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Running title: Cell Growth Promoting Activity of Ascophyllan

Effects of Sulfated Fucan, Ascophyllan, from the Brown Alga Ascophyllum nodosum on Various Cell Lines: A Comparative Study on Ascophyllan and Fucoidan

Zedong JIANG, 1 Takasi OKIMURA, 2 Takeshi YOKOSE, 1,3 Yasuhiro YAMASAKI, 4 Kenichi YAMAGUCHI, 1,4 and Tatsuya ODA1, 4*

1 Graduate School of Science and Technology, Nagasaki University, Nagasaki 852-8521, Japan

2 Research and Development Division, Hayashikane Sangyo Co., Ltd., Shimonoseki, Yamaguchi 750-8608, Japan

3 Nagasaki Prefectural Institute for Environment Research and Public Health, 11306-11, Ikeda 2-chome, Omura, Nagasaki, 856-0026, Japan

4 Division of Biochemistry, Faculty of Fisheries, Nagasaki University, Nagasaki 852-8521, Japan.

Key words: Ascophyllum nodosum; ascophyllan; fucoidan; sulfated fucan; cytotoxicity; growth promotion; MDCK cells

*To whom correspondence should be addressed. Division of Biochemistry, Faculty of Fisheries, Nagasaki University, Nagasaki 852-8521, Japan. Tel: +81-95-819-2831; Fax: +81-95-819-2831; E-mail: t-oda@nagasaki-u.ac.jp
**Abbreviations:** BSE, bovine spongiform encephalopathy; FBS, fetal bovine serum; G-CSF, granulocyte colony-stimulating factor; α-MEM, alpha-modified Eagle’s minimum essential medium; TNF-α, tumor necrosis factor-α
Abstract: The effects of fucose-containing sulfated polysaccharides, ascophyllan and fucoidan, isolated from the brown alga *Ascophyllum nodosum*, on the growth of various cell lines (MDCK, Vero, PtK₁, CHO, HeLa, and XC) were investigated. In a colony formation assay, ascophyllan and fucoidan showed potent cytotoxic effects on Vero and XC cells, while other cell lines were relatively resistant to these polysaccharides. Almost no significant effects of these polysaccharides were observed in the cell lines tested using the Alamar blue cytotoxicity assay over 48 h with varying initial cell densities (2500–20,000 cells/well) in growth medium. Interestingly, a significant growth promoting effect of ascophyllan on MDCK cells was observed, whereas treatment with fucoidan showed growth suppressive effects on this cell line under the same experimental conditions. These results suggest that ascophyllan is distinguishable from fucoidan in terms of their bioactivities. This is the first report of the growth promoting effects of a sulfated fucan on a mammalian cell line under normal growth conditions.
INTRODUCTION

Marine organisms are not only very important resources as food, feed, and energy, but they are also rich sources of structurally diverse bioactive compounds with valuable pharmaceutical and biomedical potentials. Marine algae are an especially major source of industrially important polysaccharides, including agar, carrageenan, and alginate. In addition to their importance in the food and cosmetic industries, polysaccharides derived from marine algae have been drawing much attention as important bioactive natural products. It has been shown that algal polysaccharides have a wide spectrum of activity in various biological systems depending on the compositions, entire structures, and the physicochemical properties. Some of them even show therapeutic effects on viral infection and tumor metastasis (1-4). A description of the use of seaweed products for medical purposes is found in traditional Chinese herbal medicine as early as the sixteenth century (5, 6).

A common structural feature of sulfated polysaccharides is that they contain sulfate half-esters in their sugar residues. Among the sulfated polysaccharides, the sulfated fucans found in marine brown algae, often called fucoidans, are well known to have potent anticoagulant and antithrombotic activities. Algal fucoidans are mainly found in Fucales and Laminariales, but are also present in Chordariales, Dictyotales, Dictyosiphonales, Ectocarpales, and Scytosiphonales. In fact, this kind of sulfated polysaccharide has been discovered in all the brown algae (Phaeophyceae) investigated so far, but seems to be absent in green algae (Chlorophyceae), red algae (Rhodophyceae), and golden algae (Xanthophyceae), as well as in freshwater algae and terrestrial plants (7). Previous studies have demonstrated that sulfated fucans have numerous other biological properties such as anti-viral, anti-tumor, anti-inflammatory, and apoptosis-inducing activities. All of these activities give polysaccharides a multitude of potential applications in human health care. Besides that, biomaterials derived from plant origin generally have
an advantage in that there is an absence of potential risk of contamination from animal viruses and pathogens of bovine spongiform encephalopathy (BSE).

