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# Thermally controlled biotransformation of glycyrrhizic acid *via* an asymmetric temperature-responsive polyurethane membrane†

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Separating a target product from a relatively complex bioreaction system is often difficult. In this work, a “smart” bioreaction system was developed by using the special characteristic of temperature-responsive polyurethane (TRPU). By combining solvent evaporation with a wet phase inversion technique, an asymmetric membrane consisting of an integral and dense skin layer supported by a porous sublayer was prepared from a thermally responsive polyurethane that experiences a sudden free volume increase upon heating through a phase transition temperature of 56 °C. Subsequently, the asymmetric TRPU membrane served as the carrier of an immobilized enzyme, wherein β-glucuronidase was multipoint-conjugated by using biotin and streptavidin on the porous sublayer. Then, the material-asymmetric TRPU membrane served jointly as the antennae as well as the actuator, which reversibly responds to temperature to switch (on–off) the access of the reactant glycyrrhizic acid (GL). Under the optimal temperature (40 °C) and pH (7.0) conditions, the immobilized β-glucuronidase contributed to almost 33% yield of glycyrrhetic acid 3-*O*-mono-β-*D*-glucuronide (GAMG) of the isolated counterpart for the same concentration of substrate (250 mg L<sup>-1</sup>) reaction for 24 h, while costing 1% of that of the isolated β-glucuronidase. Kinetic results showed that  $V_{\max}$  and  $K_m$  values were  $8.89 \times 10^3$  mg L<sup>-1</sup> and  $2.30 \times 10^3$  mg L<sup>-1</sup> h<sup>-1</sup>, respectively. The specific functional polymer-immobilized β-glucuronidase design serves as a bioreactor of GL into GAMG, as well as a separator deliberately irritated and controlled by temperature. This “smart” support material presents a potential facilitator for the separation of complex biotransformation reactions.

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## 1. Introduction

Synthetic polymers that exhibit a reversible response to temperature, pH, and light can be regarded as “smart” and hold great potential for applications in bioprocessing and pharmaceuticals.<sup>1–6</sup> These polymers that undergo changes in size, conformation, or hydrophobicity in response to variations in temperature or pH are gaining increased attention. In particular, there is increasing interest in their potential application ranges, including drug delivery, bioprocessing, gas or liquid transport, and leather industry.<sup>7–11</sup> Temperature-responsive polymers (TRPUs) prepared using various materials and

procedures are generally characterized by a segmented structure with a partially miscible fixed phase and a reversible phase favorably imported with a typical switching temperature. When the environmental temperature is above the phase transition temperature, the free volume in the polymeric matrix dramatically expands.<sup>12,13</sup> More advantageously, by manipulating the molecular structure or chemical composition, the transition of the reversible phase can also be interlinked to a temperature of interest for a particular application.

From a TRPU, an asymmetric TRPU (ASTRPU) membrane can be prepared using a wet phase transfer approach, which consists of a porous sublayer enabling its contribution to various potential applications. In our previous studies, an asymmetric membrane was successfully synthesized and used in transport, medical, and environmental applications.<sup>14–16</sup> In continuation, we intend to investigate its applications in the area of biotransformation by limiting the access of the substrate to the enzyme of interest, thereby controlling the reaction process.

Biotransformation is an effective means to obtain active components and the leading compounds of natural products in manufacturing biotherapeutic agents.<sup>17,18</sup> Glycyrrhizic acid (GL) and glycyrrhetic acid (GA) are important components

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extracted from the root of *Glycyrrhiza glabra* and have been widely used in traditional Chinese medicine. The biological actions of both compounds cover a wide spectrum, including anti-inflammatory, antiviral, antioxidative, and anticancer activities.<sup>19,20</sup> When GL is converted to GA, the intermediate product glycyrrhetic acid 3-*O*-mono- $\beta$ -D-glucuronide (GAMG) is often produced. GAMG exhibits even higher antitumor and anti-inflammatory activities than GA. The most commonly used approach to acquire these effective components is extraction, despite its lower efficiency and higher cost. As an alternative method, chemical transformation or biotransformation has become increasingly attractive, particularly biotransformation as it is endowed with the benefit of higher selectivity and requires less use of organic solvents. Hence, it is regarded as a sort of "green chemistry" or "green technology".<sup>21–24</sup> Obviously, the enzymatic activity of either the whole cell or an isolated enzyme dominates the reaction rate, such as hydrolysis, esterification, oxidation, reduction, and asymmetric synthesis.<sup>25,26</sup> As compared to the entire cell enzyme, the isolated enzymes contribute toward specific selectivity, although they may incur a higher cost. Therefore, the use of an immobilized enzyme is a considerably less expensive and convenient, as well as recyclable, means for the corresponding bioreaction. Among the general immobilized enzymes used, the catalytic-functions-(CFs)-immobilized enzyme is fairly promising, but few are obtained because of limitations of the carrier.<sup>27</sup>

In this work, a treated asymmetric polyurethane membrane served as a switch for biotransformation and a CFs-immobilized enzyme carrier because of its smart on-off characteristic. The multipoint immobilization using biotin and streptavidin referred to here has recently been regarded as the most rigid means of immobilization.<sup>28–31</sup>

## 2. Material and methods

### 2.1 Materials

A polycaprolactone diol with number-average molecular weight of 10 000 g mol<sup>-1</sup>, 4,4'-methylenebis(phenyl isocyanate) (MDI), 1,4-butanediol (BDO), and *N,N'*-dimethylformamide (DMF) were purchased from Sigma Chemical Co.  $\beta$ -Glucuronidase (EC.3.2.1.31, lyophilized powder from *E. coli*, 1 000 000–5 000 000 units per g), GL and GA (purity > 98%), and phosphate buffered saline (PBS) were purchased from Sigma Chemical Co. EZ-link Amine-PEG3-Biotin and EDC were purchased from Fisher, USA, and streptavidin was purchased from Invitrogen, USA. Hydrazide 4'-hydroxyazobenzene-2-carboxylic acid (HABA) was purchased from Pierce, USA.

PBS: one tablet dissolved in 200 mL of deionized water yields 0.01 M phosphate buffer, 0.0027 M potassium chloride, and 0.137 M sodium chloride at a pH of 7.4 and temperature of 25 °C.

### 2.2 Synthesis of TRPU

A 250 mL round-bottom, three-necked separable flask equipped with a mechanical stirrer, nitrogen inlet thermometer, and condenser with a drying tube was used as the reactor. The TRPU

resin was synthesized *via* solution polymerization in dimethylformamide (DMF) under an N<sub>2</sub> atmosphere *via* two steps. During the first step, poly(epsilon-caprolactone) (PCL) 10 000 and a two-fold molar ratio of MDI were charged into the reactor and stirred at 80 °C for 2 h to prepare an isocyanate-terminated prepolymer. The prepolymer was subsequently chain extended at 80 °C for another 2 h by successively adding a stoichiometric amount of MDI and chain extender, BDO. The final molar ratio of MDI to PCL10000 and BDO was maintained at 1 : 1 to yield a linear polymer. Occasionally, DMF was added into the reactor when the viscosity of the reaction mixture was too high. Finally, a viscous and transparent TRPU solution with 37.6 wt% solid content was obtained. The above synthesis procedure and molecular structure of TRPU are illustrated in Scheme 1.

### 2.3 Synthesis of ASTRPU

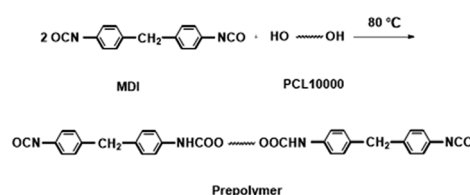
ASTRPU was prepared as per our previous work.<sup>14</sup> The reaction membrane was prepared using a wet phase inversion film formation process, *i.e.*, casting the TRPU solution on a release paper and then heating the membrane for 15 min at 50 °C (near its crystal temperature). Subsequently, the membrane was immersed in water to saturate. Then, an asymmetric membrane was obtained by washing with water to exclude the solvent and trace metal ions. Finally, the membrane was dried at 40 °C under vacuum conditions for 10 h. The physical and chemical parameters of ASTRPU are listed in Table 1.

### 2.4 Characteristic methods

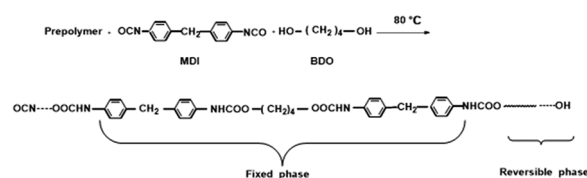
The skin and sublayer porous structure of the synthesized ASTRPU were scanned using a scanning electron microscope (SEM; JSM-6100 scanning microscope, JEOL, Japan).

Plasma (AutoGlow 100, Glow Research, USA) O<sub>2</sub> treatment was applied to activate the porous sublayer surface of the ASTRPU membrane and was immediately available to conjugate with EDC and biotin hydrazide. Using X-ray photoelectron spectroscopy (XPS; PHI Versaprobe II, Japan), the elemental composition

First step:



Second step:



Scheme 1 Synthesis procedure and molecular structure of TRPU.



Table 1 Physical and chemical parameters of ASTRPU

Fix phase content <sup>a</sup> (wt%)	Area <sup>b</sup> (m <sup>2</sup> g <sup>-1</sup> )	Temperature <sub>on</sub>	Temperature <sub>off</sub>	M <sub>n</sub> (g mol <sup>-1</sup> )	M <sub>w</sub> (g mol <sup>-1</sup> )	-NCO/-OH (mole ratio)
37.6	4.70	56 °C	<25 °C	4.47 × 10 <sup>4</sup>	6.46 × 10 <sup>4</sup>	1.2

<sup>a</sup> The fixed phase content was calculated as the mass fraction of MDI and BDO. <sup>b</sup> Measured as BET surface area.

changes in the ASTRPU membrane before and after the plasma treatment were explored. Peak analysis was performed using CasaXPS version 2.1.9 software, in which the Shirley background was subtracted from all the spectra prior to fitting.

## 2.5 Analytical methods

To determine the transformational temperature of the synthesized ASTRPU, a differential scanning calorimeter (DSC; Discovery, USA) was used. The temperature range was measured from -80 °C to 270 °C at an increasing temperature rate of 10 °C min<sup>-1</sup>. Triplicate scanning was performed; finally, an average temperature was obtained.

To measure the surface area of the ASTRPU membrane, the Brunauer-Emmett-Teller (BET; ASAP2020, USA) method was applied. The sample was pretreated by drying for 12 h. Then, the sample was measured and analyzed using the instrument software.

To test the molecular weight, the gel permeation chromatography (GPC) method was used.

To determine the effect of biotinylation on the structure of  $\beta$ -glucuronidase, circular dichroism (CD; Jasco J-1500, Japan) was used. The spectra were measured using a 0.1 cm path length cell and recorded at 25 °C in the range of 250–195 nm at a scanning speed of 50 nm min<sup>-1</sup>, step resolution of 0.1 nm, bandwidth of 1 nm, response of 4 s, and sensitivity of 200 mdeg.

To measure the bioreaction of GL, the product GA and the intermediate product GAMG were analyzed using a high-pressure liquid chromatography system (HPLC; Waters, 2487 series, MA, USA). The column was Extend C<sub>18</sub> (150 × 4.6 mm; I.D.: 5  $\mu$ m) with a flow rate of 1.00 mL min<sup>-1</sup>, detection wavelength of 254 nm, and at ambient temperature. The solvent system consisted of methanol and 3.6% acetic acid at a ratio of 85 : 15 (v/v).

To verify the degree of biotinylation, an ultraviolet visible spectrometer (UV-vis; Thermo 2010, USA) was used with HABA assay.

All the experiments were performed at the Massachusetts Institute of Technology, USA.

## 3. Results and discussion

### 3.1 Characteristics of ASTRPU membranes

DCS analysis (Fig. 1) reveal that the prepared ASTRPU undergoes a sharp phase transition at 56 °C. The BET specific area of the membrane was 4.70 m<sup>2</sup> g<sup>-1</sup>. Since the porous sublayer would make direct contact with the reactant GL and  $\beta$ -glucuronidase, the morphology of the sublayer surface was evident in Fig. 2a. Contrary to the skin layer surface (Fig. 2c), the entire sublayer appeared to be replete with interconnected

perforations. The SEM image involving the morphologies of the cross-section are shown in Fig. 2b. Fig. 2 shows that the abundant porous structure of the sublayer was available for the following enzymatic conjugation.

### 3.2 Switching or “on-off” characteristics of ASTRPU

The microstructure of the multiple finger-like pores in the porous sublayer indicated a larger area for  $\beta$ -glucuronidase loading. The transportability of TRPU was determined *via* the concentration change in GL initiated by the temperature change. By repeatedly switching the release assembly between two shaking water baths at different temperatures, the dynamic transport behaviors of GL through the ASTRPU membrane over four cycles of temperature oscillations between 25 °C and 45 °C were investigated, and the results are shown in Fig. 3.

As shown, the transport of GL through the ASTRPU membrane abruptly accelerated upon heating to 45 °C and then leveled off once the temperature reached 25 °C. Over four cycles of temperature oscillations, the on-off switching transport of GL was maintained, indicating good reproducibility and stability as the response to temperature.

### 3.3 Plasma treatment of ASTRPU membranes

To facilitate the conjugation of  $\beta$ -glucuronidase onto the membrane surface, the membrane surface was first treated by O<sub>2</sub> plasma (AutoGlow 100, Glow Research, USA) to alter the hydrophobic surface into a hydrophilic one. The compositional change in the porous sublayer was illustrated using XPS (PHI 5000 VersaProbe II, Japan) data obtained from the MIT Center for Materials Science and Engineering. Peak analysis was performed using the CasaXPS version 2.1.9 software with the Shirley background subtracted from all the spectra prior to fitting. The HR envelopes were analyzed and peak-fitted after

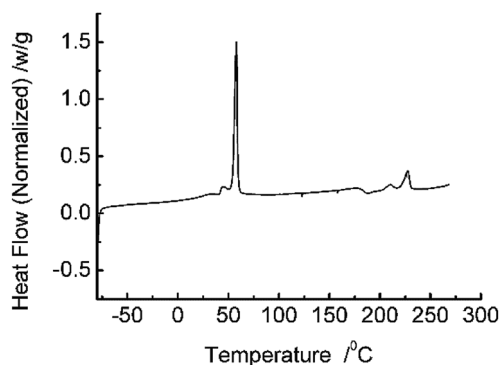


Fig. 1 DSC curve of synthesized TRPU.



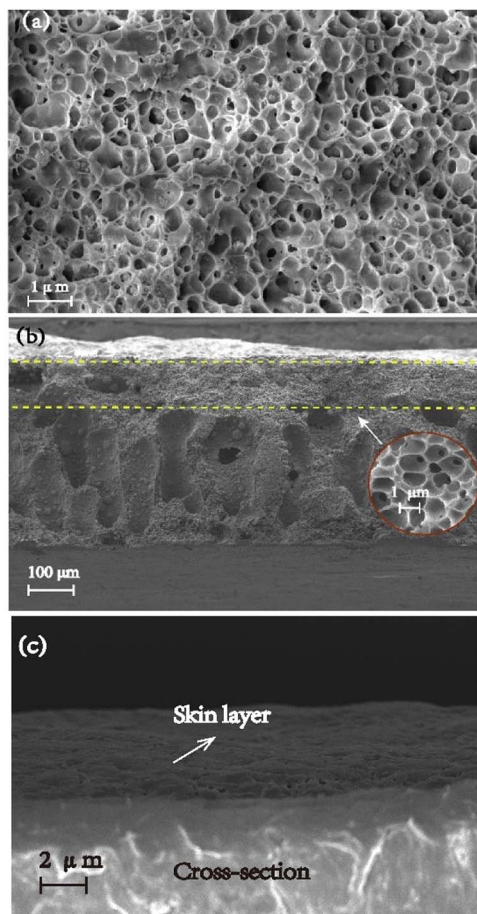


Fig. 2 SEM images of the morphologies of (a) porous sublayer surface; (b) cross-section; (c) skin layer surface of ASTRPU.

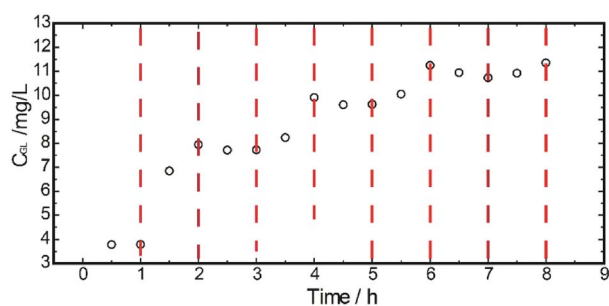


Fig. 3 Accumulative release profile of the ASTRPU membrane.

the subtraction of the Shirley background using the Gaussian-Lorentzian peak shapes obtained from the CasaXPS software package. The chemical compositions of the porous surface of ASTRPU before and after plasma O<sub>2</sub> treatment are listed in Table 2. It is obvious that the most significant change is the O content, while the N content seldom changes. As shown in Fig. 4, after treatment by O<sub>2</sub> plasma, the asymmetrical membrane became activated with the additional introduction of -C-O and -C=O groups (binding energy, BE: 286.5 eV and 289.2 eV, respectively). The O 1s lines also show an increase in -C-O (BE: 533.2 eV).<sup>32</sup>

Table 2 Chemical compositions of ASTRPU membrane before and after O<sub>2</sub> plasma treatment

Peak	Untreated ASTRPU		Plasma treated ASTRPU	
	Position BE <sup>a</sup>	Atomic (eV) conc. %	Position BE <sup>a</sup>	Atomic (eV) conc. %
O 1s	532.9	20.44	532.0	27.72
C 1s	400.4	79.24	400.0	68.29
N 1s	285.0	4.31	285.0	3.99

<sup>a</sup> BE is the binding energy.

### 3.4 Biotinylated ASTRPU and β-glucuronidase

Immediately, the ASTRPU membrane and the target β-glucuronidase were both biotinylated using EZ-link Amine-PEG<sub>3</sub>-Biotin and EDC following the Fisher protocol. The biotinylated β-glucuronidase was quantified using a Pierce Biotin Quantitation Kit supplied by Thermo Scientific. The kit contained the reagent HABA, which enables a rapid estimation of the mole-to-mole ratio of biotin to protein. Because biotin is a relatively small molecule, it can be conjugated to many proteins without altering the biological activity of the protein. A protein can be conjugated with several biotin molecules, each of which can bind one molecule of avidin, thereby considerably increasing the sensitivity of many assays. The bond formation between biotin and avidin is rapid; once formed, it remains unaffected by the most extremes pH values, organic solvents, and other denaturing agents. With regard to quantitative analysis,

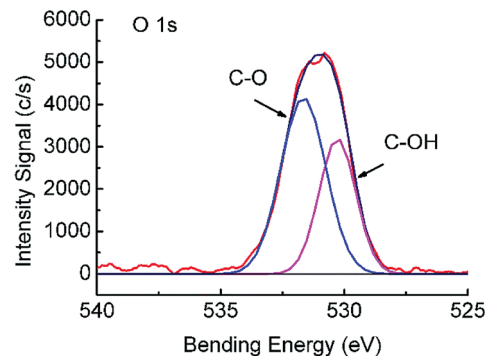
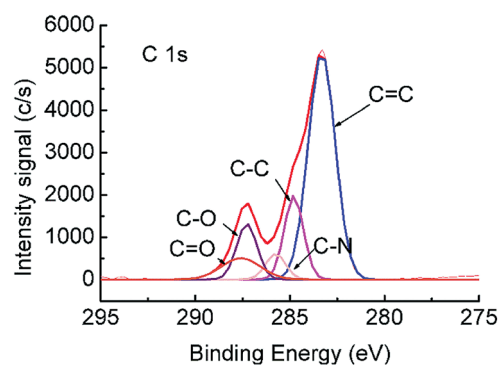


Fig. 4 XPS spectra of the C 1s and O 1s peaks of the O<sub>2</sub>-plasma-treated asymmetric TRPU. Treatment time: 1 min.



a solution containing biotinylated protein was added to a mixture of HABA and avidin. Because of the higher affinity toward avidin, biotin displaces the HABA and the absorbance at 500 nm proportionately decreases. Using this method, an unknown amount of biotin present in a solution can be quantified in a single cuvette by measuring the absorbance of the HABA–avidin solution before and after the addition of the biotin-containing sample. The change in absorbance is related to the amount of biotin in the sample *via* the extinction coefficient of the HABA–avidin complex.

