

## Epoxidation and oxidation reactions using 1,4-butanediol dimethacrylate crosslinked polystyrene-supported tertiary butyl hydroperoxide

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**Abstract.** 1,4-Butanediol dimethacrylate (1,4-BDDMA) crosslinked polystyrene-supported *t*-butyl hydroperoxide was employed in the epoxidation of olefins and oxidation of alcohols to carbonyl compounds. The reagent proved to be successful as a recyclable solid phase organic reagent with as much or more efficiency when compared to its monomeric counterpart. The extent of reaction was found to be dependent on various reaction parameters like solvent, temperature, molar concentration and presence of catalyst.

**Keywords.** Polystyrene-supported reagent; *t*-butyl hydroperoxide; solid-phase organic reagent; recyclability; epoxidation; oxidation.

### 1. Introduction

Solid-phase organic synthesis has been used as an efficient technique for the synthesis and screening of a large number of organic compounds.<sup>1</sup> Over the past years, numerous techniques have been used to study polymeric resins commonly utilized in solid phase synthesis to allow greater understanding of the chemical nature and physical properties of the supports. In addition, to overcome some of the drawbacks of existing polymer supports, several new resins and new methods of handling solid supports have been developed.<sup>2–4</sup> New methodologies have also been introduced to simplify the preparation of solid supports. A number of reactive species have been attached to polymeric supports and have been successfully used in organic synthesis.<sup>5–9</sup> Polymer-bound reagents have been reported for the oxidation of various alcohols.<sup>10</sup> Peroxy compounds are widely used as initiators for polymerisation of monomers. However, they are not popular as solid-phase organic reagents. Divinyl benzene cross-linked polystyrene-supported *t*-butyl hydroperoxide has already been developed in our laboratory, as a recyclable solid-phase organic reagent for performing various synthetic transformations.<sup>11</sup> Reactivity of the attached functional group is strongly influenced by the nature

of the polymer backbone.<sup>12–15</sup> 1,4-BDDMA cross-linked polystyrene-supported *t*-butyl hydroperoxide has also been developed for the conversion of alkyl and aryl halides to hydroxy compounds through a Grignard type reaction.<sup>16</sup> In this article, we present the epoxidation of olefins and oxidation of alcohols using 1,4-BDDMA crosslinked polystyrene-supported *t*-butyl hydroperoxide.

### 2. Experimental

#### 2.1 General

Solvents used were of reagent grade and were purified according to literature procedures. Melting points were determined on a hot stage melting point apparatus and are uncorrected. IR spectra were recorded on a Perkin–Elmer 887 IR spectrometer using KBr pellets. Thin layer chromatography (tlc) was performed on pre-coated silica gel plates.

#### 2.2 Epoxidation of olefins – General procedure

The olefin (1 mmol) in dioxan (30 ml) was stirred with a two-fold molar excess of the polymer-supported *t*-butyl hydroperoxide reagent.<sup>16</sup> The reaction was monitored by tlc as well as by estimating the residual hydroperoxide capacity. The epoxide content was determined by estimating the amount of oxirane

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oxygen by back titration with 1N HCl in the presence of dioxan. After completion of reactions, resin was filtered and washed with dioxan (10 ml  $\times$  3 times). The combined filtrate and washings, on evaporation of the solvent afforded the corresponding epoxides.

### 2.3 Oxidation of alcohols – General procedure

The alcohol (1 mmol), dissolved in chloroform (20 ml), was stirred with a five-fold molar excess of the polymer-supported *t*-butyl hydroperoxide resin. The reaction was followed by tlc. After the completion of the reaction the resin was filtered and washed with chloroform. The combined filtrate and washings, on evaporation of the solvent afforded the corresponding carbonyl compounds.

