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Characterization of immobilized tyrosinase – an enzyme that is stable in organic solvent at 100 °C†

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Tyrosinase is a copper-containing enzyme present in plant and animal tissues, which catalyzes the production of melanin and other pigments. In organic solvent, tyrosinase can convert *N*-acetyl-L-tyrosine ethyl ester (insoluble in aqueous) to a derivative of L-dopamine (a drug used for the treatment of Parkinson's disease). Thus, the performances of tyrosinase in organic solvent have attracted scientific attention since 1980. In this work, we investigated the stability of immobilized tyrosinase at high temperature in anhydrous organic solvent. Triethylaminoethyl cellulose (TEAE-Cellulose) performed the best out of six immobilization platforms. The dry immobilized tyrosinase became extremely thermostable in organic solvent, and the half-life of the dry immobilized tyrosinase in organic solvent is strongly related to the polarity of the organic solvent than their log *P* value. The immobilized tyrosinase loses its activity instantaneously in aqueous solution at 100 °C, but it keeps enzymatic activity within 10 min in hydrophilic methanol and over one month in hydrophobic hexane (log *P*: 4.66, non-polar) even incubating at 100 °C. This research provides valuable information for the design of new biocatalysts.

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Introduction

Mimicking tyrosinase has been a longstanding goal of industrial and academic chemists.¹ Tyrosinases (polyphenol oxidase, EC 1.14.18.1), copper-containing oxidoreductases, can enable aerobic oxidation of relatively inert C–H bonds, which remains difficult for synthetic catalysts. Most of the synthetic catalysts can be stable in most of the organic solvents at high temperature (mostly higher than 100 °C). The behavior of tyrosinase in organic solvents at high temperature is very interesting for mimicking tyrosinase. Tyrosinase is an enzyme with monophenolase and diphenolase activity.^{2–4} Since the 1980s, innovative research has shown that tyrosinase can function well in anhydrous media with minimal water, pioneered mainly by Klibanov *et al.*^{5,6} This research remained of broad interest in the last century and mushroom tyrosinase has been utilized as an ideal enzyme by many researchers to study the functioning of enzymes in organic media.^{7,8} The enzyme regioselectivity and

enzymatic catalysis in organic media have been systematically investigated,^{9,10} but there is no report about the thermostable behavior of tyrosinase in different pure organic solvents at high temperature.

To study the thermostable behavior of tyrosinase,^{11,12} the enzyme needs to be incubated in different organic solvents and undergo testing of its activity in the same organic solvent. In this way, the enzyme must be easy to reuse.^{13,14} As previously reported, immobilization techniques are very efficient method to recycle the enzyme.^{15–17} The enzyme immobilized on a solid carrier can make the enzyme reusability easier.^{18,19} It is one of the key steps to test the thermostability of immobilized enzymes.^{20–22} From previous reports,^{23,24} the adsorption method keeps the original structure of enzyme,^{25–27} but encounters issues with enzyme leaching. Compared with the covalent bonding,^{28,29} entrapment,^{30–32} copolymerization^{33,34} and encapsulation methods,^{35–37} the conditions of the adsorption method are much milder than the other methods. Furthermore, the glass beads,^{38,39} graphene,^{40,41} magnetic nanoparticles⁴² and metal–organic frameworks (MOFs)^{43,44} were chosen as the supports to immobilize the enzyme in previous study. In this study, the biocompatible polymers (*i.e.* cellulose) were utilized as the ideal matrixes due to their low cost, and renewable and biodegradable nature.^{45–47} Besides, we systematically studied different polymers (*i.e.* cellulose matrixes, Pluronic F68 and poly(styrene-*co*-divinylbenzene)) and traditional glass beads as enzyme support.⁴⁸

After optimizing the enzyme immobilization material, the organic solvent effect, the position of substituent group and the

