Note

Effects of Fucoidan from Mozuku on Human Stomach Cell Lines

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Received October 8, 2005; Accepted June 5, 2006

Cladosiphon okamuranus (Okinawamozuku) is a species of Mozuku known to contain the most fucoidan of any brown algae. Fucoidan is a sulfated hetero polysaccharide with various biological activities. We isolated and purified fucoidan from C. okamuranus. The yield of fucoidan was 1.8% (w/w) based on wet algae from C. okamuranus. The sulfate content and molecular weight of the fucoidan were 9.8% (w/w) and approximately 3,200,000, respectively. We examined the effects of fucoidan on human stomach cell lines. Fucoidan from C. okamuranus showed an inhibitory effect on anticancer agent 5-fluorouracil (5-FU) in Hs677st normal stomach cells with minimum inhibition of the original effect of 5-FU in cancer cells. Furthermore, fucoidan from C. okamuranus showed growth inhibition of stomach cancer cells but did not show any effects on normal cells. Thus, fucoidan from Mozuku, especially C. okamuranus, may be useful for cancer therapy.

Keywords: fucoidan, Mozuku, Cladosiphon okamuranus

Introduction

Mozuku are classified as brown algae, and most Mozuku are produced by artificially seeded culture nets in Okinawa, Japan. Among the various Mozuku, C. okamuranus and N. decipiens are edible seaweeds, and are typically served in salads with vinegar (Sunomono). Brown algae possess fucoidan and alginate in their wall matrix (Kloareg et al., 1986). Mozuku contains the most fucoidan of any brown seaweed. Fucoidan is a fucose-containing sulfated hetero polysaccharide, but details of their structure remain unknown. Fucoidan is not uniform, and their structure differ depending on the species they are derived from (Chevolut et al., 2001; Sakai et al., 2002). Fucoidan has many biological activities, including anticoagulation (Kuznetsova et al., 2003; Nishino et al., 1991), antiviral (Lee et al., 2004; Peeprame et al., 2001) and antitumor activities (Aisa et al., 2005; Koyanagi et al., 2003). The anticoagulant activity of high molecular weight fucoidan extracted from Ecklonia kurome was reported to be dependent on the sulfate content and molecular weight, as in numerous other sulfated polysaccharides. Lee et al. reported that the fucoidan from sporophyll of Undaria pinnatifida (Mekabu) showed potent antiviral activities against herpes simplex virus and human cytomegalovirus. Regarding the anti-tumor effects of fucoidan, many studies reported effects on leukemia cells and lung cancer cells both in vitro and in vivo. Recently, Aisa et al. reported that the fucoidan from Fucus vesiculosus induced apoptosis in human lymphoma HS-Sultan cell lines. In human beings, fucoidan is ingested with food. However, only a few studies have identified fucoidan in food, or have used digestive organ cells. Therefore, in the present research, we studied the effects of fucoidan on stomach cell lines with or without an anticancer agent.

Materials and Methods

Cell lines and culture conditions The stomach cell lines used in this study were obtained from the American Type Culture Collection (ATCC) and Riken Cell Bank (Saitama, Japan). According to ATCC, Hs677st (CRL-7407) cells were obtained from stomach fibroblasts in a 62-year-old Caucasian woman at 16 weeks gestation. The stomach cancer cell line MKN45 (RCB1001) were a gastric adenocarcinoma (Motoyama et al., 1986). Both cell lines were cultured in a 5% CO2 humidified incubator at 37°C. Hs677st cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM, Nihon Pharmaceutical Co., Ltd., Tokyo, Japan) containing 4.5 μg/ml glucose on collagen-coated culture plates. MKN45 cells were cultured in RPMI 1640 (Nihon Pharmaceutical Co., Ltd.). All media contained 10% cell growth promoting factor Daigo’s GF21 (Wako Pure Chemical Industries, Osaka, Japan), 60 μg/ml streptomycin (Sigma Chemical Co., Tokyo, Japan), and 50 μg/ml kanamycin (Wako Pure Chemical Industries, Osaka, Japan).
Extraction and purification of fucoidan from Mozuku

Wet and salted Mozuku (C. okamuranus), which was commercially cultured on nets in the sea around Iheya Island in Okinawa, was used in this study. Mozuku was mixed with same doses of purified water. The mixture was boiled for 1 h and filtered. The filtrate was desalted using an electrodialysis system (Micro Acilyzer S3, Asahi Kasei Corporation, Tokyo, Japan). Then, the desalted extract was freeze-dried using a lyophilizer, and was used as the crude fucoidan sample. 0.02 M or less sodium sulfate was added to the filtrate, and CTAB was added at 3 times the volume of the crude fucoidan (Fujikawa et al., 1975). The precipitate was washed using 30 mg/ml KCl solution. Some of the fucoidan was eluted with different concentrations of KCl solution. The eluted solution was precipitated using ethanol, and dried. This was used as the purified fucoidan sample.