Although several fucoidan fractions have been prepared from various brown algal species so far, their precise structures are still debated. A reason for that stems from difficulties in their extraction and isolation in pure forms (8). Furthermore, their compositions seem to change according to the algal species from which the compound is taken, the extraction process, and even the harvest seasons and local climatic conditions (7, 9). Such heterogeneity and polydispersity limit the study of their structures (8, 10). The partial characterization of fucan fractions prepared from Fucus vesiculosus suggested the existence of different types of sulfated polysaccharides that were distinguishable from fucoidans (11). In fact, ascophyllan (xylofucoglycuronan) has been isolated from the brown alga Ascophyllum nodosum as a sulfated fucan preparation distinguished from fucoidan, by the presence of a backbone of uronic acid with fucose-containing branches (3-O-D-xylosyl-l-fucose-4-sulfate) (12, 13). A. nodosum is often used in the industrial-scale preparation of alginate; however, it seems that the effective use of residual fucan fractions have not been conducted yet. In our previous study, the presence of ascophyllan was confirmed as a separated fraction from the fucoidan fraction in extracts prepared from A. nodosum. The yields of alginate, fucoidan, and ascophyllan fractions obtained from 40 g of dry milled A. nodosum were 4.7, 0.5, and 0.76 g, respectively (14). We found that both ascophyllan and fucoidan isolated from A. nodosum showed similar cytotoxic effects on U937 cells in a concentration-dependent manner. Furthermore, we found that ascophyllan was capable of inducing the secretion of tumor necrosis factor-α (TNF-α) and granulocyte colony-stimulating factor (G-CSF) in the mouse macrophage cell line RAW264.7 (14). These results suggest that ascophyllan isolated from A. nodosum is a polysaccharide with multiple bioactivities that is an attractive candidate for using in food and pharmaceutical industries. Since fucoidan has also been reported to induce TNF-α secretion from monocytes (15), it seems that there are some similarities between ascophyllan and fucoidan in their biological activities.
There may also be of some common structural features between the two polysaccharides; they both have fucose and sulfate half-ester groups as main components. On the other hand, according to previous reports, the anticoagulant activities of ascophyllan and fucoidan seem to be different (16, 17). The characterization of sulfated fucan fractions extracted from *A. nodosum* suggested that fucoidan-like polysaccharides, but not ascophyllan-like polysaccharides, are mainly responsible for the anticoagulant activity of the whole fucan preparation (16). Another study also suggested that fucoidan-like polysaccharides showed much higher anticoagulant activity than ascophyllan-like polysaccharides (17). To gain basic information on the biological activities of ascophyllan and fucoidan at the cellular level, we compared the cytotoxic effects of ascophyllan and fucoidan on various cell lines. Unexpectedly, we found that ascophyllan has growth-promoting activity on MDCK cells, while fucoidan was rather toxic to this cell line.

**MATERIALS AND METHODS**

**Preparation of ascophyllan and fucoidan samples** *A. nodosum* harvested off the coast of Norway was obtained from KAISEI Co. Shimonoseki, Japan. Ascophyllan and fucoidan were prepared from powdered *A. nodosum* using the methods reported previously (14). In brief, the alginate fraction was removed from a hot water-extract by acid precipitation and subsequent digestion with alginate lyase. Ascophyllan and fucoidan fractions were obtained from the residual fucan fraction based on their differential solubilities in aqueous ethanol. Composition analyses confirmed that ascophyllan was a heteropolysaccharide (xylofucoglycuronan) consisting of fucose, xylose, uronic acid, and sulfate half-esters in approximately equimolecular proportions, which is consistent with previous reports (14). The composition of the preparation was obviously distinct from fucoidan preparations.

