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Quantification of azide groups on a material surface and a biomolecule using a clickable and cleavable fluorescent compound†

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Here, we propose a novel method for quantifying azide groups on a solid surface and a protein using a clickable and cleavable fluorescent compound. The clickable and cleavable fluorescent compound conjugates with the azide groups on the material surface and the fluorophore is then liberated into the solvent via a cleavage reaction, which can be simply quantified with a conventional fluorometer.

Introduction

A bioorthogonal reaction is a chemical reaction that can be selectively and reliably performed in a complex biological environment without non-favorable interactions and interference with the biological system.¹⁻³ Huisgen 1,3-dipolar cycloaddition between an azide and an alkyne (the azide/alkyne cycloaddition click reaction), which is one of the most famous bioorthogonal reactions, proceeds in a high yield under mild aqueous conditions without byproducts.⁴⁻⁶ This chemistry has been applied to a number of biomolecules and synthetic compounds to create multifunction-integrated molecular systems. This click reaction has also been used to immobilize biomolecules, ligands, and polymers on substrate surfaces⁷⁻¹¹ for microarray preparation,¹² fabrication of biomolecule-conjugated nanoparticles¹³⁻¹⁶ and fluorescent labeling of living cells.¹⁷⁻¹⁹

When click chemistry is applied to sensitive analytical and biological devices, quantification of the reactive sites plays an important role in controlling the amount of azide groups (or alkyne groups) present on a substrate surface, because the surface density of the functional molecules affects the performance of the substrate.²⁰,²¹ However, to our knowledge, there have been no reports of quantification methods for clickable moieties (azide and alkyne groups) on substrate surfaces. Conventional surface analysis methods, such as X-ray photon spectroscopy (XPS), Fourier transform infrared spectroscopy and secondary-ion mass spectrometry are qualitative and provide information about the elements or functional groups not only on the outermost surface but also those at a certain depth (e.g., 10 nm for XPS) from the surface.²²,²³ These methods require a high vacuum environment or dry atmosphere, and the surfaces of polymeric materials vary depending on the environment. Therefore, it is of great importance to quantify the clickable moieties on the outermost surface in a practical environment (e.g., aqueous solution) for surface modification based on click chemistry.

In this study, we aimed to develop a method for quantification of the azide groups present on the outermost surface in an aqueous environment. We synthesized a clickable and cleavable fluorescent compound that conjugates with the azide groups on the substrate surface via Huisgen 1,3-dipolar cycloaddition (Fig. 1). Hydrolysis of the ester bond in the fluorescent compound liberates the fluorophore into the bulk solution, which allows the azide groups on a substrate surface to be quantified with a conventional fluorometer. The present methodology can be also applied to the quantification of azide groups conjugated with a protein.

Experimental

Materials

2,4-Dihydroxybenzaldehyde, 2,2-dimethyl-1,3-dioxane-4,6-dione, piperdinium acetate, 2-hydroxy-4-methoxybenzaldehyde, propargyl-PEG₄-alcohol, and azidobenzoic acid were purchased from Tokyo Chemical Industry (Tokyo, Japan). K₂CO₃, propargyl-PEG₄-alcohol, and azidobenzoic acid were purchased from Tokyo Chemical Industry (Tokyo, Japan). K₂CO₃, propargyl-

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Fig. 1 Schematic of quantification of the azide groups on substrate surfaces using the clickable and cleavable fluorescent compound.
bromide, CuSO₄, β-casein from bovine milk, rhodamine B isothiocyanate (RITC), and Tris–EDTA buffer solution (pH 7) were purchased from Sigma-Aldrich (St. Louis, MO). DBCO-s-amine was purchased from Conju-Probe LLC (San Diego, CA). The other chemicals were purchased from Wako Pure Chemical Industries, Ltd. (Osaka, Japan). NH₂-PEG-resin particles (10 μm) were purchased from Watanabe Chemical Industries (Hiratsuka, Japan). Slide-A-Lyzer G2 dialysis cassettes (20k MWCO, 15 mL) and micro BCA protein assay were purchased from Thermo Fisher Scientific (Waltham, USA). High-quality deionized (DI) water (>15 MΩ cm) produced by an Elix-5 system (Millipore, Molsheim, France) was used in the experiments.

