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A Free-standing Molecularly Imprinted Photonic Hydrogels Based on β-Cyclodextrin for Visual Detection of L-Tryptophan

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Inverse opals of molecularly imprinted photonic hydrogels (CD-MIPhs) based on β-cyclodextrin were firstly elaborated by using the colloidal crystal template method. The film can be exfoliated from the substrate in H2O to get a free-standing photonic crystal film. L-Tryptophan imprinted polyacrylamide photonic hydrogels (PAM-MIPhs) and CD-MIPhs were fabricated in this paper. The PAM-MIPhs exhibited good sensing property to L-Trp and the redshift of the Bragg diffraction peak was 83 nm. Furthermore, the recognition to L-Trp can be improved by CD-MIPhs. The Bragg diffraction peak of CD-MIPhs redshifted 126 nm when the CD-MIPhs were immersed into a series concentration of L-Trp solution, and visually perceptible color change was clear. Moreover, enantioselective recognition of Trp enantiomers was accomplished by CD-MIPhs instead of PAM-MIPhs. In short, the resultant CD-MIPhs showed rapid response, high selectivity, high sensitivity and specificity towards the imprinted molecules in an aqueous environment.

Introduction

Molecular imprinting technology (MIT) is a well-established and facile technique used to synthesize molecularly imprinted polymers (MIPs) with specific binding nanocavities. Owing to the complementarity in shape and binding sites, the created nanocavities exhibit high selectivity towards the imprinted molecules, such as protein, resveratrol, bisphenol A. Due to the unique properties of high selectivity and structure predictability, MIPs have widely been applied as the recognition elements in analytical techniques. There are many studies on the highly sensitive and selective hyphenated techniques based on the combination of MIT and other traditional methods, such as chromatography techniques, solid phase extraction (SPE), electrochemical method and hydrogel photonic crystals (HPCs).

Particularly, Molecularly Imprinted Photonic Hydrogels (MIPhs) based on the combination of MIT and HPCs have attracted considerable interest. MIPhs is characterized by a 3D-ordered interconnected macroporous structure in which numerous recognition cavities are distributed in the thin polymer wall. The created nanocavities endow MIPhs with high specificity to analyse. Moreover, the molecular recognition events of MIPhs can be directly transferred into visually perceptible color change due to the shift of photonic bandgap. Furthermore, efficient mass transport and easier accessibility to the active sites can easily be accomplished through the hierarchical porous structure. Therefore, MIPhs have been widely developed for the detection of a large and diverse set of molecules.

However, the specific recognition of all these MIPhs to target analytes was mainly based on hydrogen bonding interactions between template molecules and functional monomers. Hydrogen bonding interactions could not allow the effective detection of hydrophobic molecules when MIPhs employed in aqueous environments. In order to address this issue, β-cyclodextrin (β-CD)-based polymer had been fabricated due to the host-guest interaction between β-CD and various organic hydrophobic molecules with suitable size. As far as we are aware, little attention had been paid to the use of β-CD for the fabrication of inverse opals of MIPhs. In addition, fabricated MIPhs are deposited on a substrate. It has been demonstrated that the adhesion of the thin film can affect the swelling behavior of thin hydrogel films and thereby their sensing property. It can be envisaged that this constraint would vanished for free-standing hydrogel films that were peeled from the substrate.

Herein, novel MIPhs were constructed by copolymerizing acrylamide (AM) with β-CD derivatives. The free-standing film was formed by exfoliation in deionized water directly. L-Tryptophan (L-Trp), the typical representative of organic molecules that can interact with β-CD through supramolecular interactions, was chosen as the model analyte. The recognition properties of MIPhs were demonstrated in detail. The proposed MIPhs can provide higher sensitivity, specificity in comparison with other MIPhs.

Experimental

Materials
Acrylamide (AM), L-Tryptophan (L-Trp), D-Tryptophan (D-Trp), L-Tyrosine (L-Tyr), L-Phenylalanine (L-Phe), N,N-dimethylformamide (DMF) and maleic anhydride (MAH) were purchased from Sinopharm Chemical Reagent Co., Ltd. (China). 2,2-Diethoxyacetophenone (DEAP) was obtained from TCI (Shanghai) Chemical Industry Development Co., Ltd. N,N'-methylene bisacrylamide (BIS) and β-cyclodextrin (β-CD) were purchased from Aladdin Chemistry Co., Ltd. (China) and β-CD was purified two times by recrystallization from water, and then dried under vacuum at 110 °C for 24 h prior to use. Styrene was purified by distillation under vacuum. All other affiliated chemicals not mentioned here were used without further purification. Glass slides (76.2x24.5x1 mm) for colloidal crystal growth were well-cleaned using acetone, anhydrous ethanol and double-distilled water in an ultrasonic bath in succession.

Synthesis of β-CD based reactive monomer (MAH-β-CD)

In order to integrate β-CD units with the polymer hydrogels, MAH-β-CD carrying five vinyl carboxylic acid groups was easily prepared as elucidated in reference. Typically, 0.005 mol of β-CD (5.6820 g) was added to 30 mL DMF, and then in which 0.05 mol of MAH (4.9030g) was dissolved. Afterwards, the clear mixture was heated at 80 °C for 10 h under mechanical agitation. The reacted mixture was cooled to room temperature, and then was dropwise added into 30 mL of chloroform. Finally, a yellow precipitate was obtained by washing with large amount of acetone at least three times, and drying in a vacuum oven at 80 °C for 48 h. FT-IR spectra of β-CD and MAH-β-CD were shown in supporting information Fig. S1. And the result of element analysis showed that five primary hydroxyl groups had reacted with MAH.

Preparation of polystyrene (PS) colloidal crystals

The monodispersed PS colloids were synthesized by emulsifier-free emulsion polymerization as described in our previous work. Then, corresponding PS colloidal crystal templates were fabricated by vertical deposition method. Briefly, PS colloidal particles suspensions with a weight concentration of approximately 1 % were poured into clean containers, then glass slides were inserted vertically into the above solution. Subsequently, the containers were placed in constant temperature humidity chamber. After the solvent was evaporated completely, PS colloidal crystals were deposited on the glass slide.

Preparation of free-standing molecularly imprinted photonic hydrogels (MIPHS)

In this work, two types of MIPHS, called CD-MIPHS or PAM-MIPHS respectively, were fabricated according to whether MAH-β-CD was involved in or not. Prior to fabrication of MIPHS, a homogeneous precursor of MAH-β-CD (0.75 mmol or 0 mmol), AM (25 mmol), BIS (1.5 mmol), DEAP (50 μL), and L-Trp (0.3 mmol) in 5 mL of phosphate buffer solution (pH 7.4) was deoxygenated under nitrogen for 10 min. “Sandwich structure” was firstly obtained by covering PS colloidal crystal templates with quartz glass plate of suitable size and immobilizing tightly. Based on this structure the precursors could infiltrate in the interstitial of PS colloidal crystal templates by capillary-atraction-induced force. Polymerization of the filled mixtures was conducted in an ice bath by irradiation at 254 nm for 2 h, and composite opal film was eventually formed on the glass slide. To obtain a free-standing photonic hydrogel, composite opal film was immersed in H2O. The hydrogel film would detach from the substrate in several minutes. The exfoliated composite opal film was immersed in dimethylbenzene to completely etch the PS colloids, and subsequently incubated in methanol/acetic acid (9/1, v/v) to elute L-Trp molecules. The corresponding non-molecularly imprinted photonic hydrogels (NIPHS) were also prepared under the same preparation conditions but in the absence of L-Trp.

Characterization

Surface morphologies of PS Colloidal Crystals and inverse opals of MIPHS were characterized by field emission scanning electron microscopy (SEM, Hitachi, S-4800). The Bragg diffraction wavelength of MIPHS that were immersed in the relevant solution of various concentrations was recorded by a miniature fiber optic spectrometer (FLA 4000+, China). The corresponding color changes were photographed using a common digital camera under a daylight lamp. UV spectra of the inclusion complexes of L-Trp and β-CD of different concentrations were measured on a TU-1901 UV-vis spectrophotometer. FT-IR spectra of β-CD and MAH-β-CD were recorded on a Nicolet Nexus 470 spectrometer in the optical range of 400–4000 cm⁻¹ by averaging 32 scans at a resolution of 4 cm⁻¹.

Results and discussion

Fabrication of free-standing L-Tryptophan imprinted photonic hydrogels

Three-step procedure well-established for the construction of MIPHS was shown in Scheme 1, including the preparation of PS colloidal crystals, the infiltration and polymerization of precursors in the interstitial of colloidal crystals, and the removal of the PS template and imprinted L-Trp molecule. It should be underlined that there is a remarkable improvement in the process of infiltration of precursors. While permeating of precursors into the PS interstitials, it was tilted by about 30° against the horizontal, and then precursors was dropped on the bottom edge of “sandwich structure” using a 5 μL pipette. Spontaneously, mixtures were slowly drawn into the space of the “sandwich structure” by capillary-atraction-induced force. Through polymerization, the polymer film was deposited on the substrate. To make the film easily remove from the substrate, the sequence of soaking in water and immersing into xylene must not be reversed. Only in this case the free-standing MIPHS can be acquired and the largest fabrication
size of the film can reach 2×1 cm.