## 3. Results and discussion

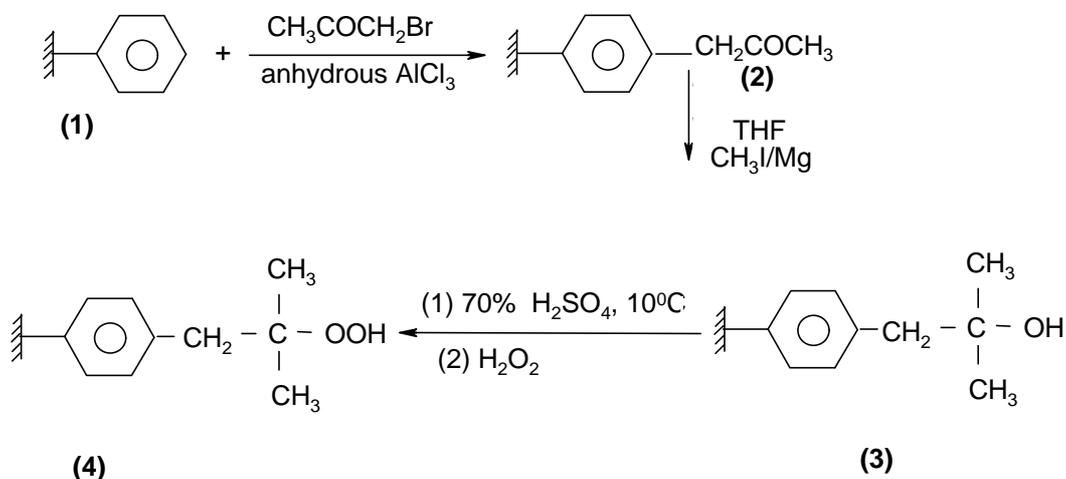
### 3.1 Preparation and characterization of the reagent

Two percent 1,4-BDDMA crosslinked polystyrene (**1**) was prepared by suspension polymerization. The white bead-shaped polymer obtained was purified by soxhlet extraction using tetrahydrofuran. It was identified by the characteristic IR spectral peaks at 1520  $\text{cm}^{-1}$  (C=C str), 906  $\text{cm}^{-1}$  (vinyl group, C–H bending), 3030  $\text{cm}^{-1}$  (C–H str) and 1740  $\text{cm}^{-1}$  (C=O ester). The polystyrene resin was functionalized with *t*-butyl hydroperoxide group by a three-step polymer analogous reaction (scheme 1). The 2-oxopropyl polystyrene (**2**) was estimated by iodoform

reaction to possess a  $\text{CH}_3\text{COCH}_2$ - capacity of 4.2 meq/g of the polymer. Spectral characterization showed intense absorption band at 1720  $\text{cm}^{-1}$  in the infrared region. When the 2-oxopropyl polystyrene resin was converted to polymeric *t*-butyl alcohol resin (**3**) the absorption observed for the keto resin at 1720  $\text{cm}^{-1}$  disappeared indicating the conversion of the keto group. The polymeric *t*-alcohol was estimated to possess a capacity of 3.2 meq/g by the gravimetric method. Polymeric *t*-butyl alcohol was converted to *t*-butyl hydroperoxide reagent by the action of  $\text{H}_2\text{O}_2$  in the cold, in the presence of 70%  $\text{H}_2\text{SO}_4$ . The resin exhibited strong absorption bands at 876  $\text{cm}^{-1}$ , 1580  $\text{cm}^{-1}$  (C=C arom str) and 3310  $\text{cm}^{-1}$  (O–H str) in the infrared region. The resin was estimated iodimetrically to possess a hydroperoxide capacity of 3.31 meq/g. The 1,4-BDDMA crosslinked polystyrene backbone was found to be more reactive towards functionalization. Utilization of the crosslinker, 1,4-BDDMA, instead of DVB, raised the hydroperoxide capacity of the polymeric reagent from 2.86 meq/g to 3.31 meq/g when the reaction was performed under identical conditions. This may be attributed to the hydrophilic and flexible nature of the 1,4-BDDMA crosslinks imparting better hydrophobic–hydrophilic balance for the macromolecular environment.

### 3.2 Epoxidation of olefins

The 1,4-BDDMA crosslinked polystyrene bound reagent was found to epoxidise various olefins as shown in table 1. Epoxidation of olefins with the *t*-butyl hydroperoxide reagent is as shown in scheme 2.

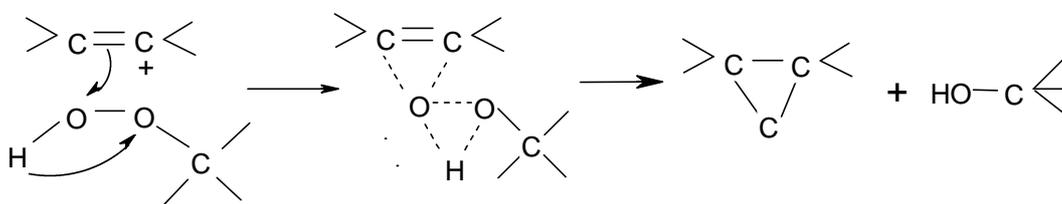


Scheme 1. Functionalisation of polystyrene resin.