**Synthesis of 7-methoxycoumarin-3-carboxylic acid.** 2-Hydroxy-4-methoxybenzaldehyde (50 mmol), 2,2-dimethyl-1,3-dioxane-4,6-dione (50 mmol), and piperidinium acetate (1 mmol) were dissolved in dehydrated ethanol (25 mL). The mixture was stirred at room temperature for 10 min and refluxed for 3 h. The reaction mixture was cooled to room temperature, and then further cooled in an ice bath for 30 min. The precipitate was collected by filtration and washed with ethanol three times. The product was dried overnight in a vacuum chamber.

1H-NMR (500 MHz, CDCl₃): δ (ppm) 6.99 (d, 1H, J = 2.2 Hz), 7.03 (dd, 1H, J = 2.6, 8.9 Hz), 7.65 (d, 1H, J = 8.8 Hz), 8.86 (s, 1H), 12.18 (s, 1H). DART-TOF/MS ([M + H]⁺) calcd for C₂₀H₂₁NO₅, 391.1; found, 391.2. Yield: 21%. This journal is © The Royal Society of Chemistry 2019

1H-NMR (500 MHz, CDC₁₃): δ (ppm) 5.97 (s, 3H), 6.93 (d, 1H, J = 2.2 Hz), 7.03 (dd, 1H, J = 2.6, 8.9 Hz), 7.65 (d, 1H, J = 8.8 Hz), 8.86 (s, 1H), 12.18 (s, 1H). DART-TOF/MS ([M + H]⁺) calcd for C₁₄H₁₄O₄, 221.0; found, 221.0. Yield: 83%.

**Click reaction of C≡C-EG₃-CMRN.** A solution of C≡C-EG₃-CMRN (2.5 μmol) and azidobenzoic acid (2.5 μmol) in DMSO (0.4 mL) was mixed and stirred with an aqueous solution (1.6 mL) of CuSO₄ (6 μmol) and sodium ascorbate (30 μmol) at 37 °C for 3 h. The solution was analyzed as described above (click reaction of C≡C-EG₃-CMRN).

1H-NMR (500 MHz, DMSO-d₆): δ (ppm) 3.54–3.61 (m, 8H), 3.71 (t, 2H), 3.88 (s, 3H), 4.34 (t, 2H, J = 2.2 Hz), 4.61 (s, 2H), 6.98 (dd, 1H, J = 2.5, 8.8 Hz), 7.01 (s, 1H), 7.82 (d, 1H, J = 8.6 Hz), 7.98 (d, 2H, J = 8.5 Hz), 8.09 (d, 2H, J = 8.9 Hz), 8.69 (s, 1H), 8.86 (s, 1H). Yield: 96%.

**Cleavage.** NaOH solution (0.1 M, 0.5 mL) was added to the solution after the click reaction of C≡C-EG₃-CMRN with azidobenzoic acid. The mixture was stirred at 37 °C for 3 h and analyzed by HPLC.

**Azidation of the resin particles.** The resin particles had 65 mmol g⁻¹ amino groups on their surfaces. A dispersion of amino-resin particles (3.8 μmol amino groups), azidobenzoic acid (380 μmol), and 4-[4-(6-dimethoxy-1,3,5-triazin-2-yl)-4-methylmorpholinium chloride (DMT-MM, 75 μmol) was mixed in DMSO (3 mL) and stirred at 37 °C for 24 h to conjugate azidobenzoic acid with the amino groups on the resin particles. The resin particles were washed four times with methanol (6 mL) and water (6 mL).

