Comparison of the methods of the phase transfer catalysis and hydroperoxide in the epoxidation of 1,5,9-cyclododecatriene

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The process of the epoxidation of *cis,trans,trans*-1,5,9-cyclododecatriene (CDT) to 1,2-epoxy-5,9-cyclododecadiene (ECDD) with the 30% aqueous hydrogen peroxide under the phase transfer conditions and with *tert*-butyl hydroperoxide under the homogeneous conditions was investigated. Onium salts such as Aliquat® 336, Arquad® 2HT, methyltrioctylammonium bromide and the $\text{Na}_2\text{WO}_4/\text{H}_3\text{PO}_4$ catalyst system are very active under the phase transfer catalysis (PTC) conditions for the selective epoxidation of *cis,trans,trans*-1,5,9-cyclododecatriene (PTC method). These catalytic systems were found to be as active and selective as the homogeneous phase system $\text{Mo}(\text{CO})_6/\text{TBHP}$ (hydroperoxide method).

Keywords: epoxidation, 1,5,9-cyclododecatriene, phase transfer catalysis, hydrogen peroxide, *tert*-butyl hydroperoxide.

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INTRODUCTION

Epoxides are useful industrial intermediates such as e.g. propylene oxide which is used at the root of the synthesis of polyether polyols used for the production of polyurethane foams, and also used for the synthesis of propylene glycol for the elaboration of unsaturated polyester resins¹. Epoxides are also one of the most important and versatile intermediates for fine chemical and specialty chemicals synthesis such as epoxy resin, surfactants, paints, adhesives, surface coating agents, and in addition to a range of bioactive substances and flavour and fragrances².

Industrial production of epoxides usually employs a different oxygen source. However, the origin of the oxygen atom is rarely dioxygen itself, thus another peroxygen atom-donor oxidant must be used. Increasingly stringent environmental legislation as well as economic factors lead to a preference of the oxidations involving dioxygen, hydrogen peroxide, and alkyl hydroperoxides such as *tert*-butyl peroxide (TBHP)¹.

For the industrial synthesis, TBHP is probably one of the best oxygen sources because it is thermally stable, not very corrosive, relatively safe, and well soluble in a non-polar solvent. Using the TBHP after its activation by an appropriate transition metal complex leads to selective oxidations. However, that requires commercialisation or recycling of a large amount of the alcohol co-product. The *tert*-butyl alcohol, co-produced from *tert*-butyl hydroperoxide, can be blended with gasoline to inhibit the engine knock or dehydrated to make isobutylene for the conversion to methyl *tert*-butyl ether, also a gasoline blending component¹. Similarly, the 1-phenylethanol obtained as a co-product from the ethylbenzene hydroperoxide reaction is dehydrated to produce styrene.

On the other hand, there is a strong need for the development of alternative epoxidation methods that are cleaner, safer, and more environmentally friendly which employ safer oxidants and produce little waste. The processes should be efficient in both the energy and raw materials consumption and produce little waste³. From this point of

view the employment of hydrogen peroxide is an attractive option both on the environmental and the economic ground⁴. It is inexpensive, relatively safe and easy to handle, and produces water as the sole co-product of reduction. Several catalytic systems have been reported for the epoxidation of a wide range of alkenes with an aqueous hydrogen peroxide as an oxidant in the last two decades. Interesting results of olefine epoxidation were obtained with tungsten catalytic systems under the phase transfer catalysis conditions (PTC)⁵⁻⁸.

In this paper the results of the studies on the epoxidation of *cis,trans,trans*-1,5,9-cyclododecatriene (CDT) by the $\text{Mo}(\text{CO})_6/\text{TBHP}$ system (hydroperoxide method) and hydrogen peroxide under the phase transfer catalysis conditions (PTC method), have been presented.

