Quantitative Estimation of Interaction between Carbohydrates and Concanavalin A by Surface Plasmon Resonance Biosensor

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We measured the affinity of more than 20 sugars with concanavalin A (ConA) by an optical biosensor (surface plasmon resonance sensor) using asialofetuin (ASF) as an immobilized binding partner of ConA. We determined kinetic parameters of the effects of sugars on the dissociation of ConA from ASF quantitatively, and the structural requirements of the functional groups of sugars for binding with ConA. We found that the affinity of ConA for sugars is dependent on its conformation induced by interaction with the binding partner. In addition, the results showed that optical biosensor system is well mimics the interaction of ConA with sugars in biomembrane.

Key words concanavalin A; asialo-fetuin; sugar interaction; optical biosensor

Concanavalin A (ConA) isolated from jack bean (Canavalia ensiformis) belongs to the legume lectin family.1) Among the monosaccharides, methyl α-D-mannopyranoside (Me α-Man, see Chart) exhibits the most potent inhibitory effect on formation of the complex of ConA with dextran,2–4) and the mannotriose 3,6-di-O-(α-D-mannosyl)-α-D-mannoside was found to be the most potent oligosaccharide, being more potent than Me α-Man.5)

Shinohara et al.5,7) examined the binding of ConA with glycosidase-treated fetuins containing the branched mannotriose moieties, which were immobilized in the chip of an optical biosensor utilizing the surface plasmon resonance (SPR) technique.6–9) Their results showed that the terminal mannotriose moiety of glucosaminidase-treated fetuin (glucosaminofetuin) was most sensitive. Galactosidase-treated fetuin (agalactosylfetuin) and sialidase-treated fetuin (asialofetuin, ASF) having glycosylated non-terminal mannotriose moieties bound more weakly than glucosaminofetuin to ConA.6)

To understand the structural requirements of sugars for binding with ConA in biomembranes, we quantitatively analyzed the effects of various sugars on the dissociation of ConA from ASF using an optical biosensor system. As ASF does not interact with ConA tightly,3) the ASF–ConA system should be favorable for determination of the affinity of sugars for ConA. In fact, we found that our system is a good mode of interaction of sugars with ConA in biomembranes.

Experimental

Materials Human and bovine fresh blood was obtained from the Blood Center of the Japan Red Cross Society and the Meat Inspection Facility, Tokushima, respectively. ConA and ASF were purchased from Sigma. Other chemicals and reagents were of the highest grade commercially available.

Preparation of the ASF-Immobile Dextran Matrix on the Biosensor Chip SPR measurements were performed with a BIACORE 1000 (Pharmacia Biosensor AB, Uppsala, Sweden). We used a CMS biosensor chip coated with carboxymethyl/dextran (CMD) matrix, and it was mounted in a flow-cell. The flow-cell was equilibrated with Hepes buffered saline (HBS) containing 90 mM NaCl and 20 mM Hepes/NaOH buffer (pH 7.0) at 25 °C, and CMD was activated with a mixture of 200 mM N-ethyl-N’-(diethylaminoethyl)-carbodiimide and 50 mM N-hydroxysuccinimide. Into the activated flow-cell 30 mg/ml ASF in 10 mM sodium acetate buffer (pH 4.7) was injected. Then, free activated carboxyl groups were masked by injection of 1 M ethanolamine at pH 8.5. Each sample was injected into the ASF-immobilized flow-cell. The net amount of the conjugated ASF was calculated by subtracting its amount in the blank flow-cell from that of the ASF-immobilized flow-cell, in terms of SPR response. The immobilization of ASF to a CMD matrix in the flow-cell increased SPR by 2300 units, which may correspond to 2.5 ng/mm² ASF. The ASF-immobilized biosensor chip was stored at 4 °C in the chemically stable package.

Inhibition of Sugars on the Binding of Con A to Immobilized ASF A solution of 50 μg/ml ConA in HBS was injected into the ASF-immobilized flow-cell, and the flow-cell was washed with HBS for 15 min. Then the test compound solution was applied for 15 min. The interaction was monitored at 25 °C as the change in the SPR response (R). The same flow-cell was used repeatedly by washing it with 150 mM HCl.