### 3.5 Conjugate $\beta$ -glucuronidase onto the ASTRPU membrane

A schematic of the immobilization process is shown in Fig. 5. The quantitative number of both biotinylated and isolated  $\beta$ -glucuronidase conjugated to the porous sublayer membrane was determined using the HABA assay:  $0.31 \text{ mg cm}^{-2}$  and  $1.50 \text{ mmol mL}^{-1}$ , respectively. To the best of our knowledge, such an approach of immobilized  $\beta$ -glucuronidase has not yet been reported. The means of immobilizing  $\beta$ -glucuronidase presented here was that of multipoint crosslinking (one streptavidin molecule offers four binding sites for trapping) and it behaves rather rigidly than that in a traditional binding approach.<sup>28</sup> To determine the effect of biotin on the structure of  $\beta$ -glucuronidase, a circular dichroism (CD; Jasco J-1500, Japan) was used. Solutions of biotinylated and original  $\beta$ -glucuronidase were prepared at a concentration of 0.50 and  $0.49 \text{ mg mL}^{-1}$  in PBS, respectively. The structural changes in the biotinylated  $\beta$ -glucuronidase was established by comparing the CD spectra. Typically, three scans were accumulated and subsequently averaged.

### 3.6 Comparison of the before and after structures of biotinylated $\beta$ -glucuronidase

CD was used to monitor the  $\beta$ -glucuronidase structure's alteration, as shown in Fig. 6. Thus far, we know that the far-UV spectra for isolated  $\beta$ -glucuronidase and biotinylated  $\beta$ -glucuronidase exhibited similar spectra with strong negative ellipticity near 220 nm, and most of the spectra of the biotinylated  $\beta$ -glucuronidase overlapped with the spectra of the original type, indicating that the secondary structure of the biotinylated  $\beta$ -glucuronidase did not change during the course of biotinylating.

### 3.7 Reaction of GL into GAMG on biotinylated $\beta$ -glucuronidase ASTRPU membranes

With regard to isolated  $\beta$ -glucuronidase, under suitable reaction conditions, namely, a pH of 7.0 (PBS modulated) and

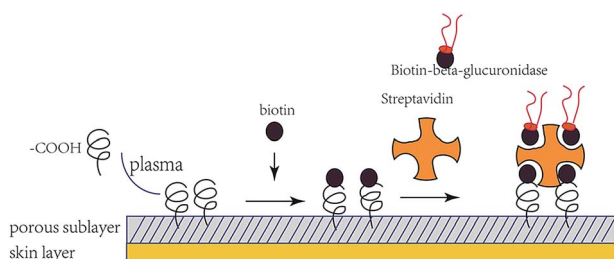


Fig. 5 Schematic of ASTRPU– $\beta$ -glucuronidase binding.

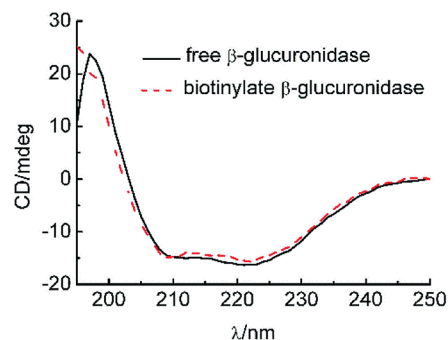


Fig. 6 CD graphs for isolated and biotinylated  $\beta$ -glucuronidase.

