Beneficial effects of fucoidan in patients with chronic hepatitis C virus infection

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Abstract

AIM: To evaluate the effects of fucoidan, a complex sulfated polysaccharide extract from marine seaweed, on hepatitis C virus (HCV) RNA load both in vitro and in vivo.

METHODS: HCV-1b replicon-expressing cells were cultured in the presence of fucoidan obtained from Cladosiphon okamuranus Tokida cultivated in Okinawa, Japan, and quantified the level of HCV replication. In an open-label uncontrolled study, 15 patients with chronic hepatitis C, and HCV-related cirrhosis and hepatocellular carcinoma were treated with fucoidan (0.83 g/d) for 12 mo. The clinical symptoms, biochemical tests, and HCV RNA levels were assessed before, during, and after treatment.

RESULTS: Fucoidan dose-dependently inhibited the expression of HCV replicon. At 8-10 mo of treatment with fucoidan, HCV RNA levels were significantly lower relative to the baseline. The same treatment also tended to lower serum alanine aminotransferase levels, and the latter correlated with HCV RNA levels. However, the improved laboratory tests did not translate into significant clinical improvement. Fucoidan had no serious adverse effects.

CONCLUSION: Our findings suggest that fucoidan is safe and useful in the treatment of patients with HCV-related chronic liver diseases. Further controlled clinical trials are needed to confirm the present findings.

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Key words: Fucoidan; Hepatitis C virus; Replicon

INTRODUCTION

Hepatitis C virus (HCV) infection often advances to chronic hepatitis due to the low viral clearance rate, leading to liver cirrhosis (LC) and subsequent development of hepatocellular carcinoma (HCC)[1,2]. The estimated global number of people infected with HCV is 170 million and more than 3.5 million new sufferers are diagnosed annually[3]. Currently, there is no vaccine available for HCV.
for prevention of HCV infection due to the extreme sequence variability within the HCV genome. The first-line treatment for chronic hepatitis C (CHC) includes the combination of pegylated IFN-α and ribavirin, a broad spectrum antiviral drug. Although the reported HCV eradication rate by this combination therapy is 75%-90% for genotypes 1 and 4 and 45%-52% for genotypes 2 and 3, these rates are still far from ideal. Because of the high rate of nonresponders among those infected with genotype 1, the predominant strain in Japan, and because antiviral treatment causes frequent, unpleasant and sometimes serious adverse effects, the establishment of a new treatment modality without serious adverse effects is desirable.

Considering the prolonged period (20-30 years) required for development of LC and HCC in individuals infected with HCV, progression of the disease might be influenced by nutritional status and diet. Although herbal supplements, including silymarin (an extract of milk thistle), are frequently used by patients with chronic liver diseases, the available scientific evidence for the beneficial effects of these supplements is limited. However, administration of EH0202, a mixture of four herbal extracts, is reported to induce IFN activity and reduce HCV RNA levels in patients with high viral titers. Furthermore, a recent study reported the hepatoprotective effect of birch bark extract in patients with CHC.

Fucoidan is a sulfated polysaccharide extracted from marine brown seaweeds that possess some biological activities including anti-inflammatories. Sulfated polysaccharides, including fucoidan, are also reported to inhibit the replication of viruses such as herpes simplex virus, Sindbis virus, human immunodeficiency virus, parainfluenza virus type II, and dengue virus. We have also reported recently that oral administration of fucoidan for 12 mo resulted in 42.4% decrease in the proviral load in patients with human T-cell leukemia virus type I-associated neurological disease. Since fucoidan shows no toxicity or irritation in humans, it may be useful also as an anti-HCV agent.

To our knowledge, there are no data on the anti-HCV effect of fucoidan. In the present study, we examined the anti-HCV activity of fucoidan extracted from the marine alga, Cladophora okamuranus Tokida (C. okamuranus Tokida) cultivated in Okinawa, Japan. Our pilot study is the first clinical trial that investigated the effect of fucoidan in patients with HCV-related chronic liver diseases.

