Structure analysis of immobilized-bovine serum albumin by means of TOF-SIMS

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Orientation of immobilized proteins on bio-devices is important to obtain their high performance. Structural change in a particular area of a protein is also very important for the study of the protein performance and the evaluation of the bio-devices. Analysis of protein structure has been studied in many fields such as x-ray diffraction analysis, NMR, Raman and Infrared spectroscopy [3–5] and mass spectrometry [6, 7]. Though the spectroscopy determines chemical structures of molecules, it is difficult to analyze partial structure because proteins have the same groups. In addition, these techniques require more than one layer of molecules on substrates. An analysis method with high sensitivity and high resolution is required in order to evaluate a bio-device with bio-molecules.

Time-of-flight secondary ion mass spectrometry (TOF-SIMS) is one of the most sensitive surface analysis techniques [8]. TOF-SIMS is able to provide information on orientation and partial structure of immobilized protein based on fragment ions composed of multiple amino acids, in principle. TOF-SIMS analyzes upper surface of one layer of molecules. Orientation of immobilized proteins can be evaluated based on determination of a partial structure, indicating ensemble of amino acids, on the surface part. Large fragment ions consisting of multiple amino acids will indicate which part of a protein is on the surface. There are probably specific and important fragment ions to determine the surface part of a molecule, because current TOF-SIMS techniques produce a variety of fragment ions. Since ion generation mechanism of SIMS has not been clarified yet, fragment ions from complicated samples such as proteins cannot be predicted. Therefore analysis techniques for TOF-SIMS spectra interpretation will be helpful to find out important peaks. In this study, spectra analysis by means of mutual information [9] is employed.

Information theory [10] has been employed to analyze TOF-SIMS data on protein-containing biomaterials [11]. This innovation is now sufficiently developed so as to be able to obtain chemical images of protein samples. Mutual information [10, 12], a technical term defined by information theory, characterizes the specificity of every peak in TOF-SIMS spectra of a sample compared with another sample, such as a reference sample. With mutual information specific, the desired peaks can be selected out of a great number of peaks which appear in TOF-SIMS spectra.

TOF-SIMS has been applied to biomaterial analysis [9, 13–19] to evaluate adsorbed protein films [20], protein folding and orientation [21, 22], and protein distribution onto medical devices [10, 23–25] and to study extracellular matrix [26, 27] and plants [28–30]. Protein distribution on artificial kidney and biosensors were evaluated in the previous studies [24, 31]. Peaks of fragment ions specific to a particular protein are found out of all peaks based on values of mutual information calculating from comparison between spectra of a sample and those of reference. Since relatively strong intensity is required to obtain clear images, peaks at higher mass are, sometimes, ignored because of their weak intensities. However, peaks of fragment ions composed of two or three amino acids, which have higher mass values, are important to determine par-

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ticular part of a protein.

In this study, in order to obtain larger fragment ions at high intensities, indium-tin oxide (ITO) glass electrode was employed as a substrate for protein immobilization, because it is often difficult to produce enough fragment ions at higher mass with insulating substrate such as glass. The model protein, bovine serum albumin (BSA), was immobilized on the substrate by covalent bonding. Two types of the oriented samples were prepared by controlling the binding part, i.e., amino groups or carboxyl groups, of the protein.

The primary structure, i.e., the amino acid sequence, of BSA has been reported [32] and its structure is very similar to human serum albumin (HSA) whose three-dimensional structure has already been reported [33] in the Protein data bank (PDB: http://www.rcsb.org/pdb/). The results from TOF-SIMS spectra analysis were compared to the amino acid sequence and 3D-structure of albumin to examine surface parts of the BSA immobilized in different ways.

II. MATERIALS AND METHODS

A. Sample Preparation

An indium tin oxide (ITO) coated glass plate (Sigma-Aldrich Co., St Louis, MO, USA) was aminosilanized with aminopropyltrimethoxysilan (Tokyo Kasei, Tokyo, Japan). The aminosilanized glass plate was activated by glutaraldehyde and then soaked in a 0.1 M-phosphate buffered saline (PBS) solution at pH 7.4 containing 1 mg/ml bovine serum albumin (BSA: Sigma-Aldrich) and allowed to react in the dark for 30 hr at 227 K. After the ITO glass plates were washed in pH 7.4 phosphate buffer saline (PBS), and then they were washed in pure water with sonic waves for 10 s to remove adsorbed proteins. These ITO glass plates were then rinsed with pure water and dried with freeze dryer (VD-250F, Taitech, Saitama, Japan) before TOF-SIMS measurement.

