Video Article

Design, Synthesis, and Photochemical Properties of Clickable Caged Compounds

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URL: <https://www.jove.com/video/60021>

DOI: [doi:10.3791/60021](https://doi.org/10.3791/60021)

Keywords: Chemistry, Issue 152, caged compounds, synthesis, photochemistry, click chemistry, prodrug, chemical biology, chemical probes

Date Published: 10/15/2019

Citation: Suzuki, A.Z., Shiraishi, Y., Aoki, H., Sasaki, H., Watahiki, R., Furuta, T. Design, Synthesis, and Photochemical Properties of Clickable Caged Compounds. *J. Vis. Exp.* (152), e60021, doi:10.3791/60021 (2019).

Abstract

Caged compounds enable the photo-mediated manipulation of the cell physiology with high spatiotemporal resolution. However, the limited structural diversity of currently available caging groups and the difficulties in synthetic modification without sacrificing their photolysis efficiencies are obstacles to expanding the repertoire of caged compounds for live cell applications. As the chemical modification of coumarin-type photo-caging groups is a promising approach for the preparation of caged compounds with diverse physical and chemical properties, we report a method for the synthesis of clickable caged compounds that can be modified easily with various functional units via the copper(I)-catalyzed Huisgen cyclization. The modular platform molecule contains a (6-bromo-7-hydroxycoumarin-4-yl)methyl (Bhc) group as a photo-caging group, which exhibits a high photolysis efficiency compared to those of the conventional 2-nitrobenzyls. General procedures for the preparation of clickable caged compounds containing amines, alcohols, and carboxylates are presented. Additional properties such as the water solubility and cell targeting ability can be readily incorporated into clickable caged compounds. Furthermore, the physical and photochemical properties, including the photolysis quantum yield, were measured and were found to be superior to those of the corresponding Bhc caged compounds. The described protocol could therefore be considered a potential solution for the lack of structural diversity in the available caged compounds.

Video Link

The video component of this article can be found at <https://www.jove.com/video/60021/>

Introduction

Caged compounds are designed synthetic molecules whose original functions are temporally masked by covalently attached photo-removable protecting groups. Interestingly, caged compounds of biologically relevant molecules provide an indispensable method for the spatiotemporal control of the cellular physiology^{1,2,3,4,5,6}. In 1977, Engels and Schlaefer reported the 2-nitrobenzyl ester of cAMP as a membrane permeable and photolabile derivative of cAMP⁷. The following year, Kaplan reported the 1-(2-nitrophenyl)ethyl ester of ATP (NPE-ATP) and named this compound "caged" ATP⁸. Since then, a range of photochemically removable protecting groups such as 2-nitrobenzyls, *p*-hydroxyphenacyls⁹, 2-(2-nitrophenyl)ethyls^{10,11}, 7-nitroindolin-1-yls^{12,13}, and (coumarin-4-yl)methyls^{14,15,16} have been used for the preparation of caged compounds.

The synthesis of caged compounds with desirable additional properties such as membrane permeability, water solubility, and cellular targeting ability would be expected to facilitate cell biological applications. Since the physical and photochemical properties of these molecules depend primarily on the chemical structure of the photochemically removable protecting groups used to prepare them, a diverse repertoire of photo-caging groups is required. However, the structural diversity of currently available caging groups that exhibit high photolysis efficiencies is limited. This could be an obstacle to increasing the use of caged compounds.

To address this issue, the repertoire of photo-caging groups has been expanded by the chemical modification of existing photoremovable protecting groups or the design of new photolabile chromophores with superior photophysical and photochemical properties. Examples include nitrobenzofuran (NDBF)¹⁷, [3-(4,5-dimethoxy-2-nitrophenyl)-2-butyl] (DMNPB)^{18,19}, a calcium-sensitive 2-nitrobenzyl photocage²⁰, substituted coumarinylmethyls (DEAC450²¹, DEAdcCM²², 7-azetidiny-4-methylcoumarin²³, and styryl coumarins²⁴), cyanine derivatives (CyEt-pan)²⁵, and BODIPY derivatives^{26,27}.

In addition, we previously developed the (6-bromo-7-hydroxycoumarin-4-yl)methyl (Bhc) group and successfully synthesized various caged compounds of neurotransmitters²⁸, second messengers^{29,30}, and oligonucleotides^{31,32,33} exhibiting large one- and two-photon excitation cross-sections. If additional properties can be installed easily into the caging groups without compromising their photosensitivity, then the repertoire of caged compounds can be expanded^{34,35,36,37,38,39}. We therefore designed modular caged compounds that comprise three parts, namely the Bhc group as a photo-responsive core, chemical handles for the installation of additional functionalities, and the molecules that are to be masked^{40,41}.

