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# Chemoenzymatic epoxidation of alkenes with *Candida antarctica* lipase B and hydrogen peroxide in deep eutectic solvents†

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Epoxides are important synthetic intermediates for the synthesis of a broad range of industrial products. This study presents a promising solution to the current limitation of enzyme instability. By using simple deep eutectic solvents such as choline chloride/sorbitol, significant stabilization of the biocatalyst has been achieved leading to more robust reactions while using environmentally more acceptable solvents as compared to ionic liquids.

## Introduction

Epoxides are important synthetic intermediates in the production of many industrial products.<sup>1</sup> Next to established chemical syntheses such as the Prileshajev epoxidation using stoichiometric peracids<sup>2</sup> or metal-catalyzed epoxidations,<sup>3,4</sup> enzymatic<sup>5-9</sup> and chemoenzymatic methods<sup>10-12</sup> have also gained relevance. Particularly, the latter methods are attractive due to the broad availability of hydrolases. In essence, chemoenzymatic methods rely on the *in situ* formation of a reactive peracid making use of the 'perhydrolyase' promiscuous activity of many lipases. The peracids formed react with the alkene forming the product of interest and re-forming the carboxylic acid, which enters a new catalytic cycle (Scheme 1).<sup>13</sup>

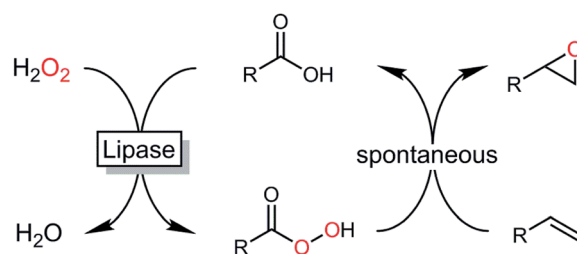
A broad range of hydrolases has been investigated for the perhydrolysis of carboxylic acids or their esters with hydrogen peroxide to produce epoxides.<sup>14-18</sup> Among them the lipase B from *Candida antarctica* (CaLB), immobilized onto an acrylic resin (tradename Novozym 435), is probably the most-widely used biocatalyst.<sup>19</sup> Several studies dealing with lipase-catalyzed *in situ* peracid formation to promote the Prileshajev epoxidation underline the preparative potential of this approach.<sup>20-23</sup>

In recent years, it has become clear that CaLB suffers from a pronounced instability against the harsh conditions of the chemoenzymatic epoxidation reaction as compared to the 'natural' esterification reactions. Arends and coworkers

reported that ionic liquids (ILs) could substantially improve the performance of chemoenzymatic Baeyer-Villiger and epoxidation reactions.<sup>24</sup> The authors suggested a two-fold effect of hydrogen-bond-donor-ILs in the reaction: first by stabilizing the biocatalyst and second, by accelerating the reaction through solvent-activation of the peracid.

Though frequently termed 'green solvents' (mostly due to their non-volatility), ILs are actually questionable from an environmental point-of-view.<sup>25</sup> For example, toxicity issues can impair the environmental friendliness as well as the sometimes rather complex and resource-consuming synthesis (*vide infra*). Therefore, it is not astonishing that current research efforts focus on environmentally less demanding substitutes such as (natural) deep eutectic solvents (DESs).

The number of studies evaluating DESs as more environmentally acceptable alternatives to ILs is steadily increasing.<sup>26</sup> For example, beneficial effects of DESs on lipase-catalysed esterification reactions have been reported.<sup>27-29</sup> Inspired by these very promising reports, we set out to investigate the possible beneficial effects of DESs on the chemoenzymatic epoxidation reaction.



**Scheme 1** Chemoenzymatic epoxidation reaction using *in situ* generated peracids. In the first step, lipase-catalyzed perhydrolysis of a carboxylic acid with H<sub>2</sub>O<sub>2</sub> forms the corresponding peracid, which then, in a non-enzymatic Prileshajev reaction forms the epoxide and the (catalytic) carboxylic acid.