Scheme 1. (a) Schematic illustration of the procedure for the preparation of inverse opals of MIPs and molecular structures of (b) β-CD and (c) L-Trp.

Fig. S1 exhibited the typical SEM photos of the adopted colloidal-crystal template and the resultant hydrogel inverse opal film with a thickness of about 100 μm. Highly ordered PS colloidal crystals had been prepared by vertical deposition of the PS colloids (210 nm) onto a glass substrate (Fig. S1a). After removal of PS colloids and L-Trp molecules, as it can be seen clearly in Fig. S1b, c, 3D highly ordered and interconnected macroporous arrays with a thin polymer wall of approximately 50 nm were obtained successfully. The hierarchical macroporous structure was essential for the easier accessibility to the specific nanocavities after etching PS colloid. Moreover, the photograph of exfoliated film was also given in Fig. S1d, showing the liberation of photonic hydrogels from the substrate successfully.

Recognition property of PAM-MIPs

Fig. 1 Optical response of PAM-MIPs (a) and PAM-NIPs (b)
upon soaking in L-Trp buffers. (c) Color change of PAM-MIPHs.

Amino acids would show different forms (cation, zwitterion and anion) in solutions with various pH. Minor structural difference of imprinting molecules in the fabrication of the MIPHs may result in different nanocavities in configuration. That would exert effect on the recognition properties of MIPHS. In order to maintain the uniformity of specific binding sites, existing state of L-Trp in precursors needs to be certain. Therefore, the pH of the precursors was constant 7.4 in our work. Furthermore, recognition properties of MIPHS were investigated in the analyze solution with the same pH (7.4). Fig. 1 displayed reflectance spectra of PAM-MIPHs and PAM-NIPHS upon soaking in buffer solutions with the L-Trp concentration at $10^{-5}$ M, $10^{-4}$ M, $10^{-3}$ M, $10^{-2}$ M, and $10^{-1}$ M. The Bragg diffraction peak of the PAM-MIPHs soaked in $10^{-5}$ M L-Trp was already redshifted by 83 nm relative to that of the original blank PAM-MIPHs in phosphate buffer solution, whereas only a slight shift (13 nm) was detected by the PAM-NIPHS. Furthermore, even upon exposure to $10^{-12}$ M L-Trp buffer solutions, the Bragg diffraction peak of PAM-MIPHs showed about 3 nm shift. Therefore, the detection limit of PAM-MIPHs approached $10^{-12}$ M.

Bragg diffraction wavelength ($\lambda_{max}$) is determined by the Bragg diffraction law\textsuperscript{13,35} as

$$\lambda_{max} = 2(d_{klm}/m)(D/D_0)(n_{eff}^2 - \sin^2 \theta)^{1/2}$$

where $d_{klm}$ is the interplanar distance of two neighboring layers along the incident direction, $n_{eff}$ is the effective refractive index, and $D/D_0$ is the degree of swelling of the gel where $D$ and $D_0$ denote the equilibrium diameters of the hydrogel at a given condition and in the reference state, respectively. For thin films of MIPHS, the top layers typically correspond to the (111) plane, so that $d_{111}=0.8165d$, where $d$ is the diameters of PS colloids. Thus, eq. 1 simplifies to give

$$\lambda_{max} = 1.633(d/m)(D/D_0)(n_{eff}^2 - \sin^2 \theta)^{1/2}$$

For specific MIPHS, $d=210$ nm, $m=1$, $\theta=0^\circ$. The effective refractive index ($n_{eff}$) can be expressed as

$$n_{eff} = [n_w^2(1-f) + n_m^2f]^{1/2}$$

in which $n_w$ and $n_m$ are the average refractive indices of the PAM walls (1.48) and medium H$_2$O (1.33), respectively. $f=\theta/74$ (fcc structure) is the filling factor of PS colloids. So the factor $n_{eff}$ affects $\lambda_{max}$ is $D/D_0$.

The change of $D/D_0$ was essentially attributed to the specific adsorption of L-Trp molecules to imprinting nanocavities. After adsorbing L-Trp molecules, concentration gradients of L-Trp between hydrogel and hydroly solution could be formed, and the more L-Trp molecules adsorbed, the greater concentration gradients formed. Then, water penetrated into hydrogel phase, resulting in the swelling of inverse opal. Therefore, the Bragg diffraction peak of PAM-MIPHs redshifted marked with the increase of L-Trp concentration. There are not specific nanocavities distributed in PAM-NIPHS. The slight redshift of the peak was due to the nonspecific adsorption of hydrogel to L-Trp molecules.