Thus, this article provides a practical method for the preparation of caged compounds of biologically relevant molecules. The present protocol describes methods for the preparation of a clickable platform for photo-caging groups, the introduction of additional functionalities to expand

the repertoire of caged compounds, the measurement of their physical and photochemical properties, and the cell-type selective targeting of a clickable caged compound for further cellular application.

Protocol

1. Synthesis of the modular caging paBhc group for clickable caged compounds^{28,41}

1. Preparation of (6-bromo-7-hydroxycoumarin-4-yl)methyl chloride (Bhc-CH₂Cl)

1. Place 4-bromoresorcinol (9.742 g, 51.5 mmol) in a 100 mL round-bottomed flask equipped with a stirrer bar.
2. Add conc. H₂SO₄ (98%, 30 mL) to the flask and stir the mixture to dissolve.
3. Add ethyl 4-chloroacetoacetate (10 mL, 74 mmol) dropwise.
4. Continue stirring the mixture at ambient temperature for 5 days.
5. Separately, place crushed ice cubes (~200 mL) in a 500 mL Erlenmeyer flask.
6. Pour the reaction mixture into the ice and stir vigorously for 30 min until a finely powdered precipitate is obtained.
7. Collect the precipitate via vacuum filtration. Wash the light brown precipitate with water five times.
8. Dry the precipitate under vacuum overnight to yield BhcCH₂Cl as a light brown powder (13.57 g, 46.9 mmol).

2. Preparation of (6-bromo-7-hydroxycoumarin-4-yl)methanol (BhcCH₂OH)

1. Place the prepared BhcCH₂Cl (1.1440 g, 3.95 mmol) and 1 M HCl (300 mL) in a 1 L round-bottomed flask equipped with a Dimroth condenser. Stir the mixture at 140 °C for 5 days. After this time, cool the mixture to ambient temperature.
 2. Remove the water from the reaction by rotary evaporation under vacuum to yield BhcCH₂OH (1) as a light brown powder (1.0359 g, 3.82 mmol, 97% yield).
- NOTE: The use of 250 mL of 1 M HCl per 1 g of BhcCH₂Cl gives a satisfactory result.

3. Preparation of paBhcCH₂OH (2) via the Mannich reaction

1. Place paraformaldehyde (446.4 mg, 14.9 mmol) in a 50 mL round-bottomed flask. Add anhydrous ethanol (5 mL) and *N*-methylpropargylamine (1.25 mL, 14.8 mmol) to the flask.
2. Stir the mixture at ambient temperature for 1 h under an Ar atmosphere.
3. Add BhcCH₂OH (1) (1.367 g, 5.04 mmol) to the flask. Heat the mixture to 80 °C with a block heater apparatus, and continue stirring the mixture at 80 °C for 2 h under an Ar atmosphere.
4. Stop the block heater and cool the reaction mixture to room temperature.
5. Collect the resulting light brown-yellow precipitate by vacuum filtration. Wash the precipitate twice with a small amount of anhydrous ethanol (1 mL each time).
6. Remove the excess ethanol under vacuum to yield paBhcCH₂OH (2) (1.393 g, 3.96 mmol).

2. Preparation of clickable caged compounds

NOTE: The following procedures can be applied to the preparation of other clickable caged compounds containing hydroxyl, amino, and carboxylate functional groups.

1. General Procedure 1: Preparation of a clickable caged amine

1. Place paBhcCH₂OH (2) (709.6 mg, 2.02 mmol) and *N,N'*-carbonyl diimidazole (CDI, 397.6 mg, 2.45 mmol) in a 30 mL round-bottomed flask. Add dry CH₂Cl₂ (6 mL) and stir the solution at ambient temperature for 1 h.
2. Add 4-dimethylaminopyridine (4-DMAP, 324.8 mg, 2.66 mmol) and *tert*-butyl (6-aminoethyl)carbamate (533.1 mg, 2.46 mmol). Stir the solution at ambient temperature for 3 h.
3. Remove the solvent and other volatile materials using a rotary evaporator under vacuum. Purify the residue directly using silica gel flash column chromatography.