**MATERIALS AND METHODS**

**Preparation of fucoidan from seaweed**

The unsalted brown seaweed C. okamuranus Tokida cultivated in Okinawa, Japan, was suspended in water, 0.57% (w/v) citric acid was added to the solution, and then heated at 90 °C for 40 min. The suspension was neutralized with NaOH and cooled to 40 °C. It was centrifuged at 3500 g by decantation centrifugal separator. The supernatant was collected, filtered using Cohlo filter, and concentrated by ultrafiltration (molecular weight cutoff 6000). The extracts were dried by spray dryer. They were composed of carbohydrates (72%), uronic acids (24%), and sulfate (8%). Total carbohydrates were determined by the phenol-HSO₄ method using fucose as the standard. Uronic acids were determined by the carbazole-HSO₄ method using D-glucuronic acid as the standard. The sulfate contents were measured by ion chromatography. The main carbohydrates were fucose. Fucoidan content determined by high-performance liquid chromatography was 83% and the molecular weight was 21-kDa. Fucoidan was dissolved in phosphate-buffered saline at a concentration of 30 mg/mL.

**Inhibition assay of HCV replicon cells by fucoidan**

Fucoidan was added to Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum of HCV subgenomic replicon cells FLR3-1 (genotype 1b, Con-1) at a final concentration of 62.5, 125, 250, 500, 1000, 2000, and 3000 μg/mL. FLR3-1 cells were established from human hepatoma HuH-7 cells by stable transfection with subgenomic selectable RNA in which the encoding HCV structural proteins were replaced by the firefly luciferase gene, the internal ribosome entry site of the Encephalomyocarditis virus, and the neomycin phosphotransferase gene. After 72 h-incubation, the cells were washed in phosphate buffered saline and lysed in reporter lysis buffer (Promega, Madison, WI). Lysates were assayed for luciferase activity with the luciferase assay system (Promega) using the instructions provided by the manufacturer. With this HCV subgenome, the efficiency of subgenomic HCV expression could be estimated by measuring luciferase activity in the replicon cells.

**Measurement of cell viability**

Cell viability was measured using the cell proliferation reagent, WST-8 (Wako Pure Chemicals, Osaka, Japan). This method relies on mitochondrial dehydrogenase cleavage of WST-8 to formazan dye to estimate the level of cell viability. Briefly, FLR3-1 cells were incubated in a 96-well microculture plate. After 24 h incubation, fucoidan was added to the cells at various concentrations. After 72 h culture, WST-8 (5 μL) was added for the last 4 h of incubation and absorbance at 450 nm was measured using an automated microplate reader. WST-8 solution was added to the media-only wells to correct for background.

**Patients**

Table 1 lists the characteristics of the patients. The subjects included in the study were 15 patients with chronic liver diseases (7 men and 8 women; age: 66.1 ± 11.1 years; mean ± SD, range, 42-86), who visited the Nakasonokazu Medical Clinic. This study was carried out as an open-label study. All patients were infected with HCV genotype 1b, with a serum viral load in excess of 10⁶ copies/mL. Nine patients had been diagnosed with CHC, 4 with HCV-related LC, and 2 with HCV-related cirrhosis and HCC.
Table 1  Characteristics of patients