Molecular weights (MWs) were determined using high-performance steric exclusion chromatography (HPSEC) in 0.15 M NaCl, 0.05 M NaH₂PO₄ buffer at pH 7.0, using pullulans (Shodex STANDARD P-82; Showa Denko K.K., Tokyo, Japan) as standards (Nardella et al., 1996; Haroun-Bouhedja et al., 2000). An Asahipak GF-710 (Showa Denko K.K., Tokyo, Japan) column was used. Polysaccharide content was calculated according to the rate of high molecular side peak area against all peaks area. Sulfate content of fucoidan was determined by the Dodgson's method (Dodgson et al., 1961).

Assay of the effects on stomach cells with or without 5-FU

Fucoidan from Fucus vesiculosus (Sigma Chemical Co., Tokyo, Japan) and sodium alginate (Kimitsu Chemical Industries) were used for comparison. Phosphate buffered saline (PBS) was used as a control in order to investigate the effects of 5-FU only.

Hs 677.st and MKN45 cells were cultured in each medium, treated with trypsin, and washed with the appropriate medium. 90 μL culture solution of the 2,000 cells/well cells were in 96-well microplates. 10 μL of each polysaccharide solution were added to microplate wells to a concentration of 1.0 mg/mL. For each cell line, the same concentration was used for eight wells per experiment.

After a 4 h pre-incubation period, 5-FU was added to the appropriate wells at 50 μg/mL, and plates were incubated for 4 days at 37°C. The growth of each cell line cultured in the presence or absence of polysaccharides with 5-FU was monitored using a modified MTT assay (Mosmann et al., 1983). After culture, 10 μL MTT (5.0 mg/mL) solution in PBS was added to each well, and incubated for 4 h at 37°C. Then, 100 μL 10% SDS solution was added to wells to dissolve the MTT crystals. We measured absorbance at 570 nm, and subtracted absorbance at 630 nm. The number of each type of stomach cells grown in culture medium only was defined as 100% relative growth. In the experiment of the effects of fucoidan on stomach cells on the presence of 5-FU, recovery rate or inhibitory rate were calculated according to the rate of increasing relative growth with each polysaccharides compared to that of the control.

Statistical analysis

Comparison of two means were made using Student's t-test (Fisher, 1958).

Results and Discussion

Extraction and purification of fucoidan from Mozuku

We prepared crude fucoidan from Mozuku, C. okamuranus. Molecular weights and sulfate content of the fucoidan are shown in Table 1. The polysaccharide content of crude fucoidan from C. okamuranus was 65%. We further purified and isolated one kind of fucoidan from Mozuku. At first, fucoidan from C. okamuranus was purified with an estimated yield of 1.0% (w/w) based on wet algae. The sulfate content and molecular weight of the fucoidan were 9.8% (w/w) and approximately 3,200,000, respectively. The molecular weights of fucoidan prepared in the present study are the highest reported to date, when compared with previous studies (Tako et al., 1996) that reported values under 500,000. In previous studies, these fucoidan were extracted in acidic conditions using hydrochloric acid for the purpose of high yield. However, in the present study, fucoidan were extracted in neutral conditions. These results show that fucoidan prepared in the present study are closer to native fucoidan in Mozuku. This indicates that we are able to estimate the biological activity of

<table>
<thead>
<tr>
<th>Polysaccharides</th>
<th>M.w.</th>
<th>Content (%)</th>
<th>Sulfate (%)</th>
<th>Yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fucoidan from F. vesiculosus</td>
<td>104,000</td>
<td>84</td>
<td>18.8</td>
<td>-</td>
</tr>
<tr>
<td>Sodium alginate</td>
<td>700,000</td>
<td>94</td>
<td>0</td>
<td>-</td>
</tr>
<tr>
<td>Crude fucoidan from C. okamuranus</td>
<td>3,080,000</td>
<td>65</td>
<td>5.6</td>
<td>1.30</td>
</tr>
<tr>
<td>Purified fucoidan from C. okamuranus</td>
<td>3,190,000</td>
<td>94</td>
<td>9.8</td>
<td>0.99</td>
</tr>
</tbody>
</table>