**Table 1.** Epoxidation of olefins using 1,4-BDDMA-crosslinked polystyrene supported *t*-butyl hydroperoxide.

Olefin <sup>a</sup>	Reaction time <sup>b</sup> (h)	Product <sup>c</sup>	Isolated yield (%)
Cinnamic acid	39	Phenyl glycidate	40
Ethyl cinnamate	23	Ethyl phenyl glycidate	58
Allyl bromide	28	Epibromohydrin	52
Styrene <sup>d</sup>	48	Styrene oxide	54
Cyclohexene	16	Cyclohexene oxide	54
Methyl methacrylate <sup>d</sup>	21	Epoxyethyl propionate	50
Acenaphthylene <sup>d</sup>	36	Acenaphthylene oxide	44
Diphenyl acetylene	36	1,2-Epoxydiphenyl ethane	52
2-Chloroacrylonitrile	32	1-Chloro-1,2-epoxycyanoethane	48

<sup>a</sup>Olefin to resin ratio, 1 : 2; solvent, dioxan; temperature, 70°C; <sup>b</sup>includes preswelling time; <sup>c</sup>characterized by comparison with authentic samples (m.p./b.p. and IR); <sup>d</sup>temperature, 30°C

**Scheme 2.** Epoxidation of olefins using polymeric *t*-butyl hydroperoxide reagent.

The uncatalysed epoxidation yielded epoxides ranging from 40 to 58%. When compared to the reaction with the DVB crosslinked system, this reagent showed more efficiency by reducing the reaction time considerably, even though an appreciable change was not observed in the yield of epoxide isolated. This was evidenced by the epoxidation of cyclohexene which required only 16 h instead of 24 h with the DVB crosslinked reagent.<sup>17</sup> The reaction time for styrene was reduced from 56 to 48 h, and that of ethyl cinnamate from 36 to 23 h. The reaction temperature of the reaction varied with the nature of the olefins. In the case of vinyl monomers, epoxidation at higher temperature yielded polymerised products instead of the epoxides. Hence, the epoxidation of substrates like methyl methacrylate was conducted at room temperature. Substrates like cinnamic acid and ethyl cinnamate were epoxidised when the temperature was raised to 70°C. The successful epoxidation of diphenyl acetylene in 52% yield after 36 h reveals the capability of the reagent to epoxidise acetylenic compounds. While cyclohexene epoxidised in 16 h, acenaphthylene required 36 h for the reaction. Hence, the reaction seems to be diffusion-controlled and depended on the molecular size of the

substrate. The products were identified in comparison with the physical constants of authentic samples and from IR spectral values.

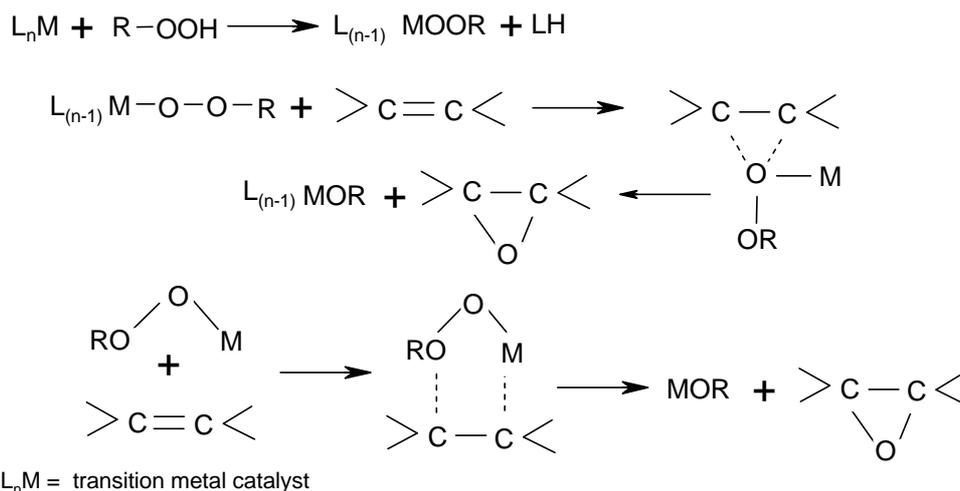
The epoxidation reactions were found to be accelerated to a great extent by the presence of catalysts such as benzyl triethyl ammonium hydroxide (BTEAH), molybdenum trioxide (MoO<sub>3</sub>) and vanadium (V) oxyacetyl acetonate [VO (acac)<sub>2</sub>]. The details of the catalysed epoxidation reactions are shown in table 2. The effect of catalyst was examined for the epoxidation of cinnamic acid as well as cyclohexene. [VO (acac)<sub>2</sub>] was found to be the most efficient catalyst among the three. The yield of epoxide was low with BTEAH. However the yield can be improved by increasing the reaction time. BTEAH being readily soluble in water, requires much simpler steps in the workup procedure when compared to [VO (acac)<sub>2</sub>]. Hence the use of BTEAH which is a phase transfer catalyst seems to suit the reaction, particularly with solid-phase reagents.

