Mesoporous Ce–Zr mixed oxides for selective oxidation of styrene in liquid phase

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Abstract

This work reports the synthesis of mesoporous $\text{Ce}_{1-x}\text{Zr}_x\text{O}_{2-\delta}$ ($x=0.5$ and 0.8) mixed oxides with distinct Ce/Zr mole ratio by inverse micelle template method and their catalytic exploration for epoxidation of styrene in isopropanol solvent using TBHP as the oxidant. Among various catalysts investigated, the $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ combination catalyst exhibited best catalytic activity with ~98% conversion and ~90% selectivity to styrene epoxide. The synthesized Ce–Zr mixed oxide catalysts were characterized by various state-of-the-art techniques. Characterization studies revealed that Ce/Zr mole ratio has an imperative influence on the physicochemical properties such as surface area, oxygen vacancy concentration, and redox nature. Interestingly, catalytic efficiency was significantly improved with the increase of Ce and decrease of Zr content in the Ce–Zr mixed oxides. Catalytic efficiency and distribution of the products for styrene oxidation under various conditions such as reaction time, solvent, temperature, and styrene to TBHP mole ratio were also evaluated. Reusability of the highly active $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ mixed oxide catalyst was also demonstrated.

Keywords Ceria · Styrene · Styrene epoxide · Oxygen vacancy · Redox properties

Introduction

Catalytic oxidation of styrene is considered as a popular reaction in the chemical and petrochemical industries, because the delivered oxygenated products are expedient and key intermediates for the synthesis of plasticizers, perfumes, drugs, agrochemicals, and pharmaceuticals [1, 2]. The literature assessment demonstrates that the catalytic oxidation of styrene produces benzaldehyde along with styrene epoxide [3]. Thus, achieving high selectivity to one of these products has been an everlasting topic in the field of heterogeneous catalysis. In the present investigation, our primary aim was to achieve high selectivity to styrene epoxide rather than benzaldehyde, owing to numerous applications of the product [4]. For example, epoxides are considered as key building blocks for the production of various organic

products including glycols, alcohols, carbonyl compounds, alkanolamines, and polymers [5]. Traditionally, styrene oxidation was accomplished by employing organic peracid as an oxidant. However, utilization of such kinds of oxidants is no longer acceptable in the chemical industry due to their harmful nature to the environment [6].

In general, hydrogen peroxide (H_2O_2), molecular oxygen (O_2), and tert-butyl hydroperoxide (TBHP) have been used as optional oxidants to explore the styrene oxidation [3, 7, 8]. But, the selection of suitable oxidant is crucial to conduct the oxidation of styrene, because it shows enormous influence on the conversion and selectivity of reaction products. It is evident from the literature that TBHP has been identified as one of the promising oxidants for styrene oxidation since it relatively furnished high selectivity to styrene epoxide compared to other oxidants [9]. The disadvantage associated with H_2O_2 is that it has the ability to decompose at high temperature, thereby shows poor selectivity in the styrene oxidation [10]. In contrast, high oxidation capability of molecular O_2 could provide more side products. Therefore, in this study, TBHP has been selected as the appropriate oxidant to perform the styrene epoxidation reaction.

In recent times, numerous heterogeneous catalysts have been explored for styrene oxidation using appropriate

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oxidants. It was found from these studies that the selectivity of products is strongly dependent on the nature of the catalytic system. For instance, Co_3O_4 [11], $\text{Mg}/\text{Co}/\text{Al}$ hydro-talcites [8], polyoxometalates [12], $\text{Mg}-\text{Cu}$ ferrite [13], and BaFe_2O_4 [7] catalysts are highly selective to benzaldehyde product. In contrast, Mn and Co oxides [3], Cu -doped hydro-talcites [14], $\text{Cu}-\text{HMS}$ [9], $\text{CuO}/\text{CoAl-HT}$ [15], $\text{Co}/\text{nitrogen}$ doped hollow spheres [16], and $\text{Au}-\text{TiO}_2$ nano particles [17] showed high selectivity to styrene epoxide. Owing to their attractive properties such as high surface area, tunable pore size, pore volume, and morphology, mesoporous metal oxides are playing a key role in the area of catalysis [3, 18]. For example, higher surface area of mesoporous metal oxides accompanying more number of active sites direct to excellent activity than that of their non-porous analogues [19, 20]. One of the favourable properties of mesoporous metal oxides is the high adsorption ability of the substrate on the catalytic surface which results in excellent catalytic performance [21].