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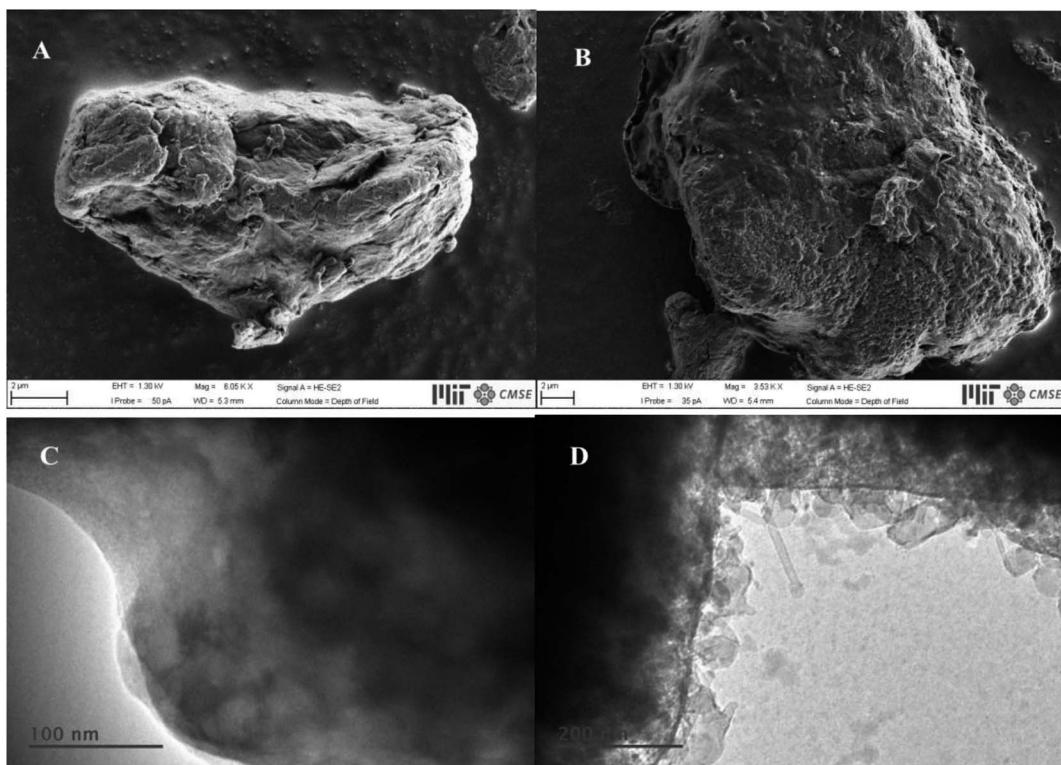


Fig. 1 SEM of (A) triethylaminoethyl cellulose (TEAE-Cellulose) and (B) TEAE-Cellulose with tyrosinase; the TEM of (C) triethylaminoethyl cellulose (TEAE-Cellulose) and (D) TEAE-Cellulose with tyrosinase.

strength of electron-donating substituent group were studied. Furthermore, the activities of immobilized tyrosinase were tested in various hydrophilic and hydrophobic organic solvents at high temperature. This study provided more reasonable and systematical explanation of tyrosinase in organic solvents.

Experimental section

Materials

p-Cresol, *p*-chlorophenol, *p*-methoxyphenol and other chemicals obtained from Sigma Aldrich (USA) were of the highest purity available. Mushroom Tyrosinase (50 kU, powder) was purchased from Sigma Aldrich (USA) as a solid with a specific activity of 2430 unit per mg (1 unit is defined as the enzyme activity resulting in an increase in absorbance at 280 nm of 0.001 at pH 6.5 at 25 °C in a 3 mL reaction volume containing L-tyrosine). Poly(styrene-*co*-divinylbenzene), glass beads, Pluronic F68, cellulose, carboxymethyl cellulose (CM-Cellulose) and triethylaminoethyl cellulose (TEAE-Cellulose) were also purchased from Sigma Aldrich (USA).

Instrumentation

Scanning electron microscope (SEM) images were obtained through high-resolution SEM (Zeiss Merlin, Germany) with a resolution of 0.8 nm at 15 kV and 1.4 nm at 1 kV. Transmission electron microscope (TEM) images were obtained from FEI Tecnai Multipurpose TEM (ThermoFisher Scientific, USA). UV-Vis spectrum was obtained by a UV-Vis Nanodrop 2000C spectroscopy (ThermoFisher Scientific, USA).