a) Determined by HPSEC using pullulans as standards. b) Content of each polysaccharide was calculated using the rate of two peak areas detected at HPSEC. c) Sulfate content is expressed as a percentage of SO₄ content per polysaccharide weight. d) Based on wet algae.
Effects of fucoidan on stomach cells in the presence of 5-FU Initially, relative growth of Hs 677.st cells in the presence of both polysaccharides and 5-FU was compared to conditions with the addition of buffer as control. Relative growth of Hs 677.st cells in conditions with addition of buffer as control was approximately 90%, and relative growth in the presence of both 5-FU and fucoidan from *F. vesiculosus* or sodium alginate were almost identical. In addition, the recovery rate of fucoidan from *F. vesiculosus* or sodium alginate was only a single-digit% level. However, relative growth with both 5-FU and fucoidan from *C. okamuranus* was increased to 100% (Fig. 1A). Furthermore, the recovery rate of fucoidan from Mozuku was 17%. These results indicate that fucoidan from *C. okamuranus* reduced the damage of anticancer agent 5-FU in normal cells.

Subsequently, the relative growth of MKN45 stomach cancer cells in the presence of polysaccharides was determined (Fig. 1B). Relative growth in the presence of both 5-FU and fucoidan from *F. vesiculosus* or *C. okamuranus* was slightly lower than relative growth with 5-FU only. The inhibitory rates of fucoidan from *F. vesiculosus* or *C. okamuranus* were −32% and −14%. These results indicate that fucoidan from *F. vesiculosus* or *C. okamuranus* hardly inhibited the original effect of anticancer agent 5-FU in cancer cells. Relative growth in the presence of both 5-FU and sodium alginate was similar to Relative growth with 5-FU only. An inhibitory rate of only 9% was observed.

These results indicate that fucoidan from *C. okamuranus* protect human stomach cells from growth inhibition by 5-FU. In other words, fucoidan from *C. okamuranus* inhibited the anticancer agent 5-FU in Hs 677.st normal stomach cells, but not in MKN45 stomach cancer cells. We suspect that high molecular weight fucoidan from *C.
okamuranus interacts with some substances on the surface of stomach cells, and therefore protects the stomach cells. Mechanisms of the effects on stomach cells in the presence of the anticancer agent 5-FU may be related to mechanisms involved in the inhibition of fucoidan in the adhesion of Helicobacter pylori (Shibata et al., 1999; Shibata et al., 2000; Nagaoka et al., 2000).

Effects of fucoidan on stomach cells  To further assess the effects of fucoidan itself on stomach cancer cells, the relative growth of MKN45 cells in the presence of crude or purified fucoidan was examined (Fig. 2A). While sodium alginate did not show any effect on the growth of stomach cancer cells, the relative growth of cancer cells with fucoidan from F. vesiculosus was reduced to approximately 30%. Fucoidan from F. vesiculosus showed inhibitory effects on the growth of stomach cancer cells. The relative growth of MKN45 cancer cells in the presence of crude fucoidan from C. okamuranus was reduced to approximately 40%. The relative growth of cancer cells with purified fucoidan from C. okamuranus was approximately 50%. In contrast, the relative growth of Hs 677.st cells in the presence of all fucoidan was similar to the control, indicating that fucoidan did not affect normal cell growth (Fig. 2B), but only affected cancer cell lines.

When we observed the morphological features of stomach cells treated with polysaccharides, MKN45 and Hs 677.st cells treated with all varieties of polysaccharides were similar to the control (data not shown).

These results indicated that fucoidan from C. okamuranus show growth inhibitory effects on stomach cancer cells. Many studies reported that fucoidan induce the apoptosis of cancer cells. However, apoptosis induction by fucoidan from Mozuku was not clearly observed in the present study. We suspect that growth inhibition by

Fig. 2. Effect of polysaccharides on human stomach cell lines without 5-FU.
A: MKN45, B: Hs 677.st. 90μl culture solution of the 2,000 cells/well cells were in 96-well microplates. 10μl of each polysaccharide solution were added to microplate wells to a concentration of 1.0μg/ml. The plates were incubated for 4 days at 37°C and the growth of each cell line was monitored using a modified MTT assay. The number of stomach cells grown in culture medium only is defined as 100% relative growth. Values represent the average±SD for relative growth (%). *P<0.01, compared with the control (Fisher, 1986).

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fucoidan from *C. okamuranus* on stomach cancer cells may relate not only to apoptosis induction but also to inhibitory effects of adhesion (Liu et al., 2000) on stomach cancer cells.

We consider that fucoidan from *C. okamuranus* may be suitable for cancer therapy, and that it could be applied to inhibitory medicine, in order to treat side effects of 5-FU anticancer agent in the future. We expect that fucoidan from Mozuku could be developed into a health food, a medicine, and an applied product. Therefore, we anticipate further research using other anticancer reagents and cell lines.

**Acknowledgements** We would like to express our appreciation to the Industrial Research Institute of Tottori Prefecture for their support for the analysis of polysaccharides and their help with the cell culture techniques.

**References**