**Cell culture** Vero (African green monkey kidney), MDCK (Madin-Darby canine
kidney), and PtK₁ (potoroo rat kangaroo kidney), XC (rat sarcoma), CHO (Chinese hamster ovary), and HeLa (human epithelia carcinoma) cells were cultured in α-minimum essential medium (α-MEM) supplemented with 10% fetal bovine serum (FBS), 10 µg/ml each of adenosine, guanosine, cytidine, and thymidine, penicillin (100 IU/ml), and streptomycin (100 µg/ml) in a humidified atmosphere with 5% CO₂ and 95% air as described previously (18).

**Measuring cytotoxicity of ascophyllan and fucoidan on mammalian cell lines** The cytotoxicity of the polysaccharides was measured in terms of the inhibition of colony formation as described previously, with slight modifications (18). In brief, adherent cells (100 cells/well) were cultured in 96-well plates, with varying concentrations of polysaccharides in the growth medium for 5 days. The number of colonies formed was counted after staining with 1% methylene blue in 50% methanol. Clusters of 30 or more cells were considered as colonies. The colony forming efficiency of all cell lines used was between 90% and 95%. The cytotoxicity of the polysaccharides was also assessed using the Alamar blue assay, as described previously (19). In brief, adherent cells (2500–20,000 cells/well) were cultured in 96-well plates varying concentrations of polysaccharides in the growth medium for 48 h. Then, Alamar blue reagent was added to the cells at a final concentration of 10%. After incubation for 3 h at 37°C, the absorbance of untreated cells and those treated with ascophyllan or fucoidan was measured at 570 nm and 600 nm in a MPR-A4i2 microplate reader (Tosoh Co., Tokyo, Japan). To check the number of viable MDCK cells, the trypan blue dye exclusion test was used.

**Statistical analyses** Data were expressed as mean ± SD. Fisher’s protected least significant difference (PLSD) for multiple comparisons, after one-way ANOVA, was used to analyze data (SPSS version 16.0, SPSS, Chicago, IL). Differences were considered significant at $p<0.01$. 
RESULTS AND DISCUSSION

To examine the effects of ascophyllan and fucoidan on the proliferation of various cell lines, colony formation assays were carried out. As shown in Fig. 1, both ascophyllan and fucoidan demonstrated potent cytotoxic effects on Vero and XC cells in a similar concentration-dependent manner, while other cell lines were relatively resistant to these polysaccharides. These results suggest that the cytotoxic potency of ascophyllan and fucoidan is dependent on the cell lines, with some cell lines appearing to be highly sensitive and others highly resistant towards these polysaccharides. The similarity between the two polysaccharides in terms of their cell line-dependent cytotoxicity may also suggest that there are common mechanisms of intoxication for the polysaccharides. Both ascophyllan and fucoidan are characterized as sulfated polysaccharides extracted from a brown alga, and they are mainly composed of L-fucose and sulfate half-esters in common. Although the exact cytotoxic mechanisms of these sulfated polysaccharides are still unclear, it has been reported that fucoidan was able to induce apoptotic cell death in the human lymphoma HS-Sultan cell line through the activation of caspase-3 (20). Our recent study has demonstrated that ascophyllan also induces apoptosis in U937 cells, as judged by typical apoptotic nuclear morphological changes and DNA fragmentation (14). Furthermore, other reports indicated that hyper-sulfation of fucoidan resulted in an increase in its anti-tumor and anti-angiogenic effects (21, 22). In contrast, it has been reported that desulfated fucoidan showed decreased biological activities compared to the natural fucoidan (23, 24). The ester sulfate has been proposed to exhibit certain biological roles for cellular recognition through specific binding to cell surface receptors (25). Based on these findings, one can speculate that the ester sulfate residues of ascophyllan and fucoidan may be active components responsible for a wide-variety of biological activities, including cytotoxicity. On the other hand, alginate isolated from same powdered *A. nodosum*, which consists of mannuronate and guluronate, showed no significant toxic effects at least on the cell lines tested in this study (Fig. 1). Although
various biological activities of alginate and its oligomers have been found in \textit{in vivo} and \textit{in vitro} systems (26-30), no cytotoxic or direct cell killing activity of alginates has been reported so far. Porphyran, an alga-derived polysaccharide containing sulfate ester, is known to show inhibitory effects on the proliferation of tumor cells (31). In this case, the sulfate esters of the bioactive polysaccharides may play an important role in the signaling pathway leading to the arrest of the proliferation of sensitive cells. A lack of cytotoxic effects of alginate may be partly due to the absence of a sulfate ester group.