2. General Procedure 2: Preparation of a clickable caged alcohol

1. Place paclitaxel (PTX, 48.7 mg, 0.057 mmol) in a 30 mL round-bottomed flask equipped with a three-way stopcock and an Ar balloon. Add dry CH₂Cl₂ (1 mL), 4-DMAP (17.1 mg, 0.14 mmol), and 4-nitrophenyl chloroformate (26.0 mg, 0.13 mmol).
2. Stir the solution at ambient temperature for 2.5 h under an Ar atmosphere.
3. Add 4-DMAP (15.7 mg, 0.13 mmol) and paBhcCH₂OH (2) (39.1 mg, 0.111 mmol) to the solution. Continue stirring the mixture at ambient temperature for 17 h.
4. Add CHCl₃ (10 mL) and 15% aqueous NaHCO₃ (5 mL) to the mixture. Stir the mixture vigorously for approximately 3 min. Remove the aqueous layer with a pipette.
5. Add 0.5 M citric acid (5 mL) to the flask containing the organic layer. Stir the mixture and remove the aqueous layer as above.
6. Separate the organic layer using a phase separation column. Remove the solvents with a rotary evaporator under vacuum. Purify the product using standard silica gel flash column chromatography.

3. General Procedure 3: Preparation of a clickable caged carboxylic acid

1. Dissolve arachidonic acid (33.0 µL, 0.100 mmol), paBhcCH₂OH (2) (39.6 mg, 0.112 mmol), and 4-DMAP (14.1 mg, 0.115 mmol) in dry CH₂Cl₂ (2 mL). Add *N,N'*-diisopropylcarbodiimide (DIPC, 17.0 µL, 0.110 mmol) and stir the solution at ambient temperature for 140 min.
2. Remove the solvent under vacuum. Purify the residue directly using silica gel flash column chromatography.

3. Installation of a functional unit into the clickable caged compounds

1. Dissolve copper(II) sulfate pentahydrate (249 mg) in ion-exchanged water (IEW, 10 mL) to give a 0.1 M CuSO₄ solution.
 2. Dissolve 2'-paBhcmoc-PTX (8.0 mg, 6.5 μmol), tris(3-hydroxypropyltriazolylmethyl)amine (THPTA, 17.5 mg, 40.3 μmol), sodium l-ascorbate (162.4 mg, 0.825 mmol), and 15-chloro-3,6,9-trioxapentadecyl azide (3.1 mg, 11 μmol) in a mixed solvent of 0.1 M phosphate buffer (2.5 mL, pH 7.2) and dimethyl sulfoxide (DMSO, 0.5 mL).
 3. Add the 0.1 M CuSO₄ solution (81.2 μL, 8.1 μmol) to the reaction mixture. Stir the mixture at ambient temperature for 80 min. Monitor the progress of the reaction using high-performance liquid chromatography (HPLC).
 4. Dissolve the precipitates by adding a 75% acetonitrile/water solution (3.5 mL). Apply the resulting solution directly to the semi-preparative HPLC system to purify the desired product.
- NOTE: Solubilization of the reaction mixture by the addition of *tert*-butanol can accelerate the progression of the reaction.

4. Photolytic uncaging reaction of the caged compounds

1. Preparation of the stock solutions

1. Dissolve the desired caged compound (5 μmol) in DMSO (500 μL) to prepare a 10 mM stock solution. Dispense an aliquot of each solution (10 μL) into a 1.5 mL microcentrifuge tube and store in a freezer (−20 °C) until just before use.
2. 6 mM K₃[Fe(C₂O₄)₃] (100 mL): Dissolve recrystallized potassium ferrioxalate (0.295 g, 0.675 mmol) in 80 mL of water. Add 0.5 M H₂SO₄ (10 mL) and an appropriate amount of IEW to make up the volume to 100 mL.
NOTE: Potassium ferrioxalate should be purified via recrystallization from hot water and stored in the dark. Recrystallized potassium ferrioxalate is obtained as the trihydrate; therefore, its formula is K₃[Fe(C₂O₄)₃]·3H₂O and a formula weight of 491.24 should be considered during preparation of the stock solution. Check the purity of the 6 mM solution by measuring its absorption at 510 nm. If the absorbance is <0.02, it is suitable for use in the experiment.
3. 0.1% Buffer-phen (30 mL): Dissolve NaOAc·3H₂O (7.35 g), 1,10-phenanthroline (phen)·H₂O (30 mg), and conc. H₂SO₄ (0.9 mL) in IEW (20 mL). Add IEW to make up the volume to 30 mL.
NOTE: The solution contains 1.8 M NaOAc, 0.54 M H₂SO₄, and 0.1% 1,10-phenanthroline.