MATERIALS AND METHODS

Chemicals

All experiments were performed with the following reagents: 46% of the TBHP solution in 1,2-dichloroethane (DCE), which was received in the extraction process of 70% TBHP water solution (Aldrich) by 1,2-dichloroethane⁹, an aqueous solutions of H_2O_2 (30%), phosphorous(V) acid (85%), and 1,2-dichloroethane were purchased from POCh, Poland; *cis,trans,trans*-1,5,9-cyclododecatriene (CDT), methyltrioctylammonium chloride (Aliquat® 336), tetraethylammonium bromide, methyltrioctylammonium bromide, and dodecane were obtained from Aldrich; sodium (tetraoxo)tungstate(VI), tetraethylammonium chloride, tetrapropylammonium bromide, tetrabutylammonium hydrogensulfate and dimethyl[diocetyl(76%) + dihexadecyl(24%)] ammonium chloride (Arquad® 2HT) were purchased from Fluka; tetrabutylammonium bromide and molybdenum hexacarbonyl, $\text{Mo}(\text{CO})_6$ (>95%) were obtained from Merck-Schuchardt.

Procedure

The catalytic epoxidation reactions with hydrogen peroxide as an oxidant were carried out under the pressure-low conditions in the apparatus described elsewhere¹⁰. In a typical experiment, sodium (tetraoxo)tungstate(VI), 30% aqueous H₂O₂, and 85% H₃PO₄ were added into the 25 cm³ flask at room temperature and then stirred for 15 min. Then the solution of ammonium salt in 1,2-dichloroethane was added dropwise with stirring. Stirring was continued for an additional 15 min. To the resultant solution, the weighed amount of CDT was added and then the flask was immersed in the ultrathermostat with a determined temperature. After reaching a desired temperature, 30% H₂O₂ was added dropwise. The reaction was maintained at 50°C for 45 min with vigorous stirring. After the reaction, the reaction mixture was rapidly cooled to room temperature in order to stop the reaction. The organic phase was analyzed chromatographically, and the concentration of CDT and ECDD in the product was determined. The H₂O₂ concentration in the aqueous phase was determined by an iodometric method¹¹.

All the experiments with the use of TBHP were performed according to the following procedure: the precisely weighed amount of CDT and the solution of TBHP in DCE were placed into the three-necked flask of the capacity of 25 cm³, fitted with a thermometer, reflux condenser and a mechanical mixer. The flask was later immersed in an oil bath. When the temperature of the reaction mixture reached 70°C, the molybdenum catalyst dissolved in the DCE was added immediately. After finishing the reaction, the flask with the post-reaction mixture was cooled quickly to room temperature to stop the reaction. The content of the flask was weighed and the chromatographic and iodometry analysis¹¹ was taken in order to make mass balance.

RESULTS AND DISCUSSION

PTC method

The investigation of the influence of the type of the phase transfer catalyst on the CDT epoxidation were per-

formed with the use of different ammonium salts (PTA): tetraethylammonium chloride, methyltriethylammonium chloride (Aliquat[®] 336), dimethyl[di-octadecyl(76%) + dihexadecyl(24%)]ammonium chloride (Arquad[®] 2HT), tetrabutylammonium hydrogensulfate, tetraethylammonium bromide, tetrapropylammonium bromide, tetrabutylammonium bromide, and methyltriethylammonium bromide. The initial parameters of the syntheses were as follows: the temperature of 50°C, the CDT:H₂O₂ mole ratio 1:1, the W:P:PTA mole ratio 5:2.5:1, the PTA content 0.7 mass %, and the reaction time was 45 min (Table 1).

In the presence of Aliquat[®] 336, Arquad[®] 2HT, and methyltriethylammonium bromide the maximum degree of the conversion of H₂O₂ (96.6 – 97.6 mole %) were obtained. In the presence of these salts the degrees of the conversion of CDT were also high (Table 1). However, the maximum ECDD selectivity both in relation to H₂O₂ and in relation to the CDT consumed were obtained in the presence of tetrabutylammonium hydrogensulfate. Somewhat lower selectivities were obtained in the presence of Aliquat[®] 336, Arquad[®] 2HT, tetrabutylammonium bromide, and methyltriethylammonium bromide. Other investigated salts were inactive as the phase transfer catalysts in the epoxidation of the CDT.

The studies demonstrate a considerable influence of the CDT:H₂O₂ mole ratio on the CDT epoxidation. The initial parameters of the syntheses were the following: PTA - Aliquat[®] 336, the PTA content 0.7 mass %, temperature 60°C, the W:P:PTA mole ratio 5:2.5:1, and the reaction time was 45 min (Table 2).