B. TOF-SIMS

Positive ion spectra obtained with TOF-SIMS, TRIFT-IV (Physical Electronics, Eden Prairie, MN) using 30 kV Au⁺ primary ion source, were acquired up to 1000 m/z while maintaining the primary ion dose at less than 10^{12} ions/cm² to ensure static conditions. All the spectra, composed of positive ion TOF-SIMS spectra, were calibrated to the CH₃⁺, C₂H₅⁺, C₃H₇⁺, and In⁺ peaks before data analysis. Intensities of secondary ion peaks were normalized to their total ion count before calculation of the mutual information in order to correct for the differences in total secondary ion yield from spectrum to spectrum.

C. Spectrum Analysis

The calculation steps and the basic concept of classification by means of mutual information were described in the previous paper [9, 24, 25]. A value of information entropy approaches 1 when the peak used for the calculation is not useful for discriminating samples. Calculation steps of the mutual information is the followings:

\[ S(A) = -\sum p(ai) \log_2 p(ai), \]

the probability \( p(ai) = \frac{n(ai)}{N}, \) \( i = 1, 2, \)

where \( N \) is the number of all TOF-SIMS spectra, \( n(ai) \) is the number of spectra belonging to category \( i, \) and \( S(A) \) is "a priori entropy", information entropy before obtaining information on intensities of a peak.

\[ S(A | B) = -\sum \sum p(bj)p(ai | bj) \log_2 p(ai | bj), \]

the probability \( p(ai | bj) = \frac{n(ai | bj)}{n(bj)}, \)

\( j = 1 \) (higher) or 2 (lower),

where \( n(ai | bj) \) is the number of spectra higher than a threshold \( V \) out of the spectra belonging to category \( i, \) \( n(bj) \) is the number of spectra higher than a threshold \( V \) out of all spectra, \( S(A | B) \) is "a posteriori entropy", information entropy after obtaining information on intensities of a peak.

\[ I(A; B) = S(A) - S(A | B), \]

where \( I(A; B) \) is mutual information calculated by subtracting "a posteriori entropy \( S(A | B) \) from "a priori entropy \( S(A). \)

Peaks of secondary ions at \( m/z = 1 \) to 1000 were used for calculation of mutual information.

D. Peak identification and matching

Peaks of fragment ions from combinations of amino acids are identified by searching every combination of amino acids based on the following hypothesis: 1) double bonds are not cut, 2) numbers of carbon, oxygen, nitrogen and sulfur are considered, 3) consider SS bonding when there are more than two sulfur atoms. Then the origin of the identified fragment ion, formed combination of amino acids, is found through the primary sequence of BSA.

III. RESULTS AND DISCUSSION

Particular fragment ions from combination of multiple amino acids, suggesting orientations of BSA molecules, were searched in the TOF-SIMS spectra of each sample. Table I summarizes the calculation results of the mutual information. There are two types of immobilized BSA: immobilized with amino groups (N-BSA) and immobilized with carboxyl groups (C-BSA). TOF-SIMS spectra of N-BSA and those of C-BSA were compared. The peaks at \( m/z = 259, 286, 372 \) and 413 are specific to N-BSA and the peak at \( m/z = 129 \) is specific to C-BSA as shown in Table I. The peak at \( m/z = 413 \) is not able to be determined because there are too many possibilities.
TABLE I: Fragment ions specific to each sample. [AB]&[CD] means A & C, A & D, B & C or B & D.

| Specific to Peaks | Mutual information (-) | Components | Amino acids |
|------------------|------------------------|------------|-------------|
| N-BSA            | 259                    | C_{15}H_{24}N_{2}O | W&[VK]      |
|                  |                        | C_{17}H_{24}N_{2}O | [FY] & [FY] |
|                  | 372                    | C_{24}H_{24}N_{2}O | M&[VL][KR] & [FY] & [FY] & C, M&K&W |
|                  |                        | C_{16}H_{24}N_{2}O | -           |
|                  |                        | C_{19}H_{24}N_{4}O | -           |
|                  | 413                    | lots of possibilities | - |
| C-BSA            | 129                    | C_{16}H_{18}N_{2}O | R, K, G&[VL][KR], [VL][IT][K][RE]&[AVL][IM][FWST][N][Q][Y][C][RH][DE] |