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## Results and discussion

In a first set of experiments, we evaluated the influence of various DESs of the efficiency of the chemoenzymatic epoxidation of 1-octadecene as a model reaction (Table 1).

From the nine DESs chosen for this experiment, the amine-based DES gave the poorest results. Essentially product formation ceased already after approx. 6 h. Possibly, inactivation of the enzyme (maybe due to deprotonation of structurally essential amino acid residues) accounted for this behavior. Polyol-based DESs gave significantly higher robustness and consequently also higher product titers. Hence, ChCl/sorbitol as solvent resulted in more than 70% conversion of the starting material. This effect has also been observed by Arends and co-workers who have pointed out the importance on hydrogen-bond-donating cosolvents (ILs) on the stabilization of the protein structure.<sup>24</sup> Similarly, Diego *et al.* have emphasized the importance of polyols such as sorbitol on protein structure.<sup>30</sup>

Encouraged by these results, we became interested in shedding some more light on the beneficial and detrimental effect of amine-based or polyol-based DESs on the stability of the biocatalyst. Therefore, the apparent half-life times of CalB in the presence of different solvents were determined (Fig. 1). In accordance to our previous observation,<sup>31</sup> the amine-based DESs significantly reduced (as compared to buffer as incubation solvent) the stability of CalB over the entire temperature range investigated whereas the polyol-based DESs significantly increased the stability of the biocatalyst.

The effect of the single solvents on the structural integrity of CalB was investigated in some more detail using circular dichroism spectroscopy (CD). As shown in Table 2, the relative amount of secondary structure elements within the enzyme decreased to some extent after incubation in phosphate buffer for 24 h, while in ChCl/urea almost all secondary structural

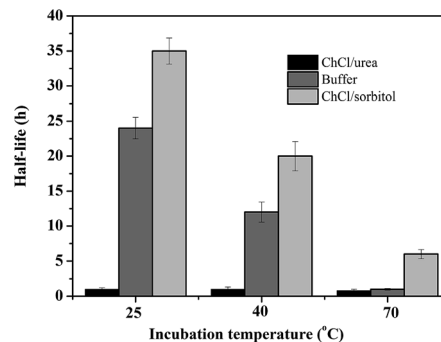


Fig. 1 CalB stability (displayed as half life time) at different temperatures in the presence of buffer, ChCl/urea and ChCl/sorbitol.

elements disappeared at the benefit of random coil. On the contrary, secondary structure elements were almost entirely preserved (as compared to the freshly dissolved enzyme) upon incubation of the enzyme in ChCl/sorbitol. This result is also in agreement with our previous study.<sup>31</sup> Hence, we conclude that the increased stability of the enzyme was due to the preservation of its structural integrity in ChCl/sorbitol.

Next, the key parameters effecting the chemoenzymatic epoxidation reaction; particularly the influence of the reaction temperature, the octanoic acid concentration and the hydrogen peroxide concentration were investigated (Fig. 2).

Quite expectedly, initial this reaction rate increased linearly with temperature (Fig. 2A). Hence, the rate-limiting step may be diffusion of either the enzyme substrates (octanoic acid and/or H<sub>2</sub>O<sub>2</sub>) or products (peroctanoic acid). It is worth pointing out here that in similar studies Törnvall *et al.*<sup>32</sup> reported a sharp decrease of the enzyme activity at 60 °C. This underlines the stabilizing effect of ChCl/sorbitol on the enzyme.