Initially, the yield of the ECDD increases with increase of the mole ratio of CDT to H₂O₂, then the yield was found to decrease with the increasing mole ratio. The maximum yield of the ECDD in relation to the CDT and H₂O₂ introduced, was obtained at the CDT:H₂O₂ mole ratio reaching 1:1 and 3:1, respectively.

At the study of the effect of the reaction time the initial parameters of the syntheses were as follows: PTA Aliquat[®] 336, the PTA content 0.7 mass %, the temperature 60°C, the CDT:H₂O₂ mole ratio 1:1, and the W:P:PTA mole ratio 5:2.5:1 (Table 3).

Table 1. The result of the CDT epoxidation under the PTC condition in relation to the ammonium salt type

	Type of ammonium salt							
	1	2	3	4	5	6	7	8
H ₂ O ₂ conversion [mole%]	0.86	97.56	97.15	17.59	1.51	1.73	6.87	96.59
CDT conversion [mole%]	8.22	78.33	82.41	20.69	13.97	2.04	8.90	80.70
ECDD selectivity/H ₂ O ₂ [mole%]	9.25	42.69	36.48	82.69	4.80	5.80	57.77	43.42
ECDD selectivity/CDT [mole%]	1.00	55.12	43.88	73.48	0.53	5.13	47.16	53.37

1 – Et₄NCl, 2 – Aliquat[®] 336, 3 – Arquad[®] 2HT, 4 – Bu₄NHSO₄, 5 – Et₄NBr, 6 – Pr₄NBr, 7 – Bu₄NBr, 8 – Me(C₈H₁₇)₃NBr

Table 2. The result of the CDT epoxidation under the PTC condition in relation to the CDT:H₂O₂ ratio

	CDT:H ₂ O ₂ mole ratio				
	0.5	1	2	3	5
Yield of ECDD/H ₂ O ₂ [mole%]	5.30	38.31	62.19	68.92	59.62
Yield of ECDD/CDT [mole%]	11.15	40.13	32.17	24.10	11.87

Table 3. The result of the CDT epoxidation under the PTC condition in relation to the reaction time

	Reaction time, min					
	15	30	45	60	90	180
W ECDD/H ₂ O ₂ [mole%]	40.12	41.54	42.09	41.18	38.31	28.21
W ECDD/CDT [mole%]	42.38	42.43	43.11	42.27	40.13	29.26

The changes in the reaction time do not have a significant influence on the yield of the ECDD in relation to the CDT and H_2O_2 introduced within a period of 15 – 60 min. However, after 60 min the yields decreased rapidly. It is worthy of note that the maximum yield was obtained in a relatively short reaction time of 15 – 45 min.

The changes of the ECDD yield in relation to the temperature are shown in Table 4. The initial parameters of the syntheses were as follows: PTA Aliquat® 336, the PTA content 0.7 mass %, the CDT: H_2O_2 mole ratio 1:1, the W:P:PTA mole ratio 5:2.5:1, and the reaction time 45 min.

As the temperature was increased from 20 to 50°C, an increase of the ECDD yield both in relation to the CDT and H_2O_2 introduced was observed from 26.03 to 43.17 mole% and from 25.44 to 41.65 mole%, respectively. Somewhat lower ECDD yields were obtained at higher temperatures.

A decrease of the ECDD yields with increasing the temperature (above 50°C) and time (above 60min) probably results from a considerable reactivity of oxirane ring.

Hydroperoxide method

The influence of the temperature in the range of 70–110°C, at the CDT:TBHP mole ratio 3:1, the $\text{Mo}(\text{CO})_6$:TBHP mole ratio $0.5 \cdot 10^{-3}$, and the reaction time 120 min were investigated (Table 5).

At the preliminary selected initial parameters, the increase of the reaction temperature increased the ECDD yield in relation to both the CDT and TBHP introduced. The highest yield of the ECDD is achieved at 90°C.

The influence of the mole ratio of CDT to TBHP on the yield of the ECDD in relation to the CDT and TBHP introduced is shown in Table 6. The studies were carried out at the temperature 90°C, the $\text{Mo}(\text{CO})_6$:TBHP mole ratio $0.5 \cdot 10^{-3}$, and the reaction time 120 min.