Ceria (CeO_2)-based mixed oxides have been employed extensively as potential candidates for many catalytic applications due to their outstanding properties [22–25]. Doping of Zr into the CeO_2 lattice is an effective strategy to manipulate the physicochemical properties like oxygen storage capacity, redox property, concentration of oxygen vacancies, and thermal stability of the ceria, thereby enlarging for better catalytic activity in many oxidation reactions [26, 27]. The main goal of this work was to explore the consequence of Ce/Zr mole ratio and morphology of mixed oxides on the physicochemical properties and catalytic activity for styrene oxidation. Accordingly, we have synthesized mesoporous $\text{CeO}_2-\text{ZrO}_2$ mixed oxides with different Ce/Zr mole ratios by inverse micelle template method. A systematic physicochemical characterisation of the synthesized catalysts has been accomplished using X-ray diffraction (XRD), Brunauer–Emmet–Teller (BET) surface area, Raman spectroscopy, X-ray photoelectron spectroscopy (XPS), Field emission scanning electron microscopy (FE-SEM), and H_2 -temperature programmed desorption (H_2 -TPR) techniques.

Experimental

Catalyst preparation

Mesoporous $\text{Ce}_{1-x}\text{Zr}_x\text{O}_{2-\delta}$ ($x=0.5$ and 0.8) mixed oxides with different Ce/Zr mole ratios along with pure CeO_2 and ZrO_2 were synthesized by adopting an inverse micelle method [28]. In a typical synthesis, appropriate amounts of corresponding metal precursors, namely, $\text{Ce}(\text{NO}_3)_4 \cdot 6\text{H}_2\text{O}$ (Aldrich, AR grade), $\text{ZrO}(\text{NO}_3)_2 \cdot x\text{H}_2\text{O}$ (Sigma Aldrich, AR grade), and butanol were taken in a beaker. Later,

desirable amount of poly(ethylene glycol)block-poly(propylene glycol)-block-poly(ethylene glycol) (P123) and concentrated nitric acid (HNO_3) were added to the above solution and stirred at room temperature until it completely dissolved. The resulting clear solution was heated at 393 K for 3 h under stirring condition. The obtained black powder was washed with excess ethanol, centrifuged, and dried in a vacuum oven overnight. At the end, the dried black powder was subjected to a heating cycle. First, they were heated at 423 K for 12 h and cooled down to room temperature under ambient conditions. The obtained material was then subjected to the following heating cycle to obtain differently calcined materials: 523 K for 3 h, 623 K for 2 h, 723 K for 2 h, and 823 K for 1 h, respectively.

Catalyst characterization

Powder XRD patterns of the prepared samples were recorded on a Rigaku diffractometer using $\text{Cu K}\alpha$ radiation (1.540 Å), operated at 40 kV and 40 mA. The diffractograms were recorded in the 2θ range of $10-80^\circ$ with a 2θ step size of 0.02° and a step time of 2.4 s. The XRD phases present in the samples were identified with the help of Powder Diffraction File-International Centre for Diffraction Data (PDF-ICDD). The average crystallite size and lattice parameter of the prepared samples were calculated using the full width at half maximum (FWHM) of the (111) peak using the Debye–Scherer equation and the cubic indexation methods, respectively.