2. Measurement of the number of photons using ferrioxalate actinometry

1. Place 6 mM K₃[Fe(C₂O₄)₃] (V₁ L) in a quartz cuvette. Irradiate the solution with 350 nm light for 5 s.
2. Transfer the irradiated solution to a Pyrex cuvette with an *l* [cm] path length.
3. Add 0.1% Buffer-phen (V₂ L) to the irradiated sample solution and mix well by pipetting. Measure the absorbance of the sample at 510 nm. Calculate the average absorption change per unit time (ΔA_{510} [s^{−1}]).
4. Calculate the number of moles of generated Fe²⁺ ions per unit time according to the following equation:

$$n\text{Fe}^{2+} [\text{mol s}^{-1}] = ((V_1 + V_2) [L] \times \Delta A_{510} [\text{s}^{-1}]) / (l [\text{cm}] \times \epsilon_{510} [\text{L mol}^{-1} \text{cm}^{-1}])$$
 where (V₁ + V₂) is the volume of the sample for the absorption measurement, *l* is the optical path length of the cuvette, and ϵ_{510} is the molar absorptivity of the Fe²⁺-phen complex at 510 nm.
 NOTE: In the typical experimental conditions, values of V₁ = 2.0 × 10^{−3} L, V₂ = 0.33 × 10^{−3} L, *l* = 1.0 cm, and ϵ_{510} = 1.1 × 10⁴ L mol^{−1} cm^{−1} were used.
5. Calculate the number of moles of photons that reach the sample (*I*₀) using the following formula:

$$I_0 [\text{einstein cm}^{-2} \text{s}^{-1}] = n\text{Fe}^{2+} / \Phi_{350}$$
 where Φ_{350} is the quantum efficiency of photoreduction of the ferrioxalate at 350 nm.
 NOTE: Although the quantum efficiency of the potassium ferrioxalate actinometer at 350 nm is not reported, the reported value of 1.25⁴² at 358 nm was employed.

3. Quantum efficiency measurements at 350 nm

1. Dilute the sample stock solution (in DMSO, 10 μL) with K-MOPS buffer (pH 7.2, 10 mL) to give a 10 μM solution in K-MOPS containing 0.1% DMSO.
NOTE: K-MOPS buffer consisted of 100 mM KCl and 10 mM 3-(*N*-morpholino)propanesulfonic acid (Mops) titrated to pH 7.2 with KOH.
2. Transfer an aliquot of the solution (V₁ L) into the same cuvette used in the photoreaction of the chemical actinometer. Irradiate the sample solution using the same setup as described in step 4.2.1.
3. Remove an aliquot (50 μL) from the irradiated solution periodically and analyze using HPLC.
4. Determine the irradiation time, in seconds, in which 90% of the starting material reacted (*t*_{90%}) by fitting plots of the time-dependent disappearance of the starting material.
NOTE: The absorbance of the irradiated sample must be maintained at <0.1 so that the inner filtering of the radiation can be neglected. It is desirable that the photolytic consumption of the starting material can be approximated by single-exponential decay so that there is no undesired secondary effect that interferes with the photolysis process.
5. Calculate the quantum yield of disappearance (Φ_{dis}) using the following equation²⁸:

$$\Phi_{\text{dis}} = 1 / (t_{90\%} \times I_0 \times \sigma_{350})$$
 where *t*_{90%} [s] is the irradiation time in which 90% of the starting material was consumed, *I*₀ [einstein cm^{−2} s^{−1}] is the number of moles of photons, and σ_{350} [cm² mol^{−1}] is the decadic extinction coefficient of the sample at 350 nm.
 NOTE: $\sigma_{350} [\text{cm}^2 \text{mol}^{-1}] = 10^3 \epsilon_{350} [\text{M}^{-1} \text{cm}^{-1}]$.

5. Targeting of a clickable caged compound with a HaloTag ligand

NOTE: Prior to use, maintain the HeLa cells in Dulbecco's modified Eagle medium (DMEM, low glucose, sodium pyruvate, l-glutamine) supplemented with 10% fetal bovine serum (FBS) containing 1% antibiotics (streptomycin sulfate, penicillin G, and amphotericin) at 37 °C and 5% CO₂.