**Recognition property of CD-MIPHs**

The prerequisite for creating specific binding nanocavities in the polymer matrix lies in the formation of stable complex between monomers and template molecules L-Trp through strong molecular interactions during the molecular imprinting process. The stronger molecular interactions are, the more specific binding nanocavities would be acquired. Considering strong supramolecular interactions between β-CD and hydrophobic molecules, an appropriate amount of β-CD based reactive monomer (MAH-β-CD) was introduced to enhance the bonding in the pre-polymerization complex. Fig. 2a displayed the sensing behavior of the resultant CD-MIPHs with a series concentration of L-Trp ($10^{-10}$ M, $10^{-9}$ M, $10^{-8}$ M, $10^{-7}$ M, $10^{-6}$ M, and $10^{-5}$ M). In contrast with PAM-MIPHs (Fig. 1a), CD-MIPHs showed a higher sensitivity at the same L-Trp concentration. L-Trp at a concentration of $10^{-5}$ M induced a Bragg diffraction redshift of 126 nm from 525 nm to 651 nm. As shown in Fig. 2b, CD-MIPHs exhibited obvious color change. Encouraged by these promising results, the effect of the content of MAH-β-CD, AM and BIS on imprinting effect were investigated (Fig. S3). The molar ratio between the functional monomers (MAH-β-CD and AM) and cross-linker (BIS) should be balanced.

![Fig. 2](image-url)
be attributed to the increase of effective binding nanocavities and the strong supramolecular interaction between binding sites and L-Trp molecules. The inclusion interaction could be analyzed by the UV spectra of the inclusion complexes. Fig. 3a presented absorbance change of L-Trp upon addition of different concentrations of MAH-β-CD to 0.05 mM L-Trp solution. The absorbance intensity of the free L-Trp increased gradually with increasing concentration of MAH-β-CD. That may be due to more L-Trp formed inclusion complex with MAH-β-CD, resulting in the diminishment of free L-Trp in solution. According to the Benesi-Hildebrand method, the stoichiometry ratio and stability constant can be determined by the following equation

\[ \frac{A - A_0}{A_0} = \frac{\Delta \xi}{\left[ C_{CD}\right]_0} + \frac{\Delta \xi}{\left[ C_{CD}\right]_0} \times k \times \left[ MAH - \beta - CD \right]_0 \]

where \( A \) and \( A_0 \) are the absorbance of Trp at each MAH-β-CD concentration and in the absence of MAH-β-CD, respectively. \( k \) is the stability constant for inclusion complexation. [MAH-β-CD] and [C_{CD}] are the original concentrations of MAH-β-CD and L-Trp molecules, respectively, and \( n \) represents the stoichiometry ratio of the inclusion complexes. \( \Delta \xi \) is the differential molar extinction coefficient of L-Trp in the absence and presence of β-CD. Kong et al. have reported the the stoichiometry ratio and stability constant for the combination of β-CD and the Trp isomers. In theory the stoichiometry ratio would not be changed, and we assumed that \( n \) kept a value of 1 in this work, and a straight line is obtained (Fig. 3b), indicating formation of the 1:1 complexes between MAH-β-CD and the L-Trp. The interaction constant \( k \) could be calculated to be 233.0 M⁻¹, which was slightly less than the value (254.6 M⁻¹) of inclusion constant between β-CD and the L-Trp. The strong supramolecular interaction allowed the β-CD based reactive monomer competent during the molecular imprinting process.

Fig. 3 (a) UV spectra of MAH-β-CD-L-Trp inclusion complexes dependent on various concentrations of MAH-β-CD: (1) 0 mM, (2) 0.5 mM, (3) 1 mM, (4) 2 mM, (5) 3 mM, (6) 5 mM, (7) 7 mM, and (8) 10 mM. (b) Benesi–Hildebrand plot for the inclusion complexes.