The polymeric byproduct was identified to be polymeric *t*-butyl alcohol resin with structure (3). The polymeric alcohol obtained in each epoxidation reaction was washed with several solvents and recycled in a single step to polymeric *t*-butyl hydroper-

**Table 2.** Catalysed epoxidation of olefins.

Catalyst	Reaction time (h) for		Isolated yield <sup>c</sup> (%) for	
	Substrate 1 <sup>a</sup>	Substrate 2 <sup>b</sup>	Substrate 1 <sup>a</sup>	Substrate 2 <sup>b</sup>
BTEAH	16	12	90	84
MoO <sub>3</sub>	14	14	82	84
VO (acac) <sub>2</sub>	12	10	86	89

<sup>a</sup>Cinnamic acid; <sup>b</sup>cyclohexene; <sup>c</sup>substrate to resin 1 : 2; solvent, dioxan, temperature, 70°C

**Scheme 3.** Probable mechanism for epoxidation of olefins.

oxide. Residual hydroperoxide capacity of the reagent was not affected considerably even after five cycles. Efficiency of the reagent remained more or less constant in terms of both yield and reaction time for the model epoxidation of cinnamic acid.

The reaction is assumed to follow first-order kinetics with respect to olefin and first order with respect to hydroperoxide, the overall reaction being a second-order process. Since large excess of the *t*-butyl hydroperoxide reagent was used it is assumed to follow pseudo first-order kinetics. The reaction is assumed to involve the nucleophilic attack of the olefin on the hydroperoxide followed by proton transfer probably by a concerted intramolecular process. The intermediate formed then decays to give the epoxide. The probable mechanism is depicted in scheme 3. The presence of electron-releasing groups on the olefin increased the rate of epoxidation, *cis* double bonds were attacked faster than *trans* double bonds.

### 3.3 Oxidation reactions

The polymeric *t*-butyl hydroperoxide resin (**4**) was employed to perform the oxidation of alcohols both

primary and secondary to aldehydes and ketones respectively in yields ranging from 54 to 85%. The alcohols used, the products isolated and reaction conditions are presented in table 3. The oxidation of alcohols by polymeric *t*-butyl hydroperoxide and its regeneration from the polymeric byproduct are shown in scheme 4. The course of the oxidation reaction was monitored by tlc. The 1,4-BDDMA crosslinked polymer support to which the *t*-butyl hydroperoxide functionality was attached exerted a marked improvement on oxidation rate when compared to the DVB crosslinked one.<sup>17</sup> Benzoin was oxidized to benzil in 85% yield within a period of 26 h while DVB crosslinked reagent produced only 82% benzil even after 36 h. Benzhydrol was also found to show a more or less similar extent of reaction as benzoin. However, cholesterol was oxidised only in 43 h. All substrates except cholesterol produced a single product on oxidation. When cholesterol was oxidized a small amount of epoxide was also obtained along with cholestenone as in the case of the DVB crosslinked polystyrene supported *t*-butyl hydroperoxide.<sup>17</sup> The reaction time was however reduced by about 14 h, when compared to the previous



**Table 4.** Catalysed oxidation using 1,4-BDDMA crosslinked polystyrene-bound *t*-butyl hydroperoxide resin.

Catalyst	Reaction time <sup>c</sup> for		Product yield <sup>c</sup>	
	Substrate 1 <sup>a</sup>	Substrate 2 <sup>b</sup>	Substrate 1 <sup>a</sup>	Substrate 2 <sup>b</sup>
BTEAH	14	12	82	85
MoO <sub>3</sub>	20	16	85	86
VO (acac) <sub>2</sub>	14	18	88	86

<sup>a</sup>Benzoin; <sup>b</sup>benzhydrol; <sup>c</sup>substrate to resin ratio, 1 : 5; solvent, chloroform; temperature, 45°C

catalysts like BTEAH. MoO<sub>3</sub> and [VO (acac)<sub>2</sub>] accelerated the epoxidation and oxidation reactions to a great extent. Dioxan was found to be the most suitable solvent for epoxidation while acetonitrile proved best for oxidation. Both the reactions studied were temperature dependent. A five-fold molar excess of the reagent showed maximum efficiency with regard to both epoxidation and oxidation reactions.

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