1. Remove the medium and trypsinize the cells by treating with trypsin-ethylenediaminetetraacetic acid (EDTA, 1 mL) at 37 °C for 1 min. Add DMEM (4 mL) to the cells and re-suspend the cells by pipetting gently. Seed approximately 5×10^5 cells per dish into 35 mm glass bottom dishes in DMEM (2 mL) 24 h before transfection.
2. For four dishes, in a 1.5 mL microcentrifuge tube, dilute the plasmid DNA (pcDNA3-Halo-EGFR, 14 µg) in the reduced serum medium (700 µL). Separately, dilute the lipofection reagent (5 µL) in the reduced serum medium (150 µL) into each of four tubes and allow them stand at ambient temperature for 5 min.
3. Add a portion of the diluted plasmid DNA (150 µL) to each of the diluted lipofection reagent samples. Incubate at ambient temperature for 5 min.
4. After maintaining the cells at 37 °C and 5% CO₂ for 24 h, aspirate the DMEM and rinse the cells with phosphate-buffered saline (PBS, 2 mL). Add the reduced serum medium (1.5 mL).
5. Add the plasmid-lipofection reagent (150 µL) complex to each dish. Maintain the cells at 37 °C and 5% CO₂ for 48 h.
6. Aspirate the medium, add a portion of freshly prepared DMEM (1 mL) containing 2 µM paBhc-hex-FITC/Halo, and incubate the cells at 37 °C and 5% CO₂ for 30 min.
7. Aspirate the medium containing the caged compound and rinse the cells twice with PBS+ (1 mL per rinse) to remove any unbound compounds. Add the reduced serum medium (500 µL) and incubate the cells at 37 °C and 5% CO₂ for 30 min to remove the compounds that entered the cells.
NOTE: PBS+ is phosphate-buffered saline supplemented with 2 mM CaCl₂ and 1 mM MgCl₂.
8. Remove the medium and rinse the cells twice with PBS+ (1 mL). Add a portion of a medium (1 mL) that does not contain phenol red. Record fluorescence images by laser scanning confocal fluorescence microscopy.

6. Photomediated modulation of a kinase localization using a clickable caged compound

NOTE: Prior to use, maintain the CHO-K1 cells in Ham's F-12 medium supplemented with 10% FBS at 37 °C and 5% CO₂.

1. Prepare a 100× working solution (1 mM) of paBhc-AA (**5**) in DMSO.
NOTE: A 10 mM stock solution of the compound is prepared and stored in a freezer (-20 °C).
2. Seed approximately 5×10^5 cells per dish into 35 mm glass bottom dishes in DMEM (2 mL) 24 h before transfection.
3. Transfect CHO-K1 cells with a plasmid coding for GFP-DGKy 48 h before the uncaging experiments.
NOTE: Transfection is performed according to steps 5.2–5.5.
4. Replace the medium with a reduced serum medium (2 mL). Add the 100× paBhc-AA working solution (20 µL) and incubate the cells at 37 °C and 5% CO₂ for between 5 min and 1 h.
NOTE: The loading time depends on the compound employed.
5. Place the cells on the objective stage of an inverted fluorescent microscope equipped with a dual light source fluorescence illuminator.
6. Take a fluorescent image every 10 s. Irradiate the cells with 330–385 nm light through a microscope objective for an appropriate time. Alternatively, irradiate the cells with 405 nm light using a Xe lamp through flexible quartz fibers.
7. Continue to record fluorescent images for 10 min.

Representative Results

Clickable caged compounds of some biologically interesting molecules, including arachidonic acid and paclitaxel, were successfully synthesized (**Figure 1**)^{28,41}. Additional properties such as the water solubility and cellular targeting ability were introduced into paBhcmoc-PTX via the copper(I)-catalyzed Huisgen cyclization ("Click" reaction) (**Figure 2**). These clickable caged PTXs were then photolyzed to produce their parent PTXs upon irradiation at 350 nm (**Figure 3**), and the physical and photochemical properties of the clickable caged compounds are summarized in Table 1. The quantum yields of clickable caged compounds 2'-glc-paBhcmoc-PTX (Φ_{dis} 0.14) and paBhc-AA (Φ_{dis} 0.083) were more than twice those of conventional Bhc caged compounds 2'-Bhcmoc-PTX (Φ_{dis} 0.040) and Bhc-AA (Φ_{dis} 0.038)⁴³. In addition, an improved water solubility was observed for 2'-glc-paBhcmoc-PTX, which contains a glucose moiety.

