A Protocol for Human Serum Fucoxanthinol Quantitation using LC-MS/MS System

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Abstract

Fucoxanthin, a marine carotenoid widely distributed in brown algae, is known to possess various health promoting activities including anti-obesity effects in humans, but little is known about its kinetics partly because of its low bioavailability. We have developed a simple and reproducible protocol for quantifying system of human serum fucoxanthinol, a fucoxanthin metabolite, used LC-MS/MS multiple reaction monitoring (MRM), and have applied for clinical trial to study fucoxanthin absorption. We established a MRM channel for fucoxanthinol (m/z 617.5 for Q1 and 109.0 for Q3 channel) with a standard curve of \( R^2 > 0.99 \) and an optimized pretreatment of human serum samples. Fucoxanthinol levels in sera obtained from subjects participating in clinical trials were then measured and a significant increase in the levels was observed at 4 hr after a single oral dose of 22 mg fucoxanthin using newly developed protocol. We propose that use of this protocol could enhance clinical investigation involving the bioavailability and functionality of fucoxanthinol.

Keywords

Fucoxanthin, Fucoxanthinol, Fucoxanthin intake, Human serum, LC-MS/MS, MRM, Clinical trial

Abbreviations

APCI: Atmospheric pressure chemical ionization; Fx: Fucoxanthin; FxOH: Fucoxanthinol; HPLC: High-performance liquid chromatography; LC: Liquid chromatography; LC-MS: Liquid chromatography-mass spectrometry; LC-MS/MS: Liquid chromatography-tandem mass spectrometry; LLOQ: The lower limit of quantification; LOD: The limit of detection; MCT: Medium-chain triglyceride; MRM: Multiple reaction monitoring; MS: Mass spectrometry; m/z: Mass-to-charge ratio; Q1: The first quadrupole; Q3: The third quadrupole.

Introduction

Fucoxanthin is a marine carotenoid widely found in edible brown algae such as wakame (Undaria pinnatifida), kombu (Laminaria japonica) and akamoku (Sargassum horneri), and contributes more than 10% of the estimated total production of carotenoids in nature [1]. Fucoxanthin exhibits numerous health benefits especially anti-cancer [2-5] and anti-obesity [6,7] activities as well as benefitting various other health related problems [8-10]. According to Abidov, et al. in a report, which focuses on the anti-obesity activity of fucoxanthin, oral administration of a dietary supplement capsule containing fucoxanthin induces weight loss and increases resting energy expenditure in obese women [11]. However, only the activities of fucoxanthin were assayed and the kinetics of possible metabolites were unidentified in the report. To determine the physiological relevance of these activities, it is essential not only to study the digestion and absorption of orally administered fucoxanthin but also to identify its metabolites and the effective relationship between fucoxanthin and its possible activity.

Dietary fucoxanthin is incorporated into the blood as fucoxanthinol after deacetylation in the digestive tract [12] (Figure 1), and at least in the liver of mice, is then metabolized to amoracoxanthin A [13]. Several detection systems have been described previously for analyzing fucoxanthin in vivo, including HPLC [14,15], LC-MS [16,17] and LC-MS/MS [18]. Low bioavailability of fucoxanthinol compared to other carotenoids in humans has been demonstrated [14,15]. Asai, et al. reported that serum fucoxanthinol concentration after wakame intake (including 6.1 mg of fucoxanthin) was close to the lower limit of quantification by HPLC [14]. Therefore, to perform valid kinetics studies a more sensitive quantifying system for serum fucoxanthinol than exists at present is required. In addition to this approach, since the presence of highly concentrated proteins and other components in serum samples complicates analyses (e.g. causing background contamination by impurities), it is needed to remove them by improving preparation and analytical protocol to determine serum fucoxanthinol properly.

We aimed to develop a protocol for quantification of serum fucoxanthinol using liquid chromatography coupled with tandem mass spectrometry (MS/MS), which has a high sensitivity and a wide dynamic range. To apply the developed protocol to a large number of serum samples obtained from clinical trials, we also examined a washing step of LC for continuous analyses and evaluated the effects of pre-analytical factors, such as a freeze-thaw cycle on fucoxanthinol stability. We here report the kinetics in humans after a single dose of fucoxanthin to confirm the validity of applying the developed analysis to various clinical researches involving fucoxanthin intervention.

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Moreover, precision and accuracy of intra- and inter-day were assessed using calibration standards of 10, 100 and 1000 ng/mL (50, 500, 5000 pg on the column) fucoxanthinol and fucoxanthin. The precision and accuracy were determined by replicated analysis of 3 samples of each concentration of standard on 8 different days. Intra-day analysis was performed twice a day (i.e. morning and afternoon).