To further examine the differences in the cell line-dependent cytotoxicity of ascophyllan and fucoidan, we conducted Alamar blue assays in which sub-confluent monolayers of cells were incubated in the presence of varying concentrations of polysaccharides. As shown in Fig. 2, no significant cytotoxic effects of these polysaccharides were observed even in Vero and XC cell. Although the exact reason for the discrepancy between the results obtained by two assays is uncertain now, this may be derived from several different experimental conditions. One possibility is that the quite different initial cell densities between the two assays may have led to different sensitivities for the polysaccharides, even within the same cell line. In the colony formation assay, the initial cell numbers were 100 cells per well, which is a much lower cell density than that used in the Alamar blue assay. However, at least under the cell densities tested (2500–20,000 cells/well in a 96-well plate), no significant differences in ascophyllan or fucoidan were observed in the six cell lines using the Alamar blue assay. Another possibility that seems more likely responsible for the different susceptibility of the cells to the polysaccharides is the different incubation periods of the two assays. Colony formation assays take 5 days, while Alamar blue assays take 48 h. It is likely that ascophyllan and fucoidan may require long contact time with the target cells to cause profound cytotoxic effects. Obviously, further studies are necessary to clarify this point as well as the cytotoxic mechanisms by which these polysaccharides act.

During the course of investigating the cytotoxicity of ascophyllan, we realized that ascophyllan promotes the proliferation of MDCK cells. In fact, the size of MDCK cell
colonies formed in the presence of ascophyllan evidently became larger than those of control colonies without ascophyllan (Fig. 3A). The Alamar blue assay confirmed that ascophyllan is capable of promoting the growth of MDCK cells in a concentration-dependent manner (Fig. 3B). Furthermore, the growth curve of MDCK cells cultured in the presence of 1000 μg/ml of ascophyllan was significantly higher than that of control cells cultured in the absence of ascophyllan (Fig. 3C). In both assays, the growth suppressive effects of fucoidan on MDCK cells were observed. These results clearly indicate that there is a difference in the bioactivities of ascophyllan and fucoidan in terms of their effects on the growth of MDCK cells. To our knowledge, this is the first report that ascophyllan or even algal-derived sulfated polysaccharides can promote the growth of certain cell lines under normal growth conditions. Regarding the effects of saccharides on the growth of certain mammalian cells, it has been reported that alginate oligosaccharides stimulate or enhance vascular endothelial growth factor-mediated growth of human endothelial cells and the growth of keratinocyte in the presence of epidermal growth factor (32, 33). Since no significant effects of alginate oligosaccharides were observed on the cells in the absence of the growth factors, it has been considered that alginate oligosaccharides may stimulate proliferative activity of growth factors by a synergistic mechanism. In addition to mammalian systems, it has been reported that enzymatically depolymerized alginate oligomers promoted the growth of bifidobacteria, whereas the original alginate polysaccharide had no effect (34). Alginate oligomers with an average molecular weight of 1800 Da, prepared with bacterial alginate lyase, increased shoot elongation after germination of komatsuna (Brassica rapa var. pervidis) seeds (35). Iwasaki et al. have reported that an alginate oligosaccharide mixture had promoting activities for the root growth of lettuce seedlings (36). Root growth-promoting activity of alginate oligomer on carrot and rice plants was also reported (37). In addition to these terrestrial plant cells, our recent study has demonstrated that the growth of a unicellular marine microalga, Nannochloropsis oculata, was promoted by alginate oligosaccharides (38). These findings suggest that alginate oligosaccharides might act on many different
plant cells as a growth-promoting agent. Furthermore, it has been suggested that uronic acid residues in certain bioactive oligosaccharides might play an important role in the initiation of certain signal-transduction pathways in plant cells (39). Farmer et al. have proposed that uronide-Ca\(^{2+}\) complexes might be the active molecular species that initiate the signal transduction pathways leading to enhance growth and proliferation (40). Chaki et al. have suggested that sodium alginate oligosaccharides exerted antagonist activity towards calcium channels, especially voltage-operated calcium channels (41). Ascophyllan is distinguishable from fucoidan in that it contains a much higher amount of uronic acids than fucoidan. Moreover, the level of uronic acid in ascophyllan is nearly equal to fucose. Taken together with these findings, it may be possible to speculate that uronic acid residues in ascophyllan may be somewhat involved in the growth-promoting activity on MDCK cells. Further studies are required to elucidate the mechanism of action of ascophyllan on this cell line, especially in terms of structure–activity relationships.