<table>
<thead>
<tr>
<th>Patient No.</th>
<th>Age (yr)</th>
<th>Gender</th>
<th>Diagnosis</th>
<th>Previous IFN therapy</th>
<th>Other medications</th>
</tr>
</thead>
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<tr>
<td>1</td>
<td>73</td>
<td>F</td>
<td>LC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>2</td>
<td>78</td>
<td>F</td>
<td>LC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>3</td>
<td>49</td>
<td>M</td>
<td>LC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>4</td>
<td>72</td>
<td>M</td>
<td>LC + HCC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>5</td>
<td>66</td>
<td>M</td>
<td>LC + HCC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>6</td>
<td>70</td>
<td>M</td>
<td>LC</td>
<td>Not eligible</td>
<td>None</td>
</tr>
<tr>
<td>7</td>
<td>70</td>
<td>F</td>
<td>CHC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>8</td>
<td>86</td>
<td>F</td>
<td>CHC</td>
<td>Not eligible</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>9</td>
<td>55</td>
<td>M</td>
<td>CHC</td>
<td>Intolerant</td>
<td>None</td>
</tr>
<tr>
<td>10</td>
<td>69</td>
<td>M</td>
<td>CHC</td>
<td>Non-responder</td>
<td>Glycyrrhizin</td>
</tr>
<tr>
<td>11</td>
<td>71</td>
<td>M</td>
<td>CHC</td>
<td>Non-responder</td>
<td>Glycyrrhizin</td>
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<tr>
<td>12</td>
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<td>CHC</td>
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<td>Glycyrrhizin</td>
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<tr>
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<td>62</td>
<td>F</td>
<td>CHC</td>
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<tr>
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<td>42</td>
<td>F</td>
<td>CHC</td>
<td>Non-responder</td>
<td>None</td>
</tr>
</tbody>
</table>

M: Male; F: Female; LC: Liver cirrhosis; CHC: Chronic hepatitis C; HCC: Hepatocellular carcinoma; IFN: Interferon.

Eight patients were not eligible for IFN treatment because of LC, complication (depression), or advanced age. Seven patients had received IFN therapy in the past. Six patients were non-responders to IFN and 1 discontinued therapy because of side effect (depression). All patients assessed the tolerability as excellent.

During fucoidan treatment, 11 patients received a glycyrrhizin preparation. Fucoidan (provided by Kanehide Bio Co., Okinawa, Japan) was given orally as capsules containing 166 mg of dry extract from C. okamuraensis Tokida per capsule in a dose of five capsules daily for 12 mo. Informed consent was obtained from all patients enrolled in the study, after a thorough explanation of the aims, risks, and benefits of this therapy.

Test parameters
The outcome parameters included the course of alanine aminotransferase (ALT), aspartate aminotransferase (AST), quantitative HCV RNA levels, subjective symptoms associated with CHC, LC, and HCC (such as fatigue, abdominal discomfort, depression, and dyspepsia), safety, and compliance. Data on all clinical parameters were documented at each visit. HCV RNA levels were determined using the AMPLICOR GT HCV Monitor test (Roche Diagnostics, Basel, Switzerland), which has a lower limit of quantitation of 0.5 kIU/mL at a linear range up to 850 kIU/mL.

Statistical analysis
Data are expressed as mean ± SD. The results of biochemical tests and HCV RNA levels were compared by the Student’s t test. A P < 0.05 was considered significant.

RESULTS
Fucoidan suppresses HCV replication
To assess the effects of fucoidan on intracellular replica-
We also measured serum IFNα levels to determine the indirect effect of fucoidan on IFNs, especially whether it increases the antiviral activity of IFNs. However, IFNα could not be detected in the serum of patients treated with fucoidan. Furthermore, fucoidan did not enhance IFNs expression in FLR3-1 replicon cells (data not shown). It has been reported that the protective effect of fucoidan is based on direct inhibition of viral replication and stimulation of both innate and adaptive immune defense functions. We are currently investigating the effect of fucoidan on the host immune system including natural killer cell cytotoxic activity.

Our study has certain limitations. First, the study comprised only a small number of patients, including 6 patients who were known non-responders to IFN therapy. Second, all patients harbored HCV virus genotype I b and 6 had cirrhosis. Thus, at least some patients in this cohort could be classified as likely non-responders to IFN therapy. Thus, the selection criteria employed in the present study may have favored a poor response to fucoidan.