The apparent dissociation rate constant k₇ for the ConA-ASF complex was experimentally obtained by Eq. 1, where R₀ and Rₜ are the SPR responses at time t and at the time of injection of the test compound q₀, respectively.6–10

\[
\ln \frac{R_0}{R} = k_7 (t - t_0)
\]

(1)

The χ²-test was used to assess the validity of the obtained k₇ value, using the operating program of the BIACORE 1000 system. ConA dissociated from the immobilized ASF at the surface of the sensor chip diffuses from the unstirred surface (stationary phase, ConAₛ) to the mobile phase (ConAₐ). These processes and the kinetics of dissociation of ConAₛ from ASF are shown by Eqs. 2 and 3, respectively.

\[
\text{ConA} \rightleftharpoons \text{ConA}_\text{s} \rightarrow \text{ConA}_\text{a} \quad k_5
\]

(2)

\[
\frac{dR}{dt} = k_5 [\text{ConA}_\text{s}] (R_0 - R) - k_7 R
\]

(3)

Under steady-state conditions, k₅ is given by Eq. 4, where k₅, k₇, and k₉ represent intrinsic association and dissociation rate constants of ASF–ConA complex and mass-transport rate constant of ConA from the stationary phase to mobile phase, respectively.11,12)

\[
k_5 = \frac{k_7}{1 + k_7 (R_0 - R)}
\]

(4)

The association constant Kₛ of the competitive inhibitor S with ConA is represented by Eq. 5.

\[
K_s = \frac{[\text{ConA} - S]}{[\text{ConA}][S]}
\]

(5)

Then, the relative concentration of free ConA α is shown by Eq. 6.

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When the binding of S to ConA proceeds immediately, Eq. 7 is obtained.

$$\frac{dR}{dt} = k_1 [\text{ConA}] (R_0 - R) - k_2 R$$

From Eqs. 3, 4 and 7, $k_1$ is expressed by Eq. 8.

$$k_1 = \frac{k_2}{1 + \frac{k_2 \alpha (R_0 - R)}{K_2}}$$

Then, Eq. 9 is obtained.

$$\frac{1}{k_d} = \frac{1}{k_1} + \frac{1}{k_0} - \frac{1}{K_2[S] + 1}$$

When $\nu = 1/k_0$ and $\kappa = K_2/k_1 (R_0 - R)$, Eq. 10 is obtained from Eq. 9 under the conditions of $[S] > 1/K_2$, showing that there is a linear relationship between $1/k_d$ and $1/[S]$.

$$\frac{1}{k_d} = \nu + \frac{1}{\kappa}$$

Values of $\kappa$ and $\nu$ are related to the association constant of S with ConA and the reciprocal of the intrinsic dissociation rate constant of ConA–ASF complex, respectively, and they are obtained by regression analysis computed with the software Q SAR developed by us using Fortran 77.13–15

**Erythrocyte Aggregation Test** Human and bovine blood was washed three times with phosphate buffered saline (PBS) containing 150 mM NaCl and 10 mM sodium phosphate buffer (pH 7.4). The washed erythrocyte suspension in PBS was prepared at 2% (v/v) of the packed cells. To the wells of a microtiter plate 50 μl of 2% erythrocyte suspension and subsequently 100 μl of test compound solution in PBS were added. The plate was kept at room temperature for 2 h, and the degree of aggregation was scored.

**Results and Discussion**

**Effects of Sugars on the ASF–ConA Complex** Prior to conducting optical biosensor measurement, we confirmed the concentration dependence of ConA on its binding with ASF, which was covalently immobilized on the dextran bed in the optical biosensor chip. The immobilized ASF showed an SPR response of 2,300 units. ConA was added to the ASF-immobilized flow-cell, and injections of 50, 100, 200, and 300 μg/ml of ConA induced the SPR responses of 2000, 4200, 8500, and 13000 units, respectively. In the blank flow-cell without immobilized ASF, no increase of the SPR response was detected. Therefore, formation of the ASF–ConA complex in the flow-cell was nearly proportional to the concentration of the injected ConA. The following experiments were performed with 50 μg/ml ConA.