In live cell experiments, the targeting of paBhc-hex-FITC/Halo to the cultured mammalian cells transiently expressing a fusion protein of a HaloTag protein and epidermal growth factor receptor (EGFR) was achieved successfully. Green fluorescence of the fluorescein moiety of paBhc-hex-FITC/Halo was observed on the cell membrane (**Figure 4**). Photo-mediated modulation of the subcellular localization of a kinase was achieved using a paBhc caged compound. The translocation of diacylglycerol kinase γ (DGKy) has been reported to be activated in the presence of arachidonic acid (AA)⁴⁴. CHO-K1 cells transiently expressing GFP-DGKy were treated with either AA or paBhc-AA (**5**). Addition of AA caused the modulation of the subcellular localization of DGKy (**Figure 5A,B**). Similar changes in the localization of DGKy were observed for the paBhc-AA-treated cells after exposure to UV light (**Figure 5C,D**).

Figure 1: Preparation of the clickable cage compounds.

(A) Reagents and conditions: a. ethyl 4-chloroacetoacetate/conc. H₂SO₄/rt/7 days/91% yield, b. 1 M HCl/reflux/3 days/97% yield. c. N-methylpropargylamine /HCHO/EtOH, then add (1) and heat at reflux for 17 h/79% yield. (B) Syntheses of the clickable cage amine, PTX, and arachidonic acid. [Please click here to view a larger version of this figure.](#)

Figure 2: Installation of functional units into clickable caged compounds.

(A) Synthesis of a water-soluble caged PTX via the copper(I)-catalyzed Huisgen cyclization. (B) Structures of clickable caged compounds containing the HaloTag ligand for cellular targeting. [Please click here to view a larger version of this figure.](#)

Figure 3: Photolysis of 2'-glc-paBhcmoc-PTX (6).

Samples (10 μ M) in K-MOPS solution (pH 7.2) were irradiated at 350 nm. (A) Typical HPLC traces for the photolysis of **6** (measured at 254 nm). Samples were analyzed at the specified irradiation time. (B) Time course for the photolysis of **6**. Blue circles show the consumption of **6**. The solid line shows the least-squares curve fit for a simple decaying exponential for **6**. Red squares show the yield of PTX. The error bars represent the standard deviation ($\pm$ SD). [Please click here to view a larger version of this figure.](#)

Figure 4: Fluorescence images of cultured mammalian cells incubated with paBhc-hex-FITC/Halo (8).

Cells transfected with pcDNA3-Halo-EGFR were incubated with a 2 μ M solution of compound **8** at 37 °C for 30 min. The images were obtained after repeated washing with PBS+. Mock-treated HEK293T cells (**A**: differential interference contrast (DIC) image and **D**: fluorescence image). HEK293T cells (**B** and **E**) and HeLa cells (**C** and **F**) transiently expressing Halo-EGFR (**B** and **C**: DIC images and **E** and **F**: fluorescence images). [Please click here to view a larger version of this figure.](#)

Figure 5: Fluorescence images after UV irradiation of the CHO-K1 cells incubated with Bhc caged arachidonic acid. CHO-K1 cells were transfected with a fusion protein DGK γ -EGFP.

(**A**) A fluorescence image of the transfected cells. (**B**) 100 s after the addition of a 10 μ M solution of arachidonic acid. (**C**) Cells were incubated with a 10 μ M solution of paBhc-AA (**5**) at 37 °C for 5 min. (**D**) 100 s after 20-s UV irradiation (330–385 nm). [Please click here to view a larger version of this figure.](#)

compounds	$\lambda_{\max}$ (nm) ^a	$\epsilon_{\max}$ (M ⁻¹ cm ⁻¹) ^b	ϕ_{dis} ^c	$\epsilon\phi_{\text{dis}}$ ^d	Solubility (μM) ^e
PTX					1.0
2'-Bhcmoc-PTX	340	10500	0.040	400	55
2'-paBhcmoc-PTX	359	9300	0.059	670	8.3
2'-glc-Bhcmoc-PTX	373	12300	0.14	1280	650
Bhc-AA	341	10800	0.038	390	
paBhc-AA	366	10300	0.083	750	

Table 1: Physical and photochemical properties of the clickable caged compounds.

a. Absorption maximum (nm), b. Molar absorptivity at $\lambda_{\max}$ (M⁻¹ cm⁻¹), c. Quantum yield of the disappearance of the starting materials at 350 nm, d. The product of molar absorptivity and the quantum yield of disappearance at 350 nm, e. The concentration of the saturated solution in K-MOPS (pH 7.2) ($\mu\text{g mL}^{-1}$).