Preparation of tyrosinase and immobilization onto TEAE-Cellulose

Tyrosinase (6 mg) was dissolved in 0.3 mL of 50 mmol L⁻¹ phosphate buffer (pH 7.0), and then 60 mg of cellulose was added. The sticky mixture was spread on a watch glass and left to dry at room temperature. Tyrosinase immobilization onto CM-Cellulose or other materials was prepared following a similar protocol as tyrosinase immobilization onto TEAE-Cellulose.

UV-Vis spectrophotometry detection of the tyrosinase-catalyzed oxidation product

The time course for tyrosinase-catalyzed oxidation of *p*-cresol in organic solvents was measured by the following procedure: *p*-cresol was dissolved in a given solvent (its concentration: 50 mmol L⁻¹). 1 mL of *p*-cresol (50 mmol L⁻¹) in organic solvent was added into a 5 mL round-bottom flask, followed by addition of 5 μ L 50 mmol L⁻¹ phosphate buffer (PBS, pH 7.0). Then, the immobilized tyrosinase was added. Subsequently, the suspension was stirred by magnetic stirrers at 250 rpm at 25 °C. Aliquots of the liquid were removed and their absorption spectrum was recorded in the range 400–600 nm. The detection procedure of other substrates in a given solvent was the same as *p*-cresol in a given solvent.

Stability of tyrosinase in organic solvent at high temperature

The immobilized tyrosinase was transferred into a round bottom flask with anhydrous organic solvent under stirring at

100 °C. After 1 hour, the anhydrous organic solvent was removed, and then 1 mL of *p*-cresol in toluene with 5 μ L 50 mmol L⁻¹ PBS (pH 7.0) was added. The detection procedure was the same as the time course for tyrosinase-catalyzed oxidation of *p*-cresol in a given solvent.

Results and discussion

Physical characterization of TEAE-Cellulose and tyrosinase immobilization onto TEAE-Cellulose

Fig. 1B displays the SEM image of tyrosinase immobilization onto TEAE-Cellulose as compared to the original TEAE-Cellulose (Fig. 1A). There was a significant change in the morphology between the original TEAE-Cellulose and tyrosinase immobilization onto TEAE-Cellulose. The original TEAE-Cellulose had a relative clear edge, and the TEAE-Cellulose with tyrosinase showed an unclear edge, rough surface and more porosity than the original one. Fig. 1C and D show the TEM image of original TEAE-Cellulose and TEAE-Cellulose with tyrosinase, respectively. The original TEAE-Cellulose surface was very smooth. After tyrosinase immobilization onto TEAE-Cellulose, its surface was rougher than the original one. Pure TEAE-Cellulose has a clear edge from the TEM and SEM images. After tyrosinase is immobilized onto TEAE-Cellulose, TEAE-Cellulose and tyrosinase can cross-link with each other through electrostatic attraction and hydrogen-bond interactions. In this way, the edge of TEAE-Cellulose becomes unclear and rough, and there is some porosity on its surface. Fig. S1A-S1E† display the SEM images of (A) Cellulose, (B) CM-Cellulose, (C) Pluronic F68, (D) poly(styrene-*co*-divinylbenzene) and (E)

glass beads. After immobilization of tyrosinase onto the surface of materials, cellulose, CM-Cellulose and Pluronic F68 morphologies totally changed, and their surfaces became rougher than before. These alterations showed that the tyrosinase had been immobilized onto these materials.

Immobilized tyrosinase biocatalytic ability

Six kinds of materials were used for the immobilization of tyrosinase: the triethylaminoethyl group of TEAE-Cellulose exhibits positive charge (pH 7.0, PBS solution); cellulose, Pluronic F68, poly(styrene-*co*-divinylbenzene) and glass beads have neutral charges; and the carboxymethyl group of CM-Cellulose has negative charge. As shown in Fig. 2, the TEAE-Cellulose could bind tyrosinase (negative charge at pH 7.0) due to charge attraction. TEAE-Cellulose is a positively charged resin used in protein purification or separation. It can lock negatively charged protein – tyrosinase onto the carrier by charge attraction and hydrogen-bond interactions. Cellulose can lock part of tyrosinase only through hydrogen-bond interactions. CM-Cellulose cannot lock tyrosinase like TEAE-Cellulose through charge attraction, but a few of tyrosinase can be locked by hydrogen-bond interactions. The tyrosinase molecules were physically adsorbed onto the surface of Pluronic F68, poly(styrene-*co*-divinylbenzene) and glass beads. The tyrosinase molecules immobilization efficiency of different materials could be arranged as: TEAE-Cellulose > cellulose = Pluronic F68 = poly(styrene-*co*-divinylbenzene) = glass beads > CM-Cellulose.