The initial rate of the overall reaction showed a saturation-type behavior with respect to the concentration of octanoic acid used (Fig. 2B). Above 1.6 mmol of octanoic acid, no significant increase of the initial rate was observed. This may be due to the Michaelis–Menten-type kinetics of the lipase-catalyzed perhydrolysis reaction with octanoic acid. However, it should be pointed out that in these experiments we could

Table 1 Epoxidation of 1-octadecene in different DESs<sup>a</sup>

Entry	Solvents (molar ratio)	Conversion (%)
1	ChCl/urea (1 : 2)	30.2
2	ChCl/acetamide (1 : 2)	13.9
3	ChCl/ethylene glycol (1 : 2)	37.7
4	ChCl/glycerol (1 : 1)	56.7
5	ChCl/xylitol (1 : 1)	65.1
6	ChCl/sorbitol (1 : 1)	72.4
7	ChCl/xylose/water (5 : 2:5)	58.4
8	ChCl/glucose/water (5 : 2:5)	65.3
9	ChCl/sucrose/water (5 : 2:5)	56.2
10	Phosphate buffer (pH6.0, 50 mM)	62.5

<sup>a</sup> Conditions: octadecene (1.6 mmol), octanoic acid (1.6 mmol) and H<sub>2</sub>O<sub>2</sub> (1.6 mmol added as 30% w/w solution) were reacted in different DESs at 40 °C using Novozym 435 (100 mg) as catalyst for 24 h with magnetic stirring (300 rpm).

Table 2 Secondary structure percentages of CalB after incubation in different media

Medium	$\alpha$ -helix [%]	$\beta$ -strand [%]	Turn [%]	Random coil [%]
Initial CalB <sup>a</sup>	19.8	13.4	31	35.8
KP <sub>i</sub> buffer <sup>b</sup>	12.3	17.5	33.5	36.7
ChCl/urea <sup>c</sup>	2.2	1.3	34.3	62.2
ChCl/sorbitol <sup>d</sup>	16.6	19.1	32.5	32

<sup>a</sup> Determined directly after complete dissolution of the enzyme.

<sup>b</sup> Conditions: CalB incubated in 50 mM pH 6.0 phosphate buffer.

<sup>c</sup> Conditions: CalB incubated in ChCl/urea and. <sup>d</sup> Conditions: CalB incubated in ChCl/sorbitol at 30 °C for 24 h.



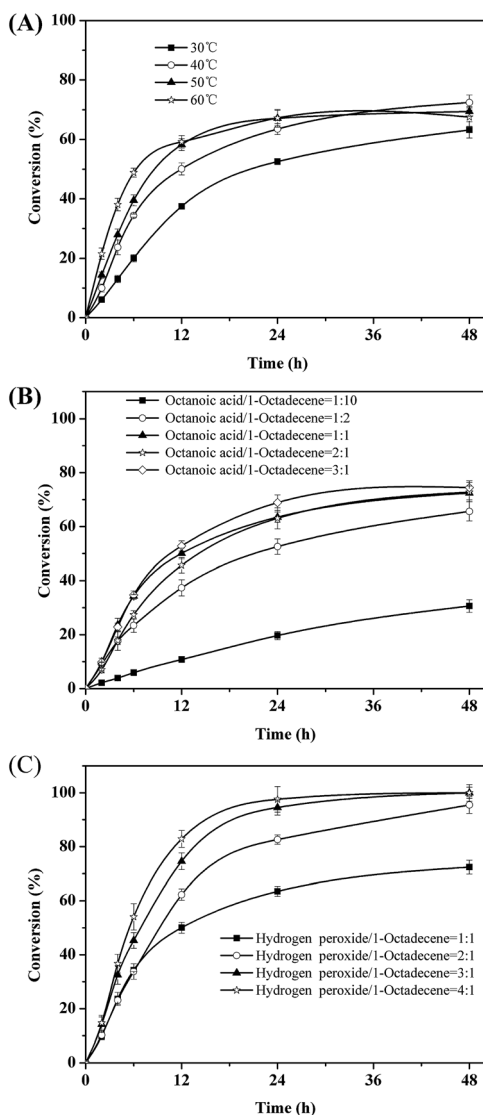


Fig. 2 Effects of temperature (A), molar ratio of octanoic acid to 1-octadecene (B) and molar ratio of hydrogen peroxide to 1-octadecene molar ratio (C) on the epoxidation of 1-octadecene.

demonstrate that octanoic acid performs as catalyst (TN = 3), pointing towards a truly catalytic system. Nevertheless, for all further experiments we used a molar ratio of 1 : 1 (acid : alkene) in order not to impair the overall reaction rate. Again, the maximal conversion reached in these experiments was approx. 70%.

Interestingly, the amount of hydrogen peroxide applied to the reaction system had almost no influence on the initial rate of the reaction (which may be explained by a comparably low  $K_M$  value for  $H_2O_2$ ) (Fig. 2C). But it had a very significant influence on the overall conversion reached. At a nominal molar ratio of 1 : 1 ( $H_2O_2$  to alkene) the maximal conversion observed in all experiments so far was approx. 70%. However, already at a molar ratio of 2 : 1 ( $H_2O_2$  to alkene) full conversion was achieved. We believe that the most likely explanation for this observation may be that the active  $H_2O_2$  content of the stock solution may have been significantly lower (approx. 70%) than

the nominal concentration. Further experiments clarifying this issue are currently underway.

We also evaluated the substrate scope of this partially optimized reaction system. As shown in Table 3, a range of (ali) cyclic alkenes as well as styrene derivatives were converted in good to excellent conversions in high selectivity as confirmed by NMR analysis of the isolated products (see ESI† for details). Only trace amounts (if any) of possible degradation (isomerization or hydrolysis) products were detectable.

The performance of the proposed chemoenzymatic reaction system using ChCl/sorbitol as ‘performance additive’ compares well with similar reactions reported in the literature.<sup>31</sup> For example, the conversion of 1-octadecene in aqueous reaction media was 68% or 30% using the lipase from *Penicillium camemberti*<sup>33</sup> or the lipase from *Malassezia globosa*,<sup>34</sup> respectively. In a three-phase system of water/methyl dichloride/ionic liquid, the conversion of cyclohexene was 41%.<sup>18</sup> Therefore, the new oxidation system proposed here represents a promising approach for preparative epoxidation reactions.

To further understand the molecular basis of the perhydrolysis reaction we modelled octanoic acid and  $H_2O_2$  into the active site of CalB (Fig. 3). Carbonyl atom of the octanoic acid is bound in the oxyanion hole of the enzyme by two hydrogen bonds to the amide nitrogens of Gln106 and Thr40. The distance between carbonyl atom of octanoic acid and  $O_\gamma$  of Ser105 is 2.9 Å, which is facilitate to nucleophilic attack carbonyl group by Ser105 in active site to form an acetyl-enzyme intermediate. Simultaneously,  $H_2O_2$  is stabilized in the active site by Gln157, Asp134 and His224. Then, hydrogen peroxide attacked acetyl-enzyme intermediate to achieve perhydrolysis reaction. This mechanism is similar to previous reports.<sup>33–35</sup>

Table 3 Substrate scope of the optimized chemoenzymatic epoxidation system<sup>a</sup>

Product	Reaction time [h]	Conversion [%]
	24	82 <sup>b</sup>
	11	85
	24	97
	11	72
	24	90
	11	90

<sup>a</sup> General conditions: 0.4 g ChCl/sorbitol, 100 mg Novozym 435, 1.6 mmol octanoic acid, 1.6 mmol alkene and 3.2 mmol 30 wt%  $H_2O_2$  at 40 °C with magnetic stirring 300 rpm. <sup>b</sup>  $T = 30$  °C.



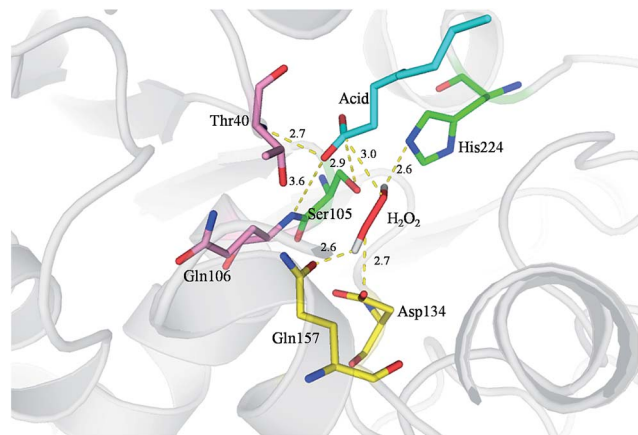


Fig. 3 The mode of CalB (PDB ID: ILBT) catalyzed epoxidation mechanism. The atom of hydrogen, oxygen and nitrogen are shown in white, red and blue, respectively. Hydrogen bonds are shown in yellow dash with distances in angstroms.

Finally, we performed a preliminary evaluation of the environmental impact of the chemoenzymatic epoxidation reaction. For this, the Sheldon's E-factor (environmental factor) analysis was used (eqn (1))<sup>36,37</sup> and compared our chemoenzymatic epoxidation of styrene with a similar reaction reported by Arends and coworkers.<sup>24</sup>

$$\text{E-factor} = \frac{\sum m(\text{waste products}) [\text{kg}]}{m(\text{product}) [\text{kg}]} \quad (1)$$

Eqn (1) calculation of Sheldon's E-factor.

It becomes clear from Table 4, that the solvent (either ChCl/sorbitol or [BMIM][BF<sub>4</sub>]) has a very significant impact on the E-factor of the reaction contributing 31% or even 75% to the total value, respectively.

Given this importance of the solvents, also their upstream processing should be considered. Following Jessop's approach of evaluating the environmental impact of solvent synthesis based on the synthesis trees<sup>38</sup> we compared those for [BMIM][BF<sub>4</sub>] and ChCl/sorbitol (Fig. S1†). This comparison reveals that

compared to [BMIM][BF<sub>4</sub>] (30 steps) the synthesis of ChCl/sorbitol with only 10 steps is significantly less complex. Furthermore, most of the solvent (55% w/w) of ChCl/sorbitol originates from renewable resources. Of course, only a full life cycle assessment (LCA) will be suitable to holistically compare the environmental impact of the solvent syntheses and the chemoenzymatic reactions, but we hold the results from this preliminary analysis to be very encouraging.

## Conclusions

In the present study, it has been demonstrated that DESs represent a true alternative to the commonly used ILs to facilitate the chemoenzymatic epoxidation of alkenes. The system excels particularly by the lower environmental impact of the cosolvents (DESs) used as compared to ILs. This advantage, however, does not come with an impaired performance compared to classical ILs. The practical applicability and efficiency of the current system is at least as good as the one of the systems of the state-of-the-art. Therefore, we are convinced that exciting future developments will arise from the application of DESs to chemoenzymatic epoxidation reactions.

## Experimental sections

### Materials

*Candida antarctica* lipase B (CalB, Novozym 435-immobilized on a polyacrylate resin) was obtained from Novo Nordisk (Denmark). The 30% (w/w) hydrogen peroxide, choline chloride (98%), urea (99%), acetamide (99%), ethylene glycol (98%), glycerol (99%), xylitol (98%), D-sorbitol (98%), D-xylose (98%), D-(+)-glucose (99.5%), sucrose (ACS grade), 1-hexene (99%), cyclohexene (99%), 1-decene (99.5%), 1-octadecene (95%), styrene (99%),  $\alpha$ -methylstyrene (99%), cyclohexene oxide (98%) and styrene oxide (98%) were bought from Aladdin Chemistry Co., Ltd (Shanghai, China). 1,2-Hexylene oxide (98%), decene oxide (98%), octadecene oxide (99%) and  $\alpha$ -methylstyrene oxide (97%) were obtained from TCI (Japan). All other reagents were of analytical grade.

### Preparation DESs

Deep eutectic solvents were formulated in different molar ratios of quaternary ammonium salt (choline chloride) to hydrogen bond donors. The eutectic mixtures were prepared by rotary evaporation of the two components at 80 °C in water bath until a homogeneous transparent liquid was formed.

### Stability of enzyme

Purified CalB enzyme was added into 50 mM phosphate buffer pH 6.0, DESs ChCl/urea or ChCl/sorbitol incubated at temperature from 25 °C to 70 °C for 12 h. The samples were extracted from reaction mixture at periodically, and then the perhydrolysis activity was followed spectrophotometrically at 290 nm as described by Peter Bernhardt.<sup>39</sup> The tests were carried out in 96-well microtiter plates with a reaction of volume 100  $\mu$ L (80  $\mu$ L assays solution to 10  $\mu$ L samples and 10  $\mu$ L hydrogen peroxide

Table 4 E-factor analysis

Entry	DESs system <sup>a,b</sup>	ILs system <sup>a24</sup>
Non-converted (styrene) conversion (%)	0.016 g (0.09) 90%	0.08 g (1) 40%
CalB (biocatalyst)	0.1 g (0.58)	0.01 g (0.13)
Octanoic acid	0.23 g (1.35)	0.029 g (0.2)
Solvent	0.4 g (ChCl/sorbitol) (2.35)	1 g ([BMIM][BF <sub>4</sub> ]) (12.5)
H <sub>2</sub> O/H <sub>2</sub> O <sub>2</sub>	0.56 g (3.29)	0.14 g (1.75)
Product	0.17 g	0.08 g
E-factor	7.7	16.7

<sup>a</sup> Numbers in parentheses represent the individual E-factor contributions (i.e. g component  $\times$  g product<sup>-1</sup>). <sup>b</sup> Results taken from Table 3.



in 0.1 M pentanoic acid buffer pH 5.0) at 40 °C for 5 minutes using microplate reader Cytation™ 5 (BioTek, USA). The blank experiments were carried out under the identical conditions adding inactivated enzymes. All tests were performed in triplicate.

### Epoxidation reaction

In a 10 mL Erlenmeyer flask were added 1.6 mmol of 1-octadecene, 1.6 mmol of octanoic acid, 1.6 mmol and 1.6 mmol of 30% hydrogen peroxide, 0.4 g of DESs and 100 mg Novozym 435 at 40 °C stirring with 300 rpm. The control without DESs was performed in the phosphate buffer. The 30% hydrogen peroxide was added in 3 portions in 1 h intervals. Samples were withdrawn in different time-points. 20 µL of solution was added into 980 µL methanol and then for gas chromatograph analysis.

### Analytical methods

Analysis of olefins and corresponding epoxides were determined using an Agilent Technology model 7890 GC (Agilent, USA), equipped with an HP-5 column (30.0 m × 0.25 mm, 0.25 µm, Macherey Nagel, Germany). Peak of olefins and corresponding epoxides in GC were identified and calculated by the calibration curves which were made with the reference standard compounds.

### Circular dichroism spectroscopy

Circular dichroism (CD) spectroscopy measurements of protein were carried out with a Chirascan spectropolarimeter (Applied Photophysics, Surrey, UK), using 0.2 mg mL<sup>-1</sup> of purified CalB enzyme in different medium. The path length was 0.5 mm. Spectra covering the wavelength range of 190–260 nm were scanned at 25 °C. The scanning rate is 1 nm per 0.5 s. Each spectrum was the average of three successive scans. Under the same conditions, the corresponding medium solution was recorded as control and subtracted from the sample spectra. The secondary structure percentages were analyzed by using the CDNN tool.

### Identification of epoxide

Epoxidized alkenes was identified by nuclear magnetic resonance (NMR) spectra (Bruker Avance 600 MHz). Samples were dissolved in 600 µL deuterated chloroform in a NMR tube and subjected to <sup>1</sup>H and <sup>13</sup>C NMR analysis.

### Structure analysis

The modeling of the ligand carboxylic acid in the binding site and complex energy minimization has been performed with MODELLER package (Discovery Studio 3.5). The structure 1LBT and the model of its open conformation<sup>40</sup> has been used for analysis.

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