The surface part of immobilized BSA is studied comparing the primary sequence of BSA with fragment ions from combination of neighbor amino acids. The sequence for BSA is summarized in the following:

MKWVTFISLLL LLFSSASYRG VFRRDTHKSE IAHRFKDLGE EHFKGLVLLIA FSQYLQCCPF
DEHVKLNIEL TEFATKVCV ESHAGCEKSL HTLFGEDELCK VASLREYGD MADCCCEKQEP
ERNECVFHSLK DDSPDLPKLK PDPMTCDEF KADEKKFVWGY YLYEIAARGHP YFYAPELLYY
ANKYNGVFQE CCQAEDKGC A LPIKETMRI KVLASSARQ R LWCSAQKF ERALKAWSVA
RLSQKFPKAE FVEVTKLVT D TKYVHEKCE GDLECAFFDR ADLAKYICDQ TQISSKLKE
CCDKPLEXK HCEAVDEKDA IPENLPIPLTA DPAEDKVDVCY NYEAKDAFLG SPFLYEYSR
HPEYAVSVIILL RLAKEYIATL EECCADDDPH ACYSTVFDKL KHLVDEPQNL IQKNDCDFKX
LGEYGFQNAL IVRYEYKTQVP VSTPTLVEVS RSLGKVGRGC CKTPESMRP CTEYDILSLIL
NRLCVLHKEK VPSEKVTKCC TESLIVNPPC FSAALTDETY VPKEAFDKL TPFADICTLP
DTEKIQKQT ALVELLKHP KATEEQLTKV MENFVAVFDK CCAADDKEAC FAVEGPPLVV
STQATALA

Underlined parts, KW (Lysine and Triptophan), WV (Triptophan and Valine), YF(Tyrosine and Phenilalanine), FY, YY, and MKW (Metionine, Lysine and Triptophan) are detected as peaks, at m/z = 259, 286 or 372, of fragment ions specific to N-BSA. These results indicate that these parts are on the surface or near the surface of N-BSA. Figure 1 shows the 3D structure of human serum albumin (HSA: 1BKE.pdb) having the very similar sequence to BSA. 3D structure of a protein will change depending on its condition. Especially, because these samples have been freeze-dried when they are measured with TOF-SIMS, they will not keep the same structure when they form in crystal. However, this 3D structure suggests, at least, a cue to identify areas facing the surface depending on the binding parts by comparing C-BSA with N-BSA treated in the same processes.

The fragment ions at m/z = 259 and 286 are produced from the circled parts of the HSA structure shown in Figure 1. Though the KW and WV parts are not shown in Figure 1, these parts will be near the NH2-terminal of the HSA molecule shown in the figure, according to the amino acid sequences of both albumins.

Since the peak at m/z = 129 specific to C-BSA has lots of possible origins, it is difficult to determine parts which produce the fragment ions. Furthermore, the peak at m/z = 129 is also observed in the TOF-SIMS spectra of N-BSA due to many possible combinations of amino acids producing the peak in many parts of BSA, though it is mainly specific to C-BSA. Therefore it is difficult to identify the surface part of C-BSA with this peak.

Albumin contains several amino acids having amino groups, i.e., asparagine (N), glutamine (Q), Lysine (K) and arginine (R), or carboxyl groups, i.e., aspartic acid (D) and glutamic acid (E), in the side chains. Therefore, there may be immobilized BSA molecules having different orientations even when they are immobilized at amino groups, because there are so many amino groups in a BSA molecule that it is difficult to identify the parts of BSA binding to the substrates. However, considering the fragment ions related to ensemble of amino acids, at least, it is indicated that most of BSA molecules immobilized with amino groups by covalent bonds have the same orientation placing the square-marked part on the surface of immobilized BSA molecules as shown in Figure 1. In addition, the NH2-terminal of the BSA molecule may not be used in the immobilization processes for N-BSA described in this study, because the NH2-terminal of the BSA molecule may be at the inside of the molecule. Further study is required to obtain detailed information on orientation of immobilized proteins.
IV. CONCLUSIONS

The surface parts of the BSA molecules immobilized on ITO glass plates with different parts of the molecule, amino groups or carboxyl groups, is evaluated by means of TOF-SIMS. Fragment-ion-generating parts of N-BSA are determined with fragment ions from particular combinations of amino acids in the sequence of BSA, and the surface part of the N-BSA is indicated. Further study is needed to obtain detailed information on orientation of immobilized proteins on bio-devices.

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