From the viewpoint of efficient utilization of unused resources remaining in the residue of *A. nodosum* after the isolation of alginate, it is worthwhile to develop ascophyllan and fucoidan as useful bioactive polysaccharides. For such applied research, the results obtained in this study may provide valuable basic information.

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Adherent cells (MDCK, Vero, PtK1, CHO, HeLa, and XC) (100 cells/well) were incubated in 96-well plates with the indicated concentrations of ascophyllan (black columns), fucoidan (white columns), or alginate (grey columns) in α-MEM containing 10% FBS at 37°C for 5 days. The number colonies formed was determined as described in the text. The columns represent the average of triplicate measurements and the bars indicate standard deviations. Data are means ± SD (n = 3). *p < 0.01.
Figure 2. Cytotoxic Effects of Ascophyllan and Fucoidan Isolated from *A. nodosum* on Various Cell Lines as Measured by Alamar Blue Assay.

Adherent cells (MDCK, Vero, PtK1, CHO, HeLa, and XC) (2,500, 5,000, 10,000, or 20,000 cells/well) were incubated in 96-well plates with 1,000 µg/ml of ascophyllan (black columns) or fucoidan (white columns) in α-MEM containing 10% FBS at 37°C for 48 h. Cell viability was examined using the Alamar blue assay as described in the text. The values obtained without ascophyllan or fucoidan treatment were taken as control (100%). The columns represent the average of triplicate measurements and the bars indicate standard deviations. Data are means ± SD (n = 3). \*p < 0.01.
Figure 3. Growth Promoting Effect of Ascophyllan Isolated from *A. nudosum* on MDCK Cells.

(A) Adherent MDCK cells (200 cells/well) were incubated in 48-well plates with the indicated concentrations of ascophyllan in α-MEM containing 10% FBS at 37°C for 5 days. The cells were then stained with 1% methylene blue in 50% methanol as described in the text and the photographs were taken. (B) MDCK cells (200 cells/well) were incubated in 96-well plates with the indicated concentration of ascophyllan (black circles)
or fucoidan (white triangles) in α-MEM containing 10% FBS at 37°C for 5 days. The viability of the treated cells was examined using the Alamar blue assay as described in the text. Each point represents the average of triplicate measurements and the bars indicate standard deviations. (C) Adherent MDCK cells (1,000 cells/well) were incubated in 24-well plates in the presence of 1,000 μg/ml of ascophyllan (black circles), fucoidan (white triangles) or in the absence of polysaccharides (white circles) in α-MEM containing 10% FBS at 37°C for the indicated periods of time. The number of viable cells treated with ascophyllan, fucoidan or control cells cultured in the absence of polysaccharides was measured using the trypan blue dye exclusion text. Each point represents the average of triplicate measurements and the bars indicate standard deviations. *p < 0.01.