The abnormally high levels of ALT tended to decrease temporarily during fucoidan treatment, suggesting a correlation between viral load and indices of hepatic dysfunction. Thus, fucoidan may be effective in the management of HCV-related chronic liver diseases, although long-term clinical improvement was not observed in the present study. Importantly, no adverse events were observed in all patients, similar to the results reported in a previously study on fucoidan, suggesting that daily oral administration of fucoidan for 12 mo is safe and tolerable.

There is no doubt that patients who fail to respond to conventional treatments often seek alternative therapies. In conclusion, our study demonstrated that fucoidan from C. okamuranus Tokida has HCV replication suppressive effects in a replicon cell system. Furthermore, our relatively small uncontrolled pilot study showed that fucoidan has temporary but beneficial effects on HCV RNA levels in HCV infected patients. The preliminary findings suggest that fucoidan may be a useful health-food additive with antiviral activity to be used in the treatment of chronic liver diseases. To suppress the viral titer as much and for as long as possible, we need to define the daily effective dosage. Further studies on the mechanism of fucoidan-induced HCV inhibition may provide alternative strategies for the design of novel anti-HCV drugs.

ACKNOWLEDGMENTS

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COMMENTS

Background

Hepatitis C virus (HCV) is a major cause of chronic liver diseases including chronic hepatitis, cirrhosis, and hepatocellular carcinoma. The standard care...
for chronic hepatitis C involves the administration of pegylated α-interferon in combination with the nucleotide analog ribavirin. However, this regimen has limited success rate for genotype 1 and 4, and unfavorable side effects. Thus, it is important to discover more effective and safer agents to improve the clinical treatment on HCV carriers. Fucoidan, a sulfated polysaccharide, has significant biological activities, such as antiviral and anti-inflammatory effects. Nevertheless, there has been no investigation on the efficacy of fucoidan against HCV infection.

Research frontiers

Natural products have been used for the treatment of various diseases as an alternative to conventional chemical agents. So far, several natural products have been screened for their antiviral effect against various viral infections. Screening of natural potent inhibitors for HCV has also become a research hotspot.

Innovations and breakthroughs

Previous studies have shown the efficacy of natural products against HCV replication in a cell-based HCV replicon system. However, there have been few clinical studies that evaluated the safety and efficacy of these natural products. In the present study, the authors investigated the anti-HCV activity of fucoidan obtained from the Cladosporium okamuranus Todoki cultivated in Okinawa, Japan, both in vitro and in vivo. This pilot study is the first clinical trial that investigated the effect of fucoidan in patients with HCV-related chronic liver diseases.

Applications

Fucoidan inhibited HCV RNA replication in the HCV replicon assay system. The experimental data on fucoidan efficiency in cell culture stimulated the rationale for clinical study. Oral fucoidan administration resulted in temporary reduction of viral loads of genotype 1b in patients with chronic HCV infection, who were not eligible for, did not respond to, or were intolerant of interferon therapy. Fucoidan is well tolerated and no serious adverse events were observed in any of the patients. Fucoidan exhibited antiviral properties against HCV both in vitro and in vivo, and would be expected to become a new strategy for HCV infection. Further controlled clinical trials will be required to confirm the present findings.

Terminology

Fucoidan is a complex sulfated polysaccharide found in the cell walls of several edible brown algae, including Fucus vesiculosus. The HCV replicon system replicates a modified HCV genome containing luciferase gene to high levels in human hepatoma cells. The efficacy of subgenomic HCV expression was estimated by measuring luciferase activity in the replicon cells. This system provides a powerful tool for studying virus replication and for screening anti-HCV drugs.

Peer review

The paper studied the effects of fucoidan, a complex sulfated polysaccharide extracted from marine seaweed, on HCV RNA load in vitro and in vivo. The research is of significance because of the high rate of nonresponders in HCV genotype 1b, which is the predominant strain in Japan. Moreover, antiviral treatment causes frequent, unpleasant and sometimes serious adverse effects, thus the search for a new treatment modality without serious adverse effects is desirable.

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