Enantioselective recognition of L-Trp and D-Trp

Fig. 4 Optical response of (a) PAM-MIPs and (b) CD-MIPs upon soaking in D-Trp buffers. (c) Plots of Bragg diffraction peak shifts of PAM-MIPs to L-Trp (□), D-Trp (○) and CD-MIPs to L-Trp (△), D-Trp (▲).

\[ y = 0.1129x + 26.32 \]

\[ R^2 = 0.9979 \]
Fig. 4 showed the sensing behavior of the resultant PAM-MIPHs and CD-MIPHs with a series concentration of D-Trp. The response of PAM-MIPHs to D-Trp (Fig. 4a) was similar to that of PAM-MIPHs to L-Trp (Fig. 1a). That indicated that PAM-MIPHs could not distinguish stereoisomer of Trp. The shift of Bragg diffraction peak of CD-MIPHs with a series concentration of D-Trp was relatively slight compared with the result (126 nm) of Fig. 2a, indicating that CD-MIPHs possessed higher stereoselectivity. It can be envisaged that the stereoselectivity of CD-MIPHs are tightly related to the incorporation of β-CD. Specific binding nanocavities with particular shape, size and binding sites were involved in the PAM-MIPHs and CD-MIPHs. But owing to the tiny configuration difference between L-Trp and D-Trp, nanocavities of PAM-MIPHs were not competent for the enantioselective recognition of L-Trp and D-Trp. Assisted by the stereoselective discrimination of Trp enantiomers by β-CD, fabricated CD-MIPHs can easily differentiate Trp enantiomers. However, due to the presence of nonspecific binding sites during the molecular imprinting process, CD-MIPHs would adsorb a certain amount of nonimprinted molecules, resulting in the minor redshift (37 nm) of Bragg diffraction peak. Although nonspecific binding of CD-MIPHs led to the redshift of Bragg diffraction peak, the effect of molecular imprinting still dominated.

Selectivity of CD-MIPHs

According to above results, fabricated CD-MIPHs exhibited the better sensitivity and stereoselectivity. Besides, selectivity of CD-MIPHs towards L-Phe and L-Tyr was also investigated. The corresponding reflectance spectra were revealed in Fig. 5. Compared with the result of CD-MIPHs to L-Trp, slight optical redshifts of CD-MIPHs induced by L-Phe and L-Tyr took place under the same measurement conditions. The reason is believed to be that imprinted nanocavities with the fixed shape, size and interaction sites can bind much more L-Trp molecules than L-Tyr and L-Phe. These results indicated that fabricated CD-MIPHs can efficiently discriminate between template and the analog owing to the cooperative effect of shape, size, and interaction sites of the imprinted nanocavities. To further elucidate imprinting effect of CD-MIPHs, reflectance spectra of CD-MIPHs upon exposure to L-Trp buffers were shown in Fig. 5c. Probably due to the nonspecific adsorption, only a slight fluctuation in the Bragg peak was observed in contrast with the sensing behavior of CD-MIPHs. In summary, CD-MIPHs had the superior ability to selective recognize template molecules.

Response speed and recoverability of CD-MIPHs

Although the thickness (100 μm) of our free-standing CD-
MIPHs was thicker than reported obviously (2 μm)\textsuperscript{32,38}, fast response could still accomplished (Fig. 6a). It only needed 30 s to nearly reach its highest Bragg diffraction peak when CD-MIPHs were immersed into the 10^{-5} M L-Trp solution. The 3D interconnected macroporous structure was favorable for molecular diffusion, and the recognition sites that are situated at the surface or in proximity to the surfaces of the ultrathin polymer wall, were easily accessible. On the other hand, the swelling behavior of exfoliated film was no longer restricted by the substrates. Therefore, free-standing CD-MIPHs exhibited rapid response speed all the same. Moreover, the responded CD-MIPHs can be easily recovered by using methanol/acetic acid to remove rebinding molecules. Fig. 6b showed the recoverability of free-standing CD-MIPHs over five cycles between pure buffer and 10^{-9} M L-Trp solution. This indicated that CD-MIPHs have good recoverability, which makes it useful for practical applications. Moreover, the free-standing film can remain intact and maintain excellent recognition property even stored for half a year. Compared to other strategies\textsuperscript{59,40} concerning the recognition of Trp, CD-MIPHs overcome the shortcomings such as high cost and time consumption. However, although the sensitivity and selectivity of CD-MIPHs can retain high levels, quantitative detection of MIPHs still needs further research.

In summary, inverse opals of molecularly imprinted photonic hydrogels (CD-MIPHs) containing β-CD were successfully developed. The CD-MIPHs demonstrated higher sensitivity and better chiral recognition for Trp enantiomers. And CD-MIPHs also exhibited highly selectivity to analogous molecules. In addition, the thin film can be exfoliated from the substrate to produce an intact free-standing CD-MIPHs film. The peeled free-standing inverse opal film might provide great convenience for the application in the real-world environments. This study not only provided a method for the detection of L-Trp, but opened an avenue for the detection of many other molecules in aqueous environments. However, further investigations still needed to minimize non-specific adsorption.

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Molecularly imprinted photonic hydrogels fabricated by colloidal crystal template method have been exfoliated from substrate to get a free-standing film.