Discussion

We previously developed Bhc caged compounds of various biologically active molecules that exhibit high photolytic efficiencies^{28,45,46,47}. With the aim of expanding the repertoire of Bhc caging groups, we also reported platforms of modular caged compounds that can be modified easily by the introduction of various functional units^{32,40,41}. The present protocol therefore represents a method for the synthesis of a clickable precursor of Bhc caging groups that can be modified via the copper(I)-catalyzed Huisgen cyclization. The synthesis of the clickable precursor, paBhcCH₂OH (**2**), was achieved via a four-step reaction sequence starting from the commercially available 4-bromoresorcinol (**Figure 1A**). The advantage of the present protocol is that no laborious purification steps (e.g., column chromatographic separations) are required.

As clickable precursor paBhcCH₂OH (**2**) can be used to mask various functional groups, clickable caged compounds of amines, alcohols, and carboxylic acids were synthesized using **2** as the precursor (**Figure 1B**). Amines were modified as their carbamates while alcohols were modified as their carbonates. In general procedures 1 and 2, CDI was used for the preparation of clickable carbamates, while 4-nitrophenyl chloroformate was used for the preparation of carbonates. As indicated by the reaction mechanism, both reagents can be used for the preparation of carbamates and carbonates. It should also be noted that the yield of the desired caged compound depends on the chemical structure of the molecule to be caged. Other examples can be seen in our previous reports^{28,30,33,48}.

Click modification was then performed using a slight modification of the reported procedure⁴⁹. The addition of tris(triazolylmethyl)amine-based ligands is necessary to obtain the desired products in good to high yields. Since a variety of azides are readily available both from commercial sources and from literature procedures, we can prepare various modular caged compounds with additional properties such as water solubility and cellular targeting ability (**Figure 2**).

The quantum yield of photolysis was then measured according to a reported procedure^{28,50}. **Figure 3** shows that the photolytic consumption of 2'-glc-paBhcmoc-PTX and the release of PTX were approximated by single-exponential decay and rise, respectively, suggesting no inner filtering of the radiation or undesired secondary effects. Improved photolysis quantum yields (Φ) and photolysis efficiencies ($\epsilon\Phi$) were observed for the clickable paBhc caged compounds compared to those of the previously reported Bhc caged compounds (**Table 1**)^{41,43}. Since the photolysis efficiencies ($\epsilon\Phi$) of Bhc caged compounds are more than one hundred times higher than those of 2-nitrobenzyl-type caged compounds⁴⁸, the marked improvement due to the presence of paBhc caging groups is clearly an advantage for this system.

As a proof-of-concept experiment, a hydrophilic moiety was introduced into 2'-paBhcmoc-PTX (**4**) and a cellular targeting ligand was introduced into compound **3** (**Figure 2**). The water solubility of 2'-glc-paBhcmoc-PTX was 650 times higher than that of the parent PTX (**Table 1**). Selective cellular targeting was achieved using a tag-probe system, and paBhcmoc-hex-FITC/Halo (**8**) bearing the HaloTag ligand was successfully targeted to the cell membrane of cultured mammalian cells expressing the HaloTag/EGFR fusion protein (**Figure 4**). Photo-mediated modulation of the subcellular localization of a kinase was also achieved using a clickable caged compound **5** (**Figure 5**).

In conclusion, we successfully demonstrated a method for the preparation of clickable platforms for photo-caged compounds of biologically interesting molecules that can be modified easily with additional properties, such as water solubility and a cellular targeting ability. Since the paBhc caging group can be used to prepare any molecules with modifiable functional groups, the application of the present protocol is not limited to the molecules described herein. Using a modular platform, namely the paBhc caging group, the desired caged compounds can be easily prepared, and their physical and chemical properties can be modulated via click modification.

Disclosures

We have nothing to disclose.

Acknowledgments

This work was supported by JSPS KAKENHI grant number JP16H01282 (TF), a Grant-in-Aid for Scientific Research on Innovative Areas "Memory Dynamism," and JP19H05778 (TF), "MolMovies."

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