**Quantiﬁcation of the azide groups on the resin particles using C≡C-CMRN.** Azidated resin particles (0.1875, 0.375, or 0.75 mg) were dispersed in water (1.8 mL containing CuSO₄ (1 μmol) and sodium ascorbate (4 μmol). The dispersion was added to 1 mM C≡C-EG₃-CMRN in DMSO (0.1 mL) and shaken at 40 °C for 24 h. After centrifugation, the supernatant was removed and the resin particles were washed three times with methanol and water. The resin particles were dispersed in NaOH aqueous solution (1 M, 0.2 mL) and then shaken at 37 °C for 24 h to liberate CMRN from the resin particles. After centrifugation, HCl solution (1 M, 0.2 mL) and phosphate buffer (0.1 M, pH 7.4, 2 mL) were added to neutralize the supernatant. The fluorescence intensity of the neutralized solution was measured with a fluorescence spectrophotometer. The excitation and emission wavelengths were 335 and 406.5 nm, respectively. The excitation and emission band widths were 5 nm. The standard curve of a liberated fluorophore in phosphate buffer is shown in Fig. S2a.† Quantification was performed in triplicate, unless specified otherwise.
Quantification of the azide groups on the resin particles using strain-promoted (Cu-free) click chemistry (Fig. S3†). Azidated resin particles (0.1875, 0.375, or 0.75 mg) were dispersed in 10 mM HCl (1.4 mL). The dispersion was added to 0.6 mL tetrahydrofuran (THF) containing 1 mM DBCO-s-s-NH₂ and shaken at 40 °C for 12 h. After centrifugation, the supernatant was removed and the resin particles were washed three times with THF/10 mM HCl aqueous solution (1/1, v/v) and once with phosphate buffer (50 mM, pH 8). The resin particles dispersed in phosphate buffer (50 mM, 1 mL) were added to phosphate buffer (50 mM, 2 mL) containing 1 mM RITC and 10 vol% trimethylamine (TEA) and then shaken at 40 °C for 3 h. The supernatant was removed and phosphate buffer (50 mM, 2 mL) containing 1 mM RITC and 10 vol% TEA was added again and shaken at 40 °C for 3 h. After centrifugation, the supernatant was removed and the resin particles were washed with methanol/10 mM NaOH aqueous solution (1/1, v/v), methanol/10 mM HCl aqueous solution (1/1, v/v) and phosphate buffer (50 mM, pH 8). After centrifugation, phosphate buffer (50 mM, pH 8, 2 mL) containing 4 mM tris(2-carboxyethyl)phosphine were added to liberate the fluorophore. The fluorescence intensity of the solution was measured with a fluorescence spectrophotometer. The excitation and emission wavelengths were 495 and 515 nm, respectively. The excitation and emission band widths were 5 nm.

Quantification of the azide groups on β-casein by TNBS. β-Casein was dissolved in 0.1 M NaHCO₃ buffer (pH 8) at concentrations of 20–200 μg mL⁻¹ to construct the standard curve. A solution of 0.01% (w/v) TNBS (0.25 mL) was added to sample solutions (0.5 mL) containing 100 μg mL⁻¹ azidated β-casein or β-casein, and the solutions were then incubated at 37 °C for 2 h. After the reaction, 10% SDS aqueous solution (0.25 mL) and 1 M HCl aqueous solution (0.125 mL) were added to the sample solution to quench the TNBS reaction. The absorbance of the solutions was measured at 335 nm.

Quantification of the azide groups on azidated β-casein. Azidated casein (50 nmol) was dissolved in 25 mM NaHCO₃ buffer (pH 8, 1.2 mL) containing CuSO₄ (2.4 μmol) and sodium ascorbate (12 μmol). C≡C–EG₃–CMRN (5 μmol) in DMSO (0.4 mL) was added to the solution and the solution was stirred at 37 °C for 3 h. After the reaction, excess C≡C–EG₃–CMRN and Cu(i) catalyst were removed by dialysis for 2 days. NaOH solution (1 M, 0.5 mL) was added to the solution and the completion of the Cu(i) complex was confirmed by HPLC-MS. When Cu(i) was not removed, Cu(i) complex was eluted between 7.7 and 14.6 min. The Cu(i) complex was not removed, NaOH aqueous solution (1 M, 0.5 mL) was added in the solution. Sodium ascorbate was added in 25 mM NaHCO₃ buffer (pH 8, 1.2 mL) containing CuSO₄ (2.4 μmol) and sodium ascorbate (12 μmol). C≡C–EG₃–CMRN (5 μmol) was added to the solution and the solution was stirred at 37 °C for 3 h. The Cu(i) complex was removed by dialysis for 2 days. The solution was neutralized with NaOH aqueous solution (1 M, 0.5 mL) and NaHCO₃ buffer (1 M, pH 8, 2 mL) was added to neutralize the solution. The fluorescence intensity of the solution was measured with a fluorescence spectrophotometer. The excitation and emission wavelengths were 335 and 406.5 nm, respectively. The standard curve of a liberated fluorophore in carbonate buffer is shown in Fig. S2b†. Quantification was performed in triplicate, unless specified otherwise.