**Extraction of fucoxanthinol from serum samples**

To quantify fucoxanthinol levels in serum samples, the serum epoxxyanthophyll fraction that included fucoxanthinol was extracted according to a method of Asai, et al. [14] with a slight modification. In brief, 1 mL serum, 0.2 mL saline, and 2 mL methanol including 5 ng fucoxanthin as an internal standard were added to glass tubes and vortexed. After adding 4 mL dichloromethane, the mixture was centrifuged at 416 g for 10 min at room temperature. The bottom layer was collected and this extraction procedure was repeated twice. The solvent of the collected fraction was removed to dry using a centrifugal concentrator (VC-96W, Taitec, Saitama, Japan) equipped with a vacuum pump and freeze trap. The residue was dissolved in n-hexane/diethyl ether (9/1, v/v), and applied to a Bond Elut ALN (100 mg, 1 mL) solid-phase extraction cartridge (Agilent Technologies, Santa Clara, CA, USA) pretreated with 1 mL n-hexane. After the cartridge was washed with the 1 mL n-hexane/diethyl ether (9/1, v/v), the epoxxyanthophylls were eluted with 1 mL diethyl ether/ethanol (4/1, v/v). The eluate was dried in vacuo, re-dissolved in methanol/acetonitrile (70/30, v/v).

After these procedures described by Asai, et al., we added to ultrafiltrate with Amicon Ultra-0.5 mL 3K devices (Merck Millipore, Billerica, MA, USA) by twice centrifugation at 14,000 g for 10 min at room temperature, due to remove proteins carried over into collected epoxxyanthophyll fraction during extraction.

The collected sample was dried and resuspended in 250 µL of methanol/acetonitrile (70/30, v/v), and then the 50 µL sample was subjected to LC-MS/MS analysis. Triplicates of each serum sample were independently extracted and analyzed.

**Quantification of fucoxanthinol in human sera**

Fucoxanthin was used as the internal standard and spiked in serum to quantify the fucoxanthinol level because its structure and solubility are similar to fucoxanthin. A previous report [15] and our preliminary experiment (Supplementary Figure 1) showed that fucoxanthin was not detected in serum after its intake. Samples were next analyzed in tuned MRM channels of both fucoxanthin and fucoxanthinol or 50 µL of serum sample). The flow rate was set for 1 mL/min at 35°C.

The LC system was coupled with triple quadruple MS/MS system (AB SCIEX, Framingham, MA, USA), which was equipped with an atmospheric pressure chemical ionization (APCI) source, and operated in positive ion mode. The MS parameters optimized were as follows: declustering potential: 76 V; entrance potential: 10 V; dwell time: 400 msec; curtain gas (nitrogen): 10 psi; ion source gas 1 (nitrogen): 80 psi; turbo gas temperature: 500°C; interface heater: on; and nebulizer current: 5.0 psi. Nitrogen was used as the collision gas with a collision energy of 52 eV and a collision cell exit potential of 7 V. Analyst software 1.5 (AB SCIEX, Framingham, MA, USA) was used for the system control, data acquisition and data processing.

**Standard curve of fucoxanthinol**

Calibration standards for fucoxanthinol were prepared at 1, 10, 100, 1000, and 10000 ng/mL by successive dilution with methanol/acetonitrile (70/30, v/v). Five microliter of each standard was subjected to LC-MS/MS as described above and the standard curve was constructed by plotting the peak area ratio of fucoxanthinol (y) versus the concentration (ng/mL) of fucoxanthinol (x).

Moreover, precision and accuracy of intra- and inter-day were assessed using calibration standards of 10, 100 and 1000 ng/mL (50, 500, 5000 pg on the column) fucoxanthinol and fucoxanthin. The precision and accuracy were determined by replicated analysis of 3 samples of each concentration of standard on 8 different days. Intra-day analysis was performed twice a day (i.e. morning and afternoon).

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for each day, samples were thawed at room temperature and extracted. The 50 μL of extracted sample was subjected to LC-MS/MS analysis. Triplicates of each concentration on freeze-thaw cycle were extracted and analyzed. The impact of freeze-thaw was analyzed by comparing it to fucoxanthin. The 50 μL of extracted sample was subjected to LC-MS/MS analysis for each day, samples were thawed at room temperature and extracted.

**Results and Discussion**

**MRM channel and standard curve of fucoxanthinol**

To obtain greater sensitivity for quantifying serum fucoxanthinol, we employed multiple reaction monitoring (MRM) [20]. Because the most intense signals for fucoxanthinol and fucoxanthin were observed in the first quadrupole (Q1) and the third quadrupole (Q3) masses (Table 1) during continuous infusion of standard solution of each compound at 1 μg/mL, we selected the channel for fucoxanthinol (m/z 617.5, 109.0) and for fucoxanthin (m/z 659.4, 109.0) as quantitation ions (Table 1), respectively. We confirmed standard fucoxanthol was detected in m/z 617.5 → 109.0 at RT 2.3 min (Figure 2A). The channel was linear in the concentration range tested of 10-10000 ng/mL (50-50000 pg on the column) with a correlation coefficient of 0.9989 (Figure 2B).