p-Cresol was chosen as the test substrate. The tyrosinase can function in chloroform whereby phenol derivatives can be converted into corresponding stable *o*-quinones. The enzymatic oxidation product of *p*-cresol is 4-methyl-1,2-benzoquinone, and its maximum absorbance peak is at 395 nm. Kinetic analysis of tyrosinase is shown in Fig. 3. The V_o values of immobilized tyrosinase onto TEAE-Cellulose, cellulose, poly(styrene-*co*-divinylbenzene), Pluronic F68, glass beads and CM-Cellulose were 6.7×10^{-2} , 5.5×10^{-2} , 4.4×10^{-2} , 3.6×10^{-2} , $2.8 \times$

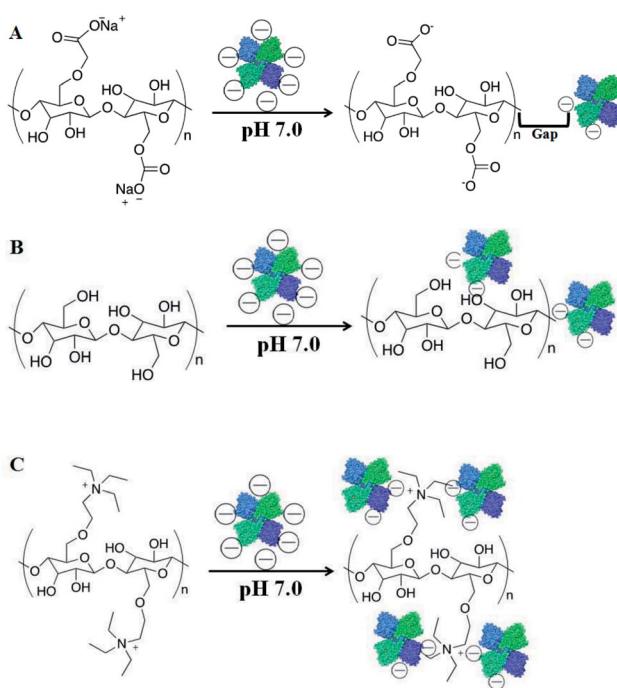


Fig. 2 Schematic diagram describing the mechanism of tyrosinase immobilization onto (A) CM-Cellulose, (B) cellulose and (C) TEAE-Cellulose.

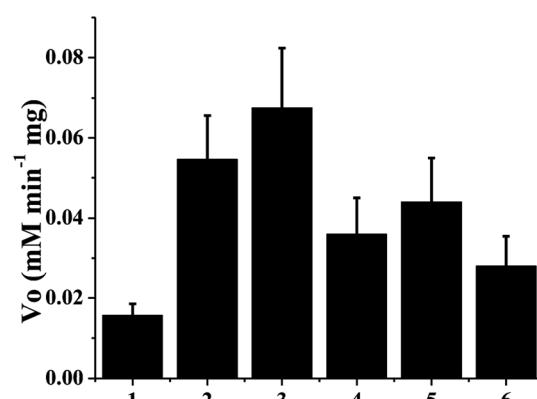


Fig. 3 V_o values of tyrosinase molecular immobilization onto CM-Cellulose, cellulose, TEAE-Cellulose, Pluronic F68, poly(styrene-*co*-divinylbenzene) and glass beads catalyzed oxidations of *p*-cresol as substrate in chloroform. *p*-Cresol concentration for these reactions is 50 mmol L⁻¹. Error bar means standard deviation.