The Brunauer–Emmet–Teller (BET) surface area of the samples was determined by N_2 adsorption on a Micromeritics Gemini 2360 instrument. Before the analysis, the samples were oven-dried at 393 K for 12 h and flushed with argon gas for 2 h to remove any surface-adsorbed residue. Surface area was calculated by utilizing the desorption data. Raman spectra of the samples were recorded on a Horiba Jobin–Yvon HR800 Raman spectrometer equipped with a liquid-nitrogen cooled charge coupled device (CCD) detector and a confocal microscope. The emission line at 638 nm from Ar^+ laser (Spectra Physics) was focused on the sample under the microscope with the diameter of the analysed spot being $\sim 1 \mu\text{m}$, under the ambient conditions. The time of acquisition was adjusted according to the intensity of Raman scattering. The wavenumber values obtained were precise to within 2 cm^{-1} .

The XPS measurements were performed on a PHI 5400 instrument. $\text{Al K}\alpha$ radiation (1486.6 eV) X-ray source was used for XPS analysis and at a pressure lower than 10^{-7} Torr. The sample preparation for XPS analysis involves mounting of a few grams of sample on a carbon tape supported with a silica plate. The resultant plate supported with a holder is placed in the vacuum chamber of Thermo K-Alpha XPS instrument before the start of analysis. Thermo-Avantage Software was

used for XPS analysis of the prepared samples. The binding energies of Ce, Zr, and O were charge-corrected with respect to the adventitious carbon (C 1 s) peak at 284.6 eV. Flood gun was used to remove the static charge that developed on the sample surface during XPS analysis. FE-SEM studies were performed using a JEOL-7610F electron microscope to examine surface morphology, and the energy dispersive X-ray analysis (EDX) was used to determine the elemental composition of samples with OXFORD Inca detector interfaced at 5 kV.

The reducibility of the synthesized samples was estimated by H_2 -TPR analysis, using a thermal conductivity detector of a gas chromatograph (Schimadzu). Approximately 30 mg of the samples were loaded in an isothermal zone of the reactor and were treated in a helium gas flow at 400 K followed by cooling to room temperature. After that, flow of the He was switched to 5% H_2 /Ar with a rate of 20 mL min^{-1} and the temperature was linearly increased to 1073 K at a heating ramp of 5 K min^{-1} , keeping all the parameters un-changed. The hydrogen consumption during the reduction process was estimated by passing the effluent gas through a molecular sieve trap to remove the produced water and was analysed by gas chromatography using a TCD.

Catalytic activity test

The catalytic efficiency of prepared mesoporous mixed oxides was evaluated for liquid phase oxidation of styrene using TBHP as the oxidant. Typically, a mixture of styrene (4 mmol), TBHP, isopropanol (3 ml), and the catalyst (50 mg) was charged into a 10-ml round-bottom flask. Then, the reaction mixture in the round-bottom flask was agitated at 1000 rpm and the process was continued for desired time at 353 K. After completion of the reaction, the liquid products and the catalyst were separated by centrifugation. The collected products were analysed by GC equipped with a BP-20 (wax) capillary column and a flame ionization detector. The products were confirmed by GC–MS equipped with a DB-5 capillary column and mass selective detector. The reaction products were also confirmed by injecting the corresponding authentic compounds in the GC. To check the reusability of the catalysts, after completion of reaction the catalyst was separated by centrifugation and washed with ethanol to remove the products adhered to the surface of the catalyst. The resulting sample was dried at 393 K for 12 h, and then reused directly for the next run.

Results and discussion

XRD and BET analysis

Figure 1 shows the XRD profiles of calcined $Ce_{1-x}Zr_xO_{2-8}$ ($x=0.5$ and 0.8) mixed oxides along with bare CeO_2 and

ZrO_2 . Interestingly, distinct XRD patterns were observed for both mesoporous mixed oxide samples. The high Ce content $Ce_{0.8}Zr_{0.2}O_2$ sample displayed XRD lines corresponding to (111), (200), (220), (311), (331), and (400) planes, which are characteristic of the fluorite like cubic structure of CeO_2 [29, 30]. Further, no XRD lines related to ZrO_2 phase were noticed for this sample indicating the formation of solid solