Supramolecular interactions are non-covalent interactions between two or more molecules. Due to their moderate bond strength and reversibility, supramolecular interactions are usually responsive to external stimuli, such as light, pH, and redox. Supramolecular interactions can be used to construct stimuli-responsive materials for various applications. Photo-responsive supramolecular materials, for example, exhibit numerous biomedical applications, including drug delivery, photo-controlled protein adsorption and cell adhesion.

Currently, most photoresponsive supramolecules efficiently absorb UV light and are thus controlled by UV light. However, UV light is problematic in biomedical applications. Compared with UV light, red and near-infrared (NIR) light in the therapeutic window (600–900 nm) are better suited for biomedical applications because they cause less photodamage and can penetrate deeper into tissue. Therefore, red- or NIR-light-responsive supramolecules are desirable for biomedical applications.

One approach to design red- or NIR-light-responsive supramolecules is based on simultaneous two-photon absorption. However, this method is inefficient even when high-intensity femtosecond lasers are used. Another approach to achieve red- or NIR-light-responsive supramolecules is by coupling UV-responsive supramolecules with upconverting nanoparticles. However, photon upconversion requires high-intensity lasers that may cause overheating problems and photodamage to biomaterials.

Red-shifting the responsive wavelength of photoresponsive supramolecules is an approach to prepare intrinsic red- or NIR-light-responsive supramolecules. This approach should be more efficient than nonlinear optical processes such as two photon absorption and photon upconversion because red or NIR light are directly absorbed by supramolecules. To construct red- or NIR-light-responsive supramolecules, we may red-shift the responsive wavelength of UV-responsive supramolecules. It is well-known that trans azobenzene can spontaneously form supramolecular complexes with α- or β-cyclodextrin (CD). UV irradiation can induce trans-to-cis isomerization of azobenzene, which leads to the disassembly of the supramolecular complexes. Recently, new azo derivatives have been synthesized to red-shift the responsive wavelength of the azo group. In particular, some azo derivatives are responsive to red or NIR light. However, the feasibility of using the azo derivatives to construct red- or NIR-light-responsive supramolecules has not been explored. Moreover, no attention has been focused on constructing intrinsic red- or NIR-light-responsive supramolecules.

Here, we fabricate red-light-responsive supramolecules using tetra-ortho-methoxy-substituted azobenzene (mAzO) and β-CD (Scheme 1). Based on the red-light-responsive supramolecules, we prepared a responsive hydrogel that showed a gel-to-sol transition under red light irradiation. The hydrogel was used as a protein carrier. Red light was demonstrated to be able to...
control protein release from the hydrogel even in deep tissue. Compared with the conventional UV-responsive supramolecular interaction of azobenzene and CD, the new red-light-responsive supramolecular interaction reported in this paper is more suitable for biomedical applications.

### Experimental section

#### Materials

2,6-Dimethoxyaniline, 12-aminododecanoic acid and β-cyclodextrin (β-CD) were purchased from Alfa Aesar. 3,5-Dimethoxyaniline, sodium nitrite, di-tert-butyl dicarbonate (BOC), triethylamine, 4-(dimethylamino)pyridine (DMAP), poly(acrylic acid) (PAA) ($M_w = 450,000 \text{ g mol}^{-1}$), 6-bromohexanoic acid, pyridine and α-cyclodextrin (α-CD) were purchased from Sigma Aldrich. Trifluoroacetic acid, $N$-($3$-dimethylaminopropyl)-$N$-$0$-ethylcarbodiimide hydrochloride (EDC) and (benzotriazol-1-yloxy) tripyrrolidino-phosphonium hexafluorophosphate (PyBOP) were purchased from Carl Roth. 3A-amino-3A-deoxy-$\beta$-cyclodextrin hydrate ($\text{NH}_2-\beta$-CD) was purchased from TCI. The fluorescently labeled protein (Bovine Serum Albumin [BSA], Alexa Fluor® 680 conjugate) was purchased from Life Technologies. All the solvents (HPLC grade) were purchased from Sigma Aldrich and were used without further purification. Milli-Q water (resistivity: 18.2 MΩ cm) provided by a Sartorius Arium 611 VF Purification System was used throughout the research.

#### Methods

$^1$H and $^{13}$C nuclear magnetic resonance (NMR) spectra were recorded on a Bruker Avance 250 MHz spectrometer and on a Bruker Avance III 700 MHz system. The $^1$H-NMR experiments were acquired with a 5 mm BBI z-gradient probe on the 700 MHz Bruker AVANCE III system. For a proton spectrum, 128 transients were used with a 10 µs long 90° pulse and a 12 600 Hz spectral width together with a recycling delay of 5 s. The $^{13}$C NMR (176 MHz) measurements were made with a J-modulated (coupling constant of 145 Hz $^1$H–$^{13}$C was used) spin-echo for C-nuclei coupled to protons to determine the number of attached protons and proton decoupling during acquisition. The 90° pulse for carbon was 14.5 µs long and 16 000 scans were taken with a relaxation delay of 2 s. The temperature was regulated at 298.3 K and calibrated with a standard $^1$H methanol NMR sample using the topspin 3.2 software (Bruker). The control of the temperature was realized with a VTU (variable temperature unit) and an accuracy of ±0.1 K. The assignment was accomplished by $^1$H, $^1$H 2D NOESY (nuclear Overhauser effect spectroscopy). The spectroscopic
widths of the homo-nuclear NOESY experiments were typically 13,600 Hz in both dimension (f1 and f2) and the relaxation delay 2 s. The mixing time used in the 2D NOESY was kept at 350 ms.

The 2D $^1$H,$^13$C-HSQC (heteronuclear single quantum correlations via double inept transfer and phase sensitive using Echo/Antiecho-TPPI gradient selection with decoupling during acquisition) and the 2D $^1$H,$^13$C-HSQC-NOESY experiments were recorded with 2048 points in f2 and 256 points in f1 dimension. A NOESY mixing time of 350 ms were used for the HSQC-NOESY.

Before Fourier transformation, the data were zero filled to 1024 points in f1 and multiplied by a window function ($q$-sine bell or sine bell) in both dimension. The spectral widths were 30,000 Hz (236 ppm) for $^13$C and 9000 Hz (18 ppm) for $^1$H, both nuclei with a relaxation delay of 2.5 s. Proton and carbon spectra were referenced using the solvents signals ($^1$H: HDO = 4.80 ppm or DMSO-d$_5$H = 2.49 ppm and $^13$C: DMSO-d$_6$ = 39.5 ppm) as an internal standard. The carbon spectra in D$_2$O were calibrated against external TMS (tetramethylsilane) reference at 0 ppm.

Mass spectra (MS) were obtained using a VG instrument ZAB 2-SE-FPD. UV/vis absorption spectra were measured on a Lambda 900 spectrometer (Perkin Elmer). Fluorescence spectra were measured on a TIDAS II spectrometer (J&M).

Photoisomerization was induced by the LEDs with the wavelengths of 470 and 625 nm (device types LCS-0470-03-22 and LCS-0625-03-22, Mightex Systems). The output intensities of the LEDs were controlled by an LED controller (device type SLC-MA04-MU, Mightex Systems) and were calibrated by an optical powermeter (Model 407A, Spectra-Physics Corporation).

**Synthesis**

The syntheses of mAzo-NH$_2$, mAzo-Py, PAA-mAzo and PAA-$\beta$-CD were shown in Scheme 2. In brief, mAzo-NH$_2$ was synthesized by coupling 2,6-dimethoxyaniline and 3,5-dimethoxyaniline. To prepare water-soluble compounds with mAzo groups, mAzo-Py and PAA-mAzo were synthesized. PAA-$\beta$-CD were prepared by grafting NH$_2$-$\beta$-CD on PAA. The details of syntheses and characterization of these compounds were provided in the ESI.$^\dagger$

**Results and discussion**

**Photoresponse of tetra-ortho-methoxy-substituted azobenzene**

We studied red-light-induced isomerization of mAzo-NH$_2$ by $^1$H-NMR and UV/vis absorption spectroscopy (Fig. 1). The n-π* transition band of trans mAzo-NH$_2$ extended to the red-light region and was separated from that of cis mAzo-NH$_2$ (Fig. 1a and b). Thus, red light could excite trans mAzo-NH$_2$ and induce trans-to-cis isomerization.$^{4,5,6}$ Approximately 85% of the trans isomers switched to cis isomers after 625 nm red light irradiation for 30 min (60 mW cm$^{-2}$) (Fig. 1c). The cis mAzo-NH$_2$ could change back to trans isomers by heating or blue light irradiation.