Table 1 The activity of tyrosinase immobilization onto TEAE-Cellulose was detected in organic solvent (i.e. CH_2Cl_2 , CCl_4 and toluene)

<i>p</i> -Cresol	V_o (mmol L ⁻¹ min ⁻¹)	Std	log P	Viscosity 10 ⁻³ Pa s
Acetonitrile	—	—	-0.34	0.34
Tetrahydrofuran	—	—	0.46	0.46
1-Octanol	$<8.3 \times 10^{-4}$	—	3	7.4
<i>tert</i> -Butyl methyl ether	8.3×10^{-4}	2.6×10^{-4}	0.94	0.36
CH_2Cl_2	1.5×10^{-3}	2.5×10^{-4}	1.25	0.42
Chlorobenzene	5.7×10^{-2}	5.0×10^{-3}	2.84	0.75
CHCl_3	6.1×10^{-2}	4.6×10^{-3}	1.97	0.54
CCl_4	6.3×10^{-2}	3.0×10^{-3}	2.83	0.9
Toluene	6.7×10^{-2}	1.4×10^{-3}	2.73	0.55

10^{-2} and 1.5×10^{-2} mmol L⁻¹ min⁻¹, respectively. The V_o value of immobilized tyrosinase onto TEAE-Cellulose was the highest among those onto CM-Cellulose, cellulose, Pluronic F68, poly(styrene-*co*-divinylbenzene) and glass beads, and the V_o value of immobilized tyrosinase onto CM-Cellulose was the lowest among these materials. The results were consistent with the mechanism shown in Fig. 2. The TEAE-Cellulose was demonstrated to be the best supporting platform for tyrosinase and it was chosen as the immobilization material in the following research work.

Immobilization enzyme kinetics in organic solvent

After optimizing the tyrosinase immobilization platform, the stability of immobilized tyrosinase was systematically assessed in different organic solvents. Organic solvents can affect the structure of enzymes at the secondary, tertiary and quaternary levels.⁷ It is important to note how organic solvents affect the immobilized tyrosinase activity. From Table 1, the data demonstrates that the immobilized tyrosinase was completely inactive in hydrophilic solvents *viz.* acetonitrile ($\log P$, -0.34) and tetrahydrofuran ($\log P$, 0.46). This can be attributed to the fact that the hydrophilic solvents could strip the essential water from tyrosinase's hydration "shell" (hence deactivate the enzyme). Moreover, the immobilized tyrosinase had biocatalytic ability in hydrophobic solvent *i.e.* *tert*-butyl methyl ether ($\log P$, 0.94), methylene chloride ($\log P$, 1.25), chloroform ($\log P$, 1.97), carbon tetrachloride ($\log P$, 2.83), chlorobenzene ($\log P$, 2.84) and toluene ($\log P$, 2.73). Octanol–water partition coefficient ($\log P$) is used in QSAR studies and rational chemical design as a measurement of molecular hydrophobicity. Our results suggested that the hydrophobicity of organic solvent played an important role in tyrosinase efficiency. As the hydrophobicity of organic solvents increased, most of the biocatalytic efficiency of the immobilized tyrosinase increased. There is an exception to

this rule. 1-Octanol is categorized as a hydrophobic solvent and the value of $\log P$ is relatively high (achieved 3, similarly with toluene and carbon tetrachloride), but the V_o value of the immobilized tyrosinase in 1-octanol is relatively low (less than 8.3×10^{-4}). In this case, the viscosity of 1-octanol achieves as high as 7.4×10^{-3} Pa s, and the mass-transfer rate of substrate in 1-octanol is very slow. This might be one of the most important parameters in determining the biocatalytic efficiency of the immobilized tyrosinase. As a result, toluene was chosen as the solvent to test the immobilized tyrosinase activity.