a temperature of  $35 \text{ }^\circ\text{C}$ , the hydrolysis of GL catalyzed by  $\beta$ -glucuronidase served as the probe model reaction to investigate the integral design of the temperature-responsive ASTRPU– $\beta$ -glucuronidase bioreaction assembly. The reaction occurred in a circular membrane surface (I.D.:  $0.68 \text{ cm}$ ; area:  $1.45 \text{ cm}^2$ ). One microliter of  $100 \text{ mg mL}^{-1}$  GL solution was added into a  $1.5 \text{ mL}$  bottle, and the porous, sublayer ASTRPU membrane served as the cover with binding  $\beta$ -glucuronidase apposite to the GL solution (*i.e.*, on the outside of the cover) and the edges of the membrane were thoroughly sealed to prevent leakage. This assembly was then hung in an inverted manner in a  $50 \text{ mL}$  conical tube containing  $1.2 \text{ mL}$  of PBS (pH 7.0) in advance. Thereafter, the  $50 \text{ mL}$  conical tube was incubated in a shaker at a certain temperature for 24 h. The products of the hydrolysis reaction in the PBS were analyzed using HPLC. The HPLC results showed that a different amount of GAMG was produced as a function of temperature, wherein the irradiated membrane released different concentrations of GL to access the  $\beta$ -glucuronidase conjugated in the porous sublayer. Fig. 7 shows the detailed results of such a tendency. The maximum GAMG product concentration was obtained at a temperature of  $50 \text{ }^\circ\text{C}$ . At a higher temperature, (*e.g.*,  $55 \text{ }^\circ\text{C}$ ), less GAMG was produced, which might have been caused by a decrease in or loss of enzymatic activity as compared to that at  $50 \text{ }^\circ\text{C}$ . Within the model reaction, the ASTRPU membrane played the role of a switch for GL approaching the  $\beta$ -glucuronidase irradiated by the temperature alteration. To investigate the kinetic behavior of the system, the reaction assembly was incubated in a shaker

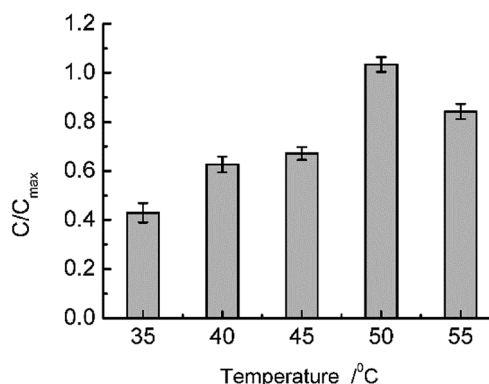


Fig. 7 Temperature-dependent GL biotransformation on an immobilized ASTRPU membrane.



at 40 °C for 24 h. The Michaelis constant,  $K_m$ , and the maximum reaction rate,  $V_{max}$ , were measured using the Lineweaver–Burk plotting method as  $8.89 \times 10^3 \text{ mg L}^{-1}$  and  $2.3 \times 10^3 \text{ mg L}^{-1} \text{ h}^{-1}$ , respectively. For isolated  $\beta$ -glucuronidase, the  $K_m$  and  $V_{max}$  values were  $131.07 \text{ mg L}^{-1}$  and  $1.64 \times 10^2 \text{ mg L}^{-1} \text{ h}^{-1}$ , respectively.  $K_m \gg C_{[GL]}$  indicated that both the isolated and immobilized enzymatic reactions were at a first-order magnitude. Although the  $K_m$  value for the immobilized version was greater than that for the isolated version (representing less affinity ability toward the substrate), the  $V_{max}$  value was nearly 10 times that of the isolated version, indicating that more product was produced more rapidly. Under the optimal temperature (40 °C) and pH (7.0), the immobilized ASTRP- $\beta$ -glucuronidase contributed to almost 33% of the GAMG of the isolated counterpart for the same concentration of substrate (*i.e.*,  $250 \text{ mg L}^{-1}$ ) for a reaction time of 24 h. However, the  $\beta$ -glucuronidase used was just 1% that of the isolate. The lost yield could form a considerable compromise given the lower consumption of  $\beta$ -glucuronidase.

## 4. Conclusions

In summary, from the perspective of controlling the bioconversion process, or alternatively, to obtain intermediate products of interest (*e.g.*, GAMG) in this work, the characteristic of TRPU was used. A catalytic bioreaction assembly coupled with the smart “on–off” switch was designed. The  $\beta$ -glucuronidase multipoint was rigidly immobilized onto ASTRP- $\beta$ -glucuronidase via biotin and streptavidin. The ASTRP- $\beta$ -glucuronidase reactor succeeded in controlling the access of the reactant GA toward  $\beta$ -glucuronidase by deliberately alternating the temperature, thereby controlling the bioconversion.

The developed reactor exhibited the potential advantage of obtaining important intermediate products or some of the steps during the reaction. Meanwhile, the reactant, product, and catalytic enzyme used were independent of each other, benefiting from the separation. Moreover, in addition to the immobilized enzyme that could be repeatedly used in most of the reaction system, the reactant could be repeatedly used in this system, which seldom occurs in most cases.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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