As shown in Fig. 1, the injection of ConA induced a hyperbolic increase in the SPR response, showing that the association between ConA and the immobilized ASF was saturating. Subsequent injection of HBS buffer caused dissociation of ConA from the immobilized ASF. Injection of d-mannose (Man, see Chart) caused dissociation of ConA. Apparent rate constant $k_d$ values for the dissociation of ConA induced by added sugars were measured and analyzed by Eq. 1.

Increases in the $k_d$ value induced by methyl $\alpha$-D-mannopyranoside (Me $\alpha$-Man, see Chart 1) are shown in Fig. 2. A similar curve was obtained for methyl $\alpha$-D-glucopyranoside (Me $\alpha$-Glc, see Chart 1). The observed dissociation of ConA from ASF was saturated at high concentrations of these sug-
The sugar to ConA (k and residual variances, respectively.

tion coefficient, standard deviation, and ratio of regression

ation of ConA–ASF complex (k) were determined to be 32.9 M

A.

Table 1. Kinetic Parameters for Sugar Derivatives

<table>
<thead>
<tr>
<th>Compound</th>
<th>ν (s)</th>
<th>k (s⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Me α-Man</td>
<td>135.0</td>
<td>32.9</td>
</tr>
<tr>
<td>Me α-Glc</td>
<td>141.3</td>
<td>14.0</td>
</tr>
<tr>
<td>Me β-Man</td>
<td>396.1</td>
<td>9.02</td>
</tr>
<tr>
<td>Me β-Glc</td>
<td>359.4</td>
<td>3.39</td>
</tr>
<tr>
<td>Man</td>
<td>123.6</td>
<td>6.02</td>
</tr>
<tr>
<td>Glc</td>
<td>207.0</td>
<td>3.74</td>
</tr>
<tr>
<td>2DOG</td>
<td>187.4</td>
<td>4.79</td>
</tr>
<tr>
<td>GlcNAc</td>
<td>187.8</td>
<td>3.32</td>
</tr>
<tr>
<td>ManNAc</td>
<td>585.0</td>
<td>3.40</td>
</tr>
</tbody>
</table>

ars. The double reciprocal plot between the measured k_d and the sugar concentration is shown in Fig. 3, and the affinity parameter k for each sugar was determined by Eq. 10. The results for Me α-Man and Me α-Glc are shown by Eqs. 11 and 12, respectively,

\[
1/k_d = 135.0 + 4.097/[Me \alpha-Man]
\]

where values in parentheses represent 95% confidence intervals, and n, r, s, and F are the number of data points, correlation coefficient, standard deviation, and ratio of regression and residual variances, respectively.

From Eqs. 11 and 12, parameters related to association of the sugar to ConA (k) and reciprocal of the intrinsic dissociation of ConA–ASF complex (ν) were determined. Table 1 lists these values for Me α-Man and Me α-Glc along with those of other sugars studied in this work described below. The value of ν for Me α-Man well agreed with that for Me α-Glc. The relative affinities k(Me α-Man) and k(Me α-Glc) were determined to be 32.9 s⁻¹ and 14.0 s⁻¹, respectively, suggesting that the affinity of Me α-Man for ConA is more than twice that of Me α-Glc.

The selectivity of Me α-Man to Me α-Glc can be expressed by the affinity ratio of these sugars, i.e., k(Me α-Man)/k(Me α-Glc)=2.35. The values of selectivity have been reported in terms of the ratio of IC₅₀ values (the sugar concentration for induction of 50% dissociation of ConA from dextran or mannosyl-lipids) by various methods. Mann and his coworkers reported the ratio 3.2 in their study on the effects of these sugars on the association of ConA with the immobilized mannosyl-lipids in an optical biosensor. Goldstein and his colleagues obtained values of 5.7 and 4.5 from their inhibitory effects on dextran precipitation induced by ConA. These results suggest that the selectivity of Me α-Man to ConA in its complex with the glucose moiety of dextran is greater than that in the complex of ConA with the mannoside moiety of mannosyl-lipids. In contrast, the lower selectivity of Me α-Man to Me α-Glc of our present results could be due to the fact that the selectivity of ConA to Me α-Man is decreased when ConA is bound to the mannotriose moiety of ASF.

Affinities of D-Mannose and D-Glucose to ConA

Next, we examined the effects of methyl β-D-mannopyranoside (Me β-Man, see Chart 1) and methyl β-D-glucopyranoside (Me β-Glc, see Chart 1) on dissociation of ASF–ConA complex. The dependences of 1/k_d on 1/[Me β-Man] and 1/[Me β-Glc] are shown in Fig. 3 and their relationships are expressed by Eqs. 13 and 14, respectively.

\[
1/k_d = 396.1 + 43.94/[Me \beta-Man]
\]

1/k_d = 359.4 + 105.6/[Me β-Glc]

The values of ν and k thus determined are shown in Table 1. We obtained the value of 3.66 for the selectivity of Me α-Man to Me β-Man (k(Me α-Man)/k(Me β-Man)) from Eqs. 11 and 13, and 4.13 for that of Me α-Glc to Me β-Glc (k(Me α-Glc)/k(Me β-Glc)) from Eqs. 12 and 14, showing that the affinity of α-configuration of the 1-OMe is about 4-times greater than the β-configuration for both methylated-Man and methylated-Glc. Although higher affinity of the α-configuration than that of the β-configuration is known, we determined the selectivity quantitatively. In addition, k(Me β-Man)/k(Me β-Glc) was determined to be 2.65 from Eqs. 13 and 14.

As the values k(Me α-Man)/k(Me α-Glc) and k(Me β-Man)/k(Me β-Glc) were very similar (2.35 and 2.65, respectively), the affinity of the configuration of the 2-OMe is more than twice that of 2α-OMe, irrespective of the configuration of 1-OMe (1α-OMe for Me α-Man and Me α-Glc, and 1β-OMe for Me β-Man and Me β-Glc).

Next, we examined the effects of Man and D-glucose (Glc, see Chart 1) on the ASF–ConA complex. For this, we allowed these solutions to stand for 2 d to attain equilibrium, and measured SPR responses. We obtained Eqs. 15 and 16.

\[
1/k_d = 123.6 + 20.62/[Man]
\]

\[
1/k_d = 207.0 + 55.33/[Glc]
\]
smaller than the values of \( \chi(\text{Me} \alpha-\text{Man})/\chi(\text{Me} \alpha-\text{Glc}) = 2.35 \) and \( \chi(\text{Me} \beta-\text{Man})/\chi(\text{Me} \beta-\text{Glc}) = 2.65 \). The equilibrium concentrations of \( \alpha- \) and \( \beta- \)-mannopyranosides in water are known to be 68% and 32%, respectively, and those of \( \alpha- \) and \( \beta- \)glucopyranosides are 36% and 64%. From the \( ^1H \) NMR spectra at a concentration of 10 mM of the anomeric mixture of these sugars in D\(_2\)O, we obtained nearly the same values (data not shown). Using these values, we obtained Eqs. 17 and 18 for the anomeric mixtures of Man and Glc, respectively.

\[
\begin{align*}
\chi(\text{Man}) &= 0.68 \chi(\alpha-\text{Man}) + 0.32 \chi(\beta-\text{Man}) \quad (17) \\
\chi(\text{Glc}) &= 0.36 \chi(\alpha-\text{Glc}) + 0.64 \chi(\beta-\text{Glc}) \quad (18)
\end{align*}
\]

As the affinities of \( 1\alpha-\text{OMe} \) and \( 1\beta-\text{OMe} \) were both independent of the configuration of the 2-OH, it can be assumed that the same relationship is observed for sugars having the 1-OH groups such as \( \alpha-\text{Man}, \beta-\text{Man}, \alpha-\text{Glc} \) and \( \beta-\text{Glc} \). Hence the following relations are derived.

\[
\begin{align*}
\chi(\alpha-\text{Man})/\chi(\beta-\text{Man}) &= \chi(\text{Me} \alpha-\text{Man})/\chi(\text{Me} \beta-\text{Man}) = 3.66 \quad (19) \\
\chi(\alpha-\text{Glc})/\chi(\beta-\text{Glc}) &= \chi(\text{Me} \alpha-\text{Glc})/\chi(\text{Me} \beta-\text{Glc}) = 4.13 \quad (20)
\end{align*}
\]

From Eqs. 17—20, we obtained \( \chi(\alpha-\text{Man}) = 7.84, \chi(\beta-\text{Man}) = 2.14, \chi(\alpha-\text{Glc}) = 7.26 \) and \( \chi(\beta-\text{Glc}) = 1.76 \).

From these values, the selectivities \( \chi(\alpha-\text{Man})/\chi(\alpha-\text{Glc}) \) and \( \chi(\beta-\text{Man})/\chi(\beta-\text{Glc}) \) were deduced as 1.08 and 1.22, respectively, being quite different from \( \chi(\text{Me} \alpha-\text{Man})/\chi(\text{Me} \alpha-\text{Glc}) = 2.35 \) and \( \chi(\text{Me} \beta-\text{Man})/\chi(\text{Me} \beta-\text{Glc}) = 2.65 \). It is noteworthy that the selectivity of the 2-OH changing greatly for the sugars with 1-OMe and 1-OH, consistent with the results that the 2\( \alpha- \)-OH group of Glc is not directly associated with the binding to ConA, whereas the 2\( \beta- \)-OH is associated with the binding. These results were consistent with the crystallographic structures of ConA-Me \( \alpha-\text{Man} \) and ConA-Me \( \alpha-\text{Glc} \) complexes.

**Affinities of Sugars and Related Compounds to ConA**

The effects of the other hexoses: \( \text{d}-\text{galactose (Gal)}, \text{d}-\text{allose, d-altrose, d-idose and d-talose on the ASF-ConA complex were also studied with the optical biosensor. Chemical structures of these sugars are shown in the Chart 1. These sugars did not affect the ConA binding to ASF even at 300 mM. The effect of \( \text{d}-\text{galulose could not be examined because of its low solubility. As these hexoses do not contain either the 3\( \beta- \)-OH or 4\( \alpha- \)-OH, these OH groups are suggested to be essential for binding with ConA. Besides these hexoses, the effects of \( 14 \) sugars and their related compounds (chemical structures, see Chart 1) on the ASF-ConA complex were further tested up to 300 mM. We found that 2-deoxy-\( \text{d}-\text{glucose (2DOG)}, \text{N}-\text{acytelyl-\( \text{d}-\text{glucosamine (GlcNAc) and N-acytelyl-\( \text{d}-\text{mannosamine (ManNAc) enhanced the dissocation of the ASF-ConA complex, but 3-deoxy-\( \text{d}-\text{glucose (3DOG), 3-O-methyl-\( \text{d}-\text{glucose (3MeGlc), methyl \( \alpha-\text{d}-\text{galactopyranoside (Me \( \alpha-\text{Gal}), methyl \( \beta-\text{d}-\text{galactopyranoside (Me \( \beta-\text{Gal}), 6-O\text{-methyl-d-galactose, N-acytelyl-\( \text{d}-\text{galactose (GalNAc), methyl \( \beta-\text{d}-\text{arabinopyranoside, methyl \( \beta-\text{l}-\text{arabinopyranoside, brefeldin A and cytochalasin B\text{) had no significant effect. Lack of the effects of 3DOG and 3MeGlc indicated that the 3\( \beta- \)-OH group of Man and Glc acts as a proton donor for hydrogen bonding with ConA.**

To clarify the role of the functional groups at the C2-position, we next examined the dose-dependent effects of 2DOG and GlcNAc, which do not have the 2-OH group, and obtained the relations shown in Eqs. 21 and 22, respectively.

\[
\begin{align*}
1/k &= 187.4 + 39.16[2DOG] 
&= (5.3) + 5.00 \quad (21) \\
&= (n=15, r=0.978, s=53.5, F_{1,13} = 141) \\
1/k &= 187.8 + 56.67[\text{GlcNAc}] 
&= (28.8) + 7.72 \quad (22) \\
&= (n=21, r=0.962, s=101, F_{1,19} = 117)
\end{align*}
\]

There is a relation of \( \chi(\text{Man})/(=6.02)) > k(2DOG) = 4.79 > \chi(\text{Glc})/(=3.74) > \chi(\text{GlcNAc}) = 3.32 \). Therefore, it is suggested that the 2\( \beta- \)-OH (Man) is associated with the binding to ConA, and the 2\( \alpha- \)-acetamido (GlcNAc) and 2\( \alpha- \)-OH (Glc) groups decrease the binding. As the values of \( \chi(\text{Man})/\chi(\text{Glc}) \) and \( \chi(\beta-\text{Man})/\chi(\beta-\text{Glc}) \) were close to 1, configuration of the 2-OH group was not associated with the binding with ConA. Therefore, the intermediate value of \( \chi(2DOG) \) could be due to the same populations of the 1\( \alpha- \) and 1\( \beta- \)-OH anomers.