**Red-light-responsive supramolecular interactions**

To investigate the supramolecular interactions between mAzo and CDs in aqueous solutions, the hydrophilic compound mAzo-Py was synthesized (Scheme 2a, Fig. S1, ESI$^\dagger$). Red light could still induce isomerization of mAzo-Py in the presence of CDs (Fig. S5, ESI$^\dagger$). The supramolecular interactions between mAzo-Py and $\beta$-CD in D$_2$O were studied by NMR spectroscopy (Fig. 2, Fig. S6 and S7, ESI$^\dagger$). In two-dimensional nuclear Overhauser effect spectroscopy (2D NOESY), the correlation between the protons of trans mAzo-Py (Proton 1 and 2) and the inner proton of $\beta$-CD (Proton f) demonstrated the
host–guest interaction between \( \text{trans mAzo-Py} \) and \( \beta\text{-CD} \) (Fig. 2, left). Moreover, adding \( \text{trans mAzo-Py} \) to the solution of \( \beta\text{-CD} \) caused the shift of the signals of the inner protons of \( \beta\text{-CD} \) (Fig. S6, ESI†), suggesting that \( \text{trans mAzo-Py} \) entered the cavity of \( \beta\text{-CD} \). Furthermore, the correlation between the carbons of \( \text{trans mAzo-Py} \) and the inner protons of \( \beta\text{-CD} \) also proved the host–guest interaction between \( \text{trans mAzo-Py} \) and \( \beta\text{-CD} \) (Fig. S7, ESI†). To study the interaction between \( \text{cis mAzo-Py} \) and \( \beta\text{-CD} \), \( \text{trans mAzo-Py} \) was converted to \( \text{cis mAzo-Py} \) by red light irradiation. No dipole–dipole (2D NOESY) correlation between the protons of \( \text{cis mAzo-Py} \) (Proton 3 and 4) and the inner protons of \( \beta\text{-CD} \) indicated that red light irradiation triggered the disassembly of the host–guest complex (Fig. 2, right). Thus, the supramolecular interaction between \( \text{mAzo-Py} \) and \( \beta\text{-CD} \) was red-light-responsive.

The association constants between the mAzo group and CDs were measured by \(^1\text{H-NMR} \) spectroscopy to quantify the supramolecular interactions using mAzo-Py or PAA-mAzo (Fig. S9–S13 and S16–S20, ESI†). The association constants between the normal azobenzene group and CDs were also measured for comparison using azobenzene-grafted PAA (PAA-Azo) (Fig. S21, ESI†). The association constants are listed in Table 1. For normal azobenzene, both \( \alpha\text{-} \) and \( \beta\text{-CDs} \) had high association constants with the \( \text{trans isomer} \) and low association constants with the \( \text{cis isomer} \). Thus, both the azobenzene/\( \alpha\text{-CD} \) and azobenzene/\( \beta\text{-CD} \) supramolecules were UV-responsive. For mAzo, \( \alpha\text{-CD} \) had low association constants with both the \( \text{trans} \) and \( \text{cis isomers} \), indicating that mAzo could not form a stable host–guest complex with \( \alpha\text{-CD} \). \( \beta\text{-CD} \) had a high association constant with \( \text{trans mAzo} \) and a low association constant with \( \text{cis mAzo} \). Therefore, the mAzo/\( \beta\text{-CD} \) supramolecule was red-light-responsive. The results in Table 1 show that the supramolecular interactions between mAzo and CDs were different from those between azobenzene and CDs. We interpreted that the differences in supramolecular interactions came from the different features of mAzo and azobenzene, such as the molecular sizes, shapes and hydrophilicities/hydrophobicities.

**Red-light-responsive supramolecular hydrogels**

To demonstrate the potential applications of the mAzo/\( \beta\text{-CD} \) interaction in stimuli-responsive materials, we fabricated supramolecular hydrogels. The gelators PAA-mAzo and PAA-\( \beta\text{-CD} \) were synthesized by grafting PAA with mAzo and \( \beta\text{-CD} \).
moieties, respectively (Scheme 2, Fig. S2 and S3, ESI†). Similar to mAzo-NH₂ and mAzo-Py, PAA-mAzo showed red-light-induced isomerization (Fig. 3a). Red light induced trans-to-cis isomerization of PAA-mAzo because the n–π* band of the trans isomer extended to the red-light region. Both blue light and heating can induce cis-to-trans isomerization. Approximately 95% of trans PAA-mAzo was obtained after cis PAA-mAzo was irradiated by blue light for 1 min (60 mW cm⁻²). Approximately 100% of trans PAA-mAzo could be obtained after cis PAA-mAzo was heated at 60 °C for 10 h. The isomerization can be switched reversibly (Fig. 3b).

Supramolecular hydrogels were prepared by mixing trans PAA-mAzo and PAA-β-CD in PBS buffer in the dark (ESI†). Gelation occurred because the mAzo/β-CD supramolecular complexes cross-linked the polymers (Fig. 4a). Red light irradiation (625 nm, 60 mW cm⁻²) for 30 min triggered the disassembly of the supramolecular complexes and a gel-to-sol transition (Fig. 4b). The gel-to-sol transition was reversible. Blue light irradiation or heating could induce the sol-to-gel transition. Our red-light-responsive hydrogel is better suited for biomedical applications than the conventional UV-responsive hydrogel1 because red light can penetrate deeper into tissue and cause less photodamage to biomaterials. Thus, it pushes photoresponsive hydrogels one step further toward biomedical applications.

Red-light-controlled protein release

The red-light-responsive supramolecular hydrogel was applied as a platform for controlled release of proteins (Scheme 3). Fluorescently labeled bovine serum albumin (BSA) was loaded in the supramolecular hydrogel (ESI†). Red light irradiation induced the collapse of the hydrogel and the release of BSA (Fig. 5a). The amounts of released BSA under different irradiation conditions were quantified by fluorescence spectroscopy (Fig. 5b, Fig. S22 and S23, ESI†). Only 9% BSA was released when the BSA-loaded hydrogel was kept in the dark for 60 min. Blue light irradiation (470 nm, 60 mW cm⁻²) for 60 min induced the release of only 15% BSA. Approximately 45% BSA was released after red light irradiation (625 nm, 15 mW cm⁻²) for 60 min. The release rate increased, and 83% BSA was released after higher-intensity (60 mW cm⁻²) red light irradiation for 60 min. We could also control the protein release kinetics “on-demand” via alternating red and blue light irradiation (Fig. 5c).

To test whether the supramolecular hydrogel maintained the red light responsiveness in deep tissue, a piece of pork...
tissue was placed between the light source and the sample (Fig. 6a). Red light (625 nm, 15 or 60 mW cm\(^{-2}\)) still induced protein release after passing through the tissue (Fig. 6b). Importantly, the light intensity used in this experiment (15 and 60 mW cm\(^{-2}\)) was an order of magnitude lower than the maximum permissible exposure of skin at 625 nm (200 mW cm\(^{-2}\)).\(^6\) This light intensity was also much lower than that for upconverting-nanoparticle-assisted photochemistry.\(^3\) Therefore, photodamage to the tissue was prevented (Fig. S24, ESI\(^\dagger\)).

Conclusions

In summary, we demonstrated the red-light-responsive supramolecular interaction between mAzo and β-CD. Supramolecular hydrogels were prepared by mixing the mAzo-functionalized polymer and the β-CD-functionalized polymer in an aqueous solution. Red light irradiation induced the disassembly of the mAzo/β-CD complexes and a gel-to-sol transition. The supramolecular hydrogel was used as a protein carrier. Red light could precisely control the protein release from the hydrogel. The red-light-responsive supramolecules exhibited several advantages over the conventional UV-responsive supramolecules\(^1\) for biomedical applications: (i) they could be controlled in deep tissue; (ii) controlling the supramolecules using low-intensity red light prevented photodamage to biomaterials. We expect the mAzo/β-CD interaction to be useful not only for constructing hydrogels but also for constructing red-light-responsive supramolecular surfaces, polymers, colloids and macroscopic assemblies. Additionally, we expect that not only azo derivatives but also other visible- or NIR-light-responsive compounds\(^7\) should be suitable for constructing supramolecular assemblies via rational design. We envision using red- or NIR-light-responsive supramolecules for drug delivery, protein therapy, optogenetics, and control adhesion and migration of cells.

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