Results and discussion

First, we designed and synthesized two different types of clickable and cleavable fluorescent compounds: C≡C–CMRN and C≡C–EG₃–CMRN (Fig. 2a). The alkyl group acts as the clickable part and the ester bond acts as the cleavable part in hydrolysis. C≡C–CMRN has no spacer between the ester bond and alkyl, but C≡C–EG₃–CMRN has an oligo (ethylene glycol) spacer. The click reactivities of C≡C–CMRN and C≡C–EG₃–CMRN were investigated in a homogeneous solution. Azido-benzoic acid was used as a model azide compound and the click reaction was performed with a copper catalyst in a mixture of water/dimethylformamide (DMF) (5/1, v/v). High-performance liquid chromatography (HPLC) measurements of C≡C–CMRN indicate simultaneous synthesis of compound 1 at a retention time (RT) of 15.3 min and production of 7-methylcoumarin-3-carboxylic acid (Fig. S1†), indicating that hydrolysis of the ester bond is probably due to chelation of a Cu ion at the neighboring site. HPLC measurements of C≡C–EG₃–CMRN reveal that the peak at RT = 14.6 min disappears and a peak at RT = 7.7 min appears (Fig. 2b). ¹H-NMR analysis of these fractions reveals that the peaks at RT = 14.6 and 7.7 min correspond to C≡C–EG₃–CMRN and compound 1 (the product of the click reaction), respectively. These results indicate that C≡C–EG₃–CMRN can be used as a substrate in the azide/alkyne click reaction and that the reaction proceeds in high yield (>96%).

Next, we investigated hydrolysis of the ester bond of C≡C–EG₃–CMRN. NaOH aqueous solution was added to the solution containing compound 1 to give a final NaOH concentration of 0.1 M, and the mixture was then stirred at 37 °C for 3 h. HPLC measurements show that the peak of compound 1 (at RT = 7.7 min) completely disappears and a new peak of 7-methylcoumarin-3-carboxylic acid appears at RT = 5.4 min (Fig. 2b). These results demonstrate that compound 1, which is the product of the C≡C–EG₃–CMRN/azidobenzoic acid click reaction, can be hydrolyzed by aqueous NaOH to liberate its fluorophore.
We then attempted to quantify the azide groups on resin particles using \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN. In a previous study, we reported that the resin particles have 65 nmol mg\(^{-1}\) amino groups.\(^{24}\) Before quantification, we produced azide groups on the resin particles by the condensation reaction between azidobenzoic acid and the amino groups on the resin particles (azidation, Fig. 3a). The azide groups on the resin particle surfaces were quantified using \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN. Azidated resin particles (0, 0.19, 0.38, or 0.75 mg) were dispersed in water (1.8 mL) containing CuSO\(_4\) (1 \( \mu \)mol) and sodium ascorbate (4 \( \mu \)mol). The dispersion was mixed with dimethyl sulfoxide (DMSO, 0.1 mL) containing 1 mM \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN and shaken at room temperature for 24 h. After washing the resin particles, the resin particles were dispersed in NaOH aqueous solution (final concentration of 0.1 M) to liberate the CMRN fluorophore from the resin particles. After centrifugation and neutralization, the fluorescence intensity of the solution was measured with a conventional fluorescence spectrophotometer. There is high linearity between the amount of particles and the fluorescence intensity of the liberated fluorophore (Fig. 2b). The quantified value of the azide groups on the resin particles is 55 nmol mg\(^{-1}\), which is comparable with the amount of amino groups on the resin particles. The slightly smaller value for the azide groups is because of the yield of azidation of the amino groups on the resin particles. These results suggest successful quantification of the azide groups on the resin particles using the proposed method. The standard curve (Fig. S4†) shows that the limit of detection of the present method was 5 nmol ml\(^{-1}\), which means that azide groups of 5 pmol on a surface was detected when using 1 mL aqueous solution of liberated fluorophore.