The biocatalytic efficiency of the immobilized tyrosinase for different substituent groups of *para*-cresol isomers in hydrophobic organic solvent

After optimizing the immobilized tyrosinase solvent conditions, the relationship between the structure of the substrate and the activity of the immobilized tyrosinase was systematically studied. How the substituent group position (three isomers of *p*-cresol) influenced the selectivity of the immobilized tyrosinase in toluene was examined. Table 2 shows that the V_o value for tyrosinase in the toluene medium, using *p*-cresol as substrate, was 8.7-fold higher than those obtained with *m*-cresol as substrate and more than 30-fold higher than those with *o*-cresol as substrate (the V_o number was too small to check). The *ortho*-substituent group was not significantly catalyzed by the immobilized tyrosinase because such substrates sterically hinder the accessibility of the phenolic hydroxyl moiety to the active-site of tyrosinase. The steric influence of the *para* substituent group of phenol for the biocatalysis of tyrosinase was less than that of *meta* or *ortho* substituents.

It was essential to know how the different *para* substituent groups influence the catalytic efficiencies of the immobilized tyrosinase. Different *para*-substituent groups of phenol were tested in toluene. Table 3 shows that the V_o value of tyrosinase

Table 2 The bioactivity of tyrosinase was tested for different substituent sites in different solvent (i.e. CH_2Cl_2 , CCl_4 and toluene)

Substrate	<i>p</i> -Cresol (V_o)	<i>p</i> -Cresol (Std)	<i>m</i> -Cresol (V_o)	<i>m</i> -Cresol (Std)	<i>o</i> -Cresol (V_o)	<i>o</i> -Cresol (Std)
Toluene	6.7×10^{-2}	1.4×10^{-3}	7.7×10^{-3}	2.2×10^{-4}	$<8.3 \times 10^{-4}$	—
Chlorobenzene	5.7×10^{-2}	5.0×10^{-3}	6.5×10^{-3}	1.9×10^{-5}	$<8.3 \times 10^{-4}$	—
CHCl_3	6.1×10^{-2}	4.6×10^{-3}	7.0×10^{-3}	1.2×10^{-4}	$<8.3 \times 10^{-4}$	—
CCl_4	6.3×10^{-2}	3.0×10^{-3}	7.2×10^{-3}	1.2×10^{-4}	$<8.3 \times 10^{-4}$	—



Table 3 The bioactivity of tyrosinase was detected for six kinds of *para*-substituents of phenol in toluene

Substrate	V_o (mmol L ⁻¹ min ⁻¹)	Std	$\log K_{ow}$	Solubility in water (mol m ⁻³)
<i>p</i> -Methyl	6.7×10^{-2}	1.4×10^{-3}	1.96	184.9
<i>p</i> -Ethyl	4.1×10^{-2}	3.2×10^{-3}	2.50	65.3
<i>p</i> -Propyl	2.8×10^{-2}	2.1×10^{-3}	3.20	12.7
<i>p</i> -Chloro	2.1×10^{-2}	1.8×10^{-3}	2.40	210.0
<i>p</i> -Bromo	9.2×10^{-3}	8.5×10^{-4}	2.59	85.54
<i>p</i> - <i>t</i> -Butyl	2.0×10^{-3}	2.0×10^{-5}	3.65	3.861