The increase of the CDT:TBHP mole ratio decreases the ECDD yield in relation to the CDT introduced. A decrease of the ECDD yield is associated with a decrease of the CDT conversion. With increasing the CDT:TBHP mole ratio the concentration of the TBHP in reaction mixtures decreases. This causes a decline of the epoxidation rate. However, the ECDD yield in relation to the TBHP introduced is increased with the increase of the CDT:TBHP mole ratio.

The changes of the $\text{Mo}(\text{CO})_6$:TBHP mole ratio in the range 0.0001 – 0.001 do not have a significant influence on the yield of the ECDD.

The influence of the reaction time on the yield of the ECDD in relation to the CDT and TBHP introduced is shown in Table 7. The studies were carried out at the

Table 4. The result of the CDT epoxidation under the PTC condition in relation to the temperature

	Temperature, °C					
	20	35	50	60	70	80
W ECDD/ H_2O_2 [mole%]	25.44	37.48	41.65	38.31	38.74	37.92
W ECDD/CDT [mole%]	26.03	38.12	43.17	40.13	39.75	38.66

Table 5. The result of the CDT epoxidation by the $\text{Mo}(\text{CO})_6$ /TBHP system in relation to the reaction temperature

	Temperature [°C]				
	70	80	90	100	110
CDT/TBHP mole ratio	3:1	3:1	3:1	3:1	3:1
$\text{Mo}(\text{CO})_6$ /TBHP mole ratio	0.0005	0.0005	0.0005	0.0005	0.0005
Reaction time [h]	2	2	2	2	2
ECDD yield /TBHP [mole%]	68.8	76.9	78.9	77.0	76.9
ECDD yield /CDT [mole%]	22.9	25.7	26.6	26.3	26.1

Table 6. The result of the CDT epoxidation by the $\text{Mo}(\text{CO})_6$ /TBHP system in relation to the CDT/TBHP mole ratio

	CDT/TBHP mole ratio				
	1	3	5	7	10
Temperature [°C]	90	90	90	90	90
$\text{Mo}(\text{CO})_6$ /TBHP mole ratio	0.0005	0.0005	0.0005	0.0005	0.0005
Reaction time [h]	2	2	2	2	2
ECDD yield /TBHP mole%	50.2	26.6	16.6	12.5	9.5
ECDD yield /CDT [mole%]	48.9	78.9	83.3	89.5	96.4

Table 7. The result of the CDT epoxidation by the $\text{Mo}(\text{CO})_6$ /TBHP system in relation to the reaction time

	Reaction time [min]		
	60	120	180
CDT/TBHP mole ratio	3:1	3:1	3:1
$\text{Mo}(\text{CO})_6$ /TBHP mole ratio	0.0005	0.0005	0.0005
Temperature [°C]	70		
ECDD yield /TBHP [mole%]	54.4	68.0	78.5
ECDD yield /CDT [mole%]	18.3	22.9	26.4
Temperature [°C]	90		
ECDD yield /TBHP [mole%]	77.6	78.9	79.9
ECDD yield /CDT [mole%]	26.1	26.6	26.9
Temperature [°C]	110		
ECDD yield /TBHP [mole%]	76.0	76.9	77.4
ECDD yield /CDT [mole%]	25.8	26.1	26.3

temperatures 70, 90, and 110°C, at the CDT:TBHP mole ratio 3:1, and at the Mo(CO)₆:TBHP mole ratio 0.5·10⁻³.

At low temperatures (70°C), an increase in the reaction time increases the ECDD yield in relation to both the CDT and TBHP introduced. At higher temperatures (90 and 110°C) prolongation of the reaction time do not have a significant influence on the yield of the ECDD.

CONCLUSIONS

The two phase catalytic epoxidation of *cis,trans,trans*-1,5,9-cyclododecatriene with the aqueous hydrogen peroxide in the presence of the phase transfer catalyst was found to be an equally good method as the homogeneous catalytic epoxidation by the TBHP in the presence of Mo(CO)₆. Comparison of the influence of the technological parameter of these two methods of the 1,2-epoxy-5,9-cyclododecadiene synthesis indicates that the epoxidation proceeds somewhat better with the use of the TBHP.

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