**Precision and accuracy of standard analyses**

Table 2 shows the results for precision and accuracy analyzed twice in one day or on 8 different days. The intra- and inter-day precisions in fucoxanthinol and fucoxanthin were shown in table 2. Generally, it is recommended that precision should be < 20% and accuracy between 80-120% of the theoretical value. However, in the analysis of 10 ng/mL (50 pg on the column) of fucoxanthinol, both precision and accuracy were decreased in both intra- and inter-day analysis. On the other hand, for the concentrations of 100 and 1000 ng/mL (500 and 5000 pg on the column) of fucoxanthol, the precision was < 17% and accuracy was 99.4-115.3%. Moreover, in all the concentrations of fucoxanthol, the precision was < 7% and the accuracy was 99.4-115.3%. The final washing and reconditioning steps were shown in Table 2. Twice in one day or on 8 different days. The intra- and inter-day precisions in fucoxanthol and fucoxanthin were shown in Table 2. Generally, it is recommended that precision should be < 20% and accuracy between 80-120% of the theoretical value. However, in the analysis of 10 ng/mL (50 pg on the column) of fucoxanthol, both precision and accuracy were decreased in both intra- and inter-day analysis. On the other hand, for the concentrations of 100 and 1000 ng/mL (500 and 5000 pg on the column) of fucoxanthol, the precision was < 17% and accuracy was 99.4-115.3%. Moreover, in all the concentrations of fucoxanthol, the precision was < 7% and the accuracy was 99.4-115.3%.

**Examination of LC column washing step for continuous analyses of serum samples**

To remove impurities from serum and shorten the total run time, we examined the effects of washing with dichloromethane and a reconditioning step from RT 5 min to 60 min. As shown in figure 3, peaks detected in MRM channels of fucoxanthinol (blue line; m/z 617.5, 109.0, red line; m/z 617.5, 67.0) were observed at least until 320 min under no washing step conditions. This was dramatically improved by washing with dichloromethane from 5 to 50 min. Figure 3B shows no peaks after 52 min. We thus successfully removed impurities and reduced total run time to less than 60 min. The final washing and reconditioning steps were as follows; solvent A (methanol/acetonitrile, 70/30, v/v) and solvent B (dichloromethane);
Validation of fucoxanthinol and fucoxanthin in serum sample was shown in Table 3. The mean CV of peak area of spiked fucoxanthinol and fucoxanthin were calculated as 9.2% and 19.2%, respectively. It indicated that this protocol kept good precision throughout serum sample preparations and analyses. In the recovery of fucoxanthinol and fucoxanthin, they were over 100% (Table 3). It may be due to the effect of serum contents. The LOD and LLOQ was fucoxanthinol and fucoxanthin from serum sample were 71.1 pg and 234.6 pg (in fucoxanthinol, respectively), and 36.2 pg and 119.5 pg (in fucoxanthin, respectively).

Increase of serum fucoxanthinol levels after fucoxanthin intake

After a single dose of 22 mg fucoxanthin, its metabolite, fucoxanthinol, was measured in human sera at 0, 4, 24 and 48h. No peak representing fucoxanthinol was detected in the MRM chromatogram at 0 h, while a rapid increase was observed at 4 h. Peaks declined at 24 to 48h (Figure 5A). Fucoxanthinol was quantified using fucoxanthin as an internal control. Serum fucoxanthinol levels were markedly elevated at 4 h in all subjects and were gradually decreasing in 24 h to 48 h (Figure 5B). However, the levels did not return to the baseline of 0 h (0 h, 0.4 ± 0.2 nM; 4 h, 44.2 ± 14.9 nM; 24 h, 6.2 ± 0.9 nM; and 48h, 1.6 ± 0.4 nM), suggesting that orally administered fucoxanthin is absorbed as fucoxanthinol within 4h and is not metabolized completely even after 48h.

Precision of serum sample analyses

Using the developed protocol, we analyzed clinical sample. The validation of fucoxanthinol and fucoxanthin in serum sample was shown in Table 3. The mean CV of peak area of spiked fucoxanthinol and fucoxanthin were calculated as 9.2% and 19.2%, respectively. It indicated that this protocol kept good precision throughout serum sample preparations and analyses. In the recovery of fucoxanthinol and fucoxanthin, they were over 100% (Table 3). It may be due to the effect of serum contents. The LOD and LLOQ was fucoxanthinol and fucoxanthin from serum sample were 71.1 pg and 234.6 pg (in fucoxanthinol, respectively), and 36.2 pg and 119.5 pg (in fucoxanthin, respectively).

