Meco demonstrated that melanoma cells were highly sensitive to leucine deprivation, which caused inhibition of autophagy.

Reference Intake for Japanese set by the Ministry of Health, Labour and Welfare is 39 mg of leucine, 20 mg of isoleucine, and 26 mg of valine per 1 kg of body weight. These amounts roughly correspond to 2 packs of Inner Power® per day for a 60 kg human. The BCAA dose used in our study was in accordance with the recommended volume for mice, 0.1 mL/10 g BW, which contained 0.7336 mg of leucine, 0.6664 mg of isoleucine, and 0.6 mg of valine per 10 g of body weight. In animal studies, the equivalent doses of test substances are often tens of times higher than standard doses in humans. The dose we used in the current study was relatively low (equivalent to 2 to 3 times the standard dose for humans). While effect of BCAA alone on protein synthesis in cancer patients has been evaluated, data are scarce for complex mixtures such as FDS. We therefore used the original composition of Inner Power® in this study.

In future study, the dose load should be optimized by extracting individual components of FDS and creating various combinations of the components.

The results suggest the possibility that oral administration of FDS in tumor-bearing mice enhances the maintenance of supraphyroid muscles, resulting in an extended feeding period and suppression of tumor growth and metastasis.

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Conflict of Interest The authors declare no conflict of interest.

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