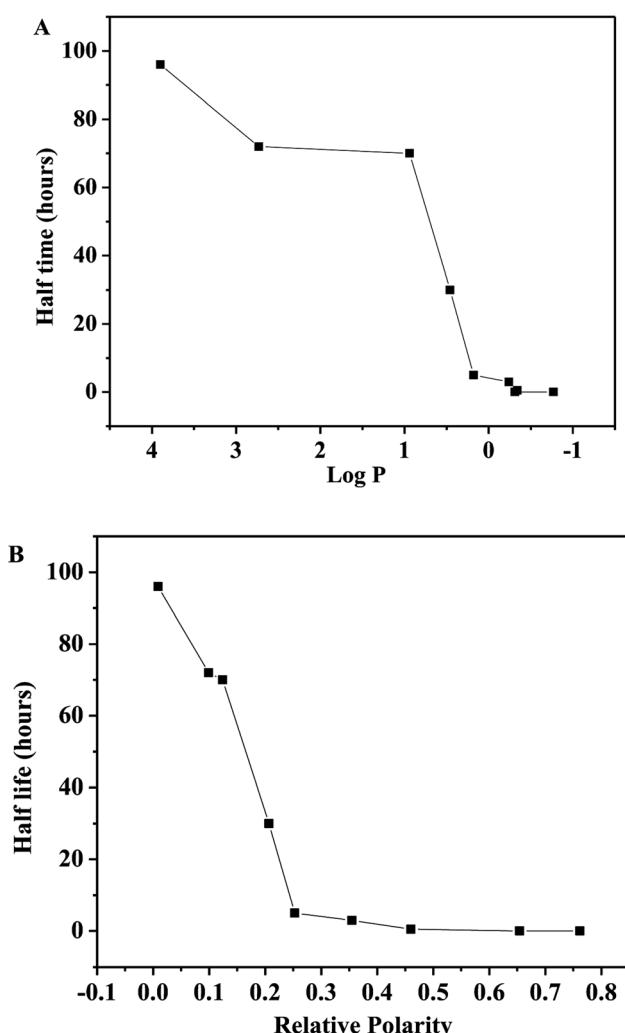


Fig. 4 (A) The half-life of the immobilized tyrosinase vs. the $\log P$ values of the organic solvent, (B) the half-life of the immobilized tyrosinase vs. the relative polarity values of the organic solvent. The organic solvent from right to left is methanol, ethanol, acetonitrile, acetone, methyl acetate, tetrahydrofuran, methyl *t*-butyl ether, toluene and hexane.

biocatalysis for *para* substituent were: *p*-methyl > *p*-ethyl > *p*-propyl > *p*-chloro > *p*-bromo > *p*-*t*-butyl. The methyl, ethyl, propyl and *t*-butyl groups are electron-donating substituent groups. By increasing the electron-donating ability and the steric size of *para*-substituents, the corresponding V_o value of

the immobilized tyrosinase decreased. It indicated that the tyrosinase activity was related to the electron-donating ability and steric size of the substituent. In contrast, *p*-chloro and *p*-bromo are electron-withdrawing substituent groups; their corresponding V_o value of the immobilized tyrosinase decreased with their deactivating ability. The conclusion was consistent with the quantitative assessment of enzymatic catalysis by Dordick *et al.*⁴⁹

After optimizing the immobilized platform, the substrate and the solvent, the behavior of tyrosinase in anhydrous organic solvent at high temperature was studied. In this way, the immobilized tyrosinase was treated as a synthetic catalyst. Nine kinds of anhydrous organic solvents were used for storing the immobilized tyrosinase at 100 °C. The nine anhydrous organic solvents are methanol ($\log P$, -0.77; relative polarity, 0.762), ethanol ($\log P$, -0.31; relative polarity, 0.654), acetonitrile ($\log P$, -0.34; relative polarity, 0.46), acetone ($\log P$, -0.24; relative polarity, 0.355), methyl acetate ($\log P$, 0.18; relative polarity, 0.762), tetrahydrofuran ($\log P$, 0.46; relative polarity, 0.207), methyl *t*-butyl ether ($\log P$, 0.94; relative polarity, 0.124), toluene ($\log P$, 2.73; relative polarity, 0.099) and hexane ($\log P$, 3.9; relative polarity, 0.009). As shown in Fig. 4, the half-life of the immobilized tyrosinase is strongly related to the relative polarity value of the storage solvent than their $\log P$ values. After the immobilized tyrosinase is immersed in anhydrous hexane at 100 °C for 96 hours, the immobilized tyrosinase still retains over 60% activity. Conversely, the immobilized tyrosinase loses the activity immediately in water at 100 °C. This indicated that the immobilized tyrosinase could keep its catalytic structure and be used as a synthetic catalyst in anhydrous organic solvent at high temperature. This research provides a lot of information for mimicking enzyme reactions.

Conclusion

Triethylaminoethyl cellulose (TEAE-Cellulose) was selected as the best immobilization platform from six kinds of materials. Compared with traditional immobilization materials (glass beads), the activity of tyrosinase immobilization onto TEAE-Cellulose was two times higher than that of immobilization onto glass beads. The immobilized tyrosinase showed potential to be a catalyst in organic solvent without water. The half-life of the dry immobilized tyrosinase in organic solvent was strongly related to the polarity of the solvent rather than the $\log P$ value of the organic solvent. This research could be valuable for the design of new biocatalysts.



Conflicts of interest

There are no conflicts to declare.

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