While the above investigations employed Cu(i) as a catalyst for 1,3-dipolar cycloaddition of azide and alkyne, a Cu-free click reaction (strain-promoted azide–alkyne cycloaddition) using a cycloalkyne has attracted a broad range of attention because of no use of a catalyst.\(^{25}\) We then attempted to quantify the azide groups on resin surfaces using a Cu-free click reaction (Fig. S3†). Although a dibenzocyclooctyne (DBCO) derivative is a useful functional molecule for a Cu-free click reaction, its water solubility is very low. We conjugated DBCO-s-s-amine with azide groups on resin surfaces and then the resultant amino groups were conjugated with rhodamine B isothiocyanate (RTIC). The disulfide bond was cleaved by a reducing agent (TCEP) to liberate the fluorophore, which was measured using a fluorometer. Although the liberated fluorophore increased linearly with the amount of azidated resin particles (Fig. S5†), we detected only 4.2 nmol azide groups per 1 mg resin. The remarkably low amount of azide groups quantified might be explained by the following possible reasons. (i) The sterically bulky structure of DBCO-s-s-amine affected its conjugation with an azide group on a surface, leading to a low conjugation ratio. (ii) A mixture of THF/water, which was used to dissolve DBCO-s-s-amine for a Cu-free click reaction, altered the surface composition of resin, resulting in the inhibition of the azide group quantification. These results suggest that a relatively small fluorescent compound, which has water solubility and avoids steric hindrance, is appropriate for the quantification of azide groups on a surface. The following experiments were carried out using \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN.

In the last decade, clickable moieties have often been introduced to biomolecules (e.g., proteins and nucleic acids) to create functionally engineered biomolecules and immobilize them on various surfaces.\(^{26–30}\) Therefore, we then investigated quantification of azide groups introduced to a protein. \( \beta \)-Casein was chosen as a model protein and azide groups were introduced to the amino groups of \( \beta \)-casein by \( N \)-hydroxy succinimide ester chemistry (Fig. 4a). \( \beta \)-Casein has 12 amino groups.\(^{31}\) The TNBS method\(^{32–33}\) shows that the azidation procedure conjugates 86% of the amino groups with azidobenzoic acid. Azidated \( \beta \)-casein (50 nmol) in an aqueous buffer (pH 7.0, 1.2 mL) containing CuSO\(_4\) (2.4 \( \mu \)mol) and sodium ascorbate (12 \( \mu \)mol) was mixed with DMSO (0.4 mL) containing \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN (5 \( \mu \)mol) and then shaken at 37 °C for 3 h. After dialysis to remove excess \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN and the Cu catalyst, NaOH aqueous solution was added to the solution to liberate the CMRN moiety from azidated \( \beta \)-casein.

Fluorescence measurement of the solution shows 410 nmol CMRN moieties per milligram of casein (Fig. 4b), meaning that 81% of the amino groups of \( \beta \)-casein are azidated. The obtained azidation ratio is comparable with the result of the TNBS method.

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**Fig. 3** (a) Schematic of azidation of the amino groups on the resin particles. (b) Quantification of the azide groups on the resin particles using \( \text{C} \equiv \text{C} \equiv \text{EG}_3 \)-CMRN.

**Fig. 4** (a) Schematic of azidation of the amino groups on \( \beta \)-casein. (b) Quantification of the azide groups on \( \beta \)-casein.
Conclusions

In conclusion, we have proposed a novel approach to quantify the azide groups on a material surface and on a protein using a clickable and cleavable fluorescent compound (C≡C-EG1-CMRN). This is the first time that quantification of the azide groups on a solid material and a biomolecule has been achieved. Quantitative analysis of the clickable moieties on a surface will play an important role in surface chemistry and extend the applicability of the azide/alkyne cycloaddition click reaction on surfaces and at interfaces.

Conflicts of interest

There are no conflicts to declare.

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