Table 3: The validation of fucoxanthinol and fucoxanthin in serum sample analyses.

<table>
<thead>
<tr>
<th></th>
<th>Fucoxanthinol</th>
<th>Fucoxanthin</th>
</tr>
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<tbody>
<tr>
<td>Precision (%CV)</td>
<td>9.2</td>
<td>19.2</td>
</tr>
<tr>
<td>Recovery (%)</td>
<td>136.6</td>
<td>117.3</td>
</tr>
<tr>
<td>LOD (pg on column)</td>
<td>71.1</td>
<td>36.2</td>
</tr>
<tr>
<td>LLOQ (pg on column)</td>
<td>234.6</td>
<td>119.5</td>
</tr>
<tr>
<td>LOD as serum sample*</td>
<td>0.58 nM</td>
<td>0.27 nM</td>
</tr>
<tr>
<td>LLOQ as serum sample*</td>
<td>1.91 nM</td>
<td>0.91 nM</td>
</tr>
</tbody>
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*Following the developed protocol in current study.

0-5 min with 0% B; 5-25 min with 0-100% B with a linear gradient of B; 25-50 min with 100% B and 50-60 min of 0% B. By adding the washing steps, we have developed a protocol (Figure 4) and achieved continuous analyses of serum fucoxanthinol in this study without any trouble such as a clogging of LC column. Although Zhang, et al. have reported the development of the system quantifying plasma fucoxanthinol in rats by LC-MS/MS, they did not mention washing conditions of LC column in continuous sample analyses [18]. Therefore, our examined washing protocol would provide the useful information for reasonable sample analyses.
Impact of freeze-thaw cycles on fucoxanthinol stability in human sera

To prepare for handling freeze-thawed blood samples in clinical research, we examined the impact of freeze-thaw cycles by spiking standard fucoxanthinol. The 0 cycle of adjusted recovery of fucoxanthinol was expressed as 100%. As shown in Figure 6, in serum samples of 5000 pg, the recovery of fucoxanthin was significantly lower than 100%. However, the recovery of fucoxanthinol remained at 100% even after 5 freeze-thaw cycles. This indicates that fucoxanthinol is more stable than fucoxanthin during freeze-thaw cycles.

Figure 4: A protocol for human serum fucoxanthinol quantitation.
decreased by freeze-thaw. Moreover, it seemed that 1 or 2 freeze-thaw cycles tended to decrease in each concentration, although a significant difference was not observed due to the wide SD of the 0 cycle sample. Previous reports indicate that retinol, alpha-tocopherol, trans-lycopene, and trans-beta-carotene in reconstituted lyophilized serum stored at -20°C were stable for at least 3 days with minimal (<5) freeze/thaw cycles [21], however, fucoxanthinol in serum samples was less stable during freeze-thawing in our study. These results showed that repeated cycles of freeze-thaw should be avoided for fucoxanthinol quantification. It is impractical in clinical experiments to keep collected blood samples fresh until analysis. Unified freeze-thaw cycles would be important for analysis of a series of clinical samples.

**Conclusions**

We have developed a quantification protocol for fucoxanthinol in human sera as detected by LC-MS/MS and observed a change of fucoxanthinol levels after a single dose fucoxanthin. In this protocol, we have achieved continuous analyses of serum fucoxanthinol in this study without any trouble. These results may be applied to clinical investigations to elucidate the biological activity of fucoxanthin in humans.

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Ethical Statement

This study was conducted according to the guidelines of the Declaration of Helsinki, and our study protocol and all procedures involving human subjects were approved by the ethical review committee of Sapporo Medical University (#24-2-91).

Competing Interests

The authors have no conflict of interests.

Author’s Contribution

NM designed the research and performed experiments, analyzed and interpreted the data and wrote the manuscript. MH and MA designed the research, especially LC-MS/MS analysis, interpreted the data and revised the manuscript. KM formulated the study conception, designed the research, especially LC-MS/MS analysis, interpreted the data and wrote the manuscript. YK supervised the work, designed the research, interpreted the data and wrote the manuscript. HS interpreted the data and revised the manuscript. HS interpreted the data and revised the manuscript. All authors read and approved the final manuscript.

References

Supplementary Figure 1: Fucoxanthin was not detected in serum after its administration. After 4 hours of 22 mg of fucoxanthin intake, we analyzed fucoxanthin and fucoxanthinol in human serum by LC-MS/MS system. The peak of fucoxanthinol was detected at 2.3 min, that of fucoxanthin was not at 2.7 min. (A) Mixed solution of fucoxanthin and fucoxanthinol standard; (B) Serum sample after 4 hours of fucoxanthin administration.