We also obtained Eq. 23 for ManNAc.

\[
\begin{align*}
1/k &= 585.0 + 172.0[\text{ManNAc}] 
&= (2471) + 52.2 \quad (24) \\
&= (n=10, r=0.935, s=293, F_{1,8} = 27.8)
\end{align*}
\]

The value of \( \chi(\text{ManNAc}) \) was determined as 3.40, being much less than \( \chi(\text{Man}) \). Therefore, substitution of 2\( \beta- \)-OH of Man by the acetamido-group decreased the binding affinity, possibly due to the steric hindrance of the acetamido-group. The \( v \) value of ManNAc was 585, much greater than those of other sugars in the range of 120—400 (see, Table 1). This could be because \( K_\text{i} \) of ManNAc was not great enough to satisfy the requirement of the experimental conditions \( [S] = 1/K_\text{i} \) in Eq. 9. Therefore, it is possible that the value of \( \chi(\text{ManNAc}) \) has not been determined exactly by Eq. 23.

**Effect of Sugars on the Erythrocyte Aggregation Induced by ConA**

To know the roles of the functional groups at the C2-position of sugars in interaction with ConA by a method other than SPR sensor, we examined the effects of the sugars Man, Glc, GlcNAc and ManNAc on the aggregation of erythrocytes induced by ConA, which interacts tightly with the mannotriose moieties of the erythrocyte membrane surface. We found that the sugar chains of the band-3 protein in bovine erythrocytes were not highly sensitive to lectins, but those of human erythrocytes are sensitive (data not shown), possibly because the population of mannotriose moiety on human erythrocyte membrane is higher than that on bovine erythrocyte membrane.

The addition of 30 mg/ml ConA to human erythrocytes at 0.7% (v/v) suspended in saline induced cell aggregation. The effects of sugars on the ConA-induced aggregation were assayed in 96-well microtiter plates. Man, Glc, GlcNAc, and ManNAc inhibited the ConA-induced cell aggregation. The minimum inhibitory concentration (MIC) was estimated as 0.3 mCl for Man, 1.5 mCl for Glc, 5.0 mCl for GlcNAc, and 20 mCl for ManNAc. It is noteworthy that ManNAc showed the inhibitory effect, although its effect was not potent.

We also observed the aggregation of bovine erythrocytes (0.7% (v/v)) on addition of the higher concentration of 300 mg/ml of ConA, due to lower ConA affinity to bovine erythrocytes than to human erythrocytes. The MIC values were 24 mCl for Man, 0.1 mCl for Glc, 0.15 mCl for GlcNAc, and
0.3 M for ManNAc, being much higher than those in human erythrocytes.

Using dextran–ConA precipitation test, Poretz and Goldstein\(^5\) reported the IC\(_{50}\) values of 2.0 mM for Man, 11.3 mM for Glc, and approximately 13 mM for GlcNAc. ManNAc showed only a partial effect. These results suggest that the affinity of sugars to ConA is dependent on the binding partner of ConA, i.e., mannotrioses in erythrocyte membrane and the Glc moiety of dextran in the assay system of Poretz and Goldstein.\(^5\) Therefore, our results showed that the sugar recognition of ConA is more sensitive in the conformation of ConA induced by binding with mannotrioses than that induced by dextran. As ASF contains mannotriose moiety, our present system well mimics biomembranes.

In conclusion, we found that the affinity of sugars for ConA is dependent on the binding partner of ConA. The conformation of ConA possibly changes on binding with the binding partner, and this conformational change then affects the binding to sugars. In the present ConA–ASF system, the binding affinity of \(\alpha\)-Glc was closer to that of \(\alpha\)-Man, and introduction of the non-methylated group to the 1-OH of these sugars considerably decreased the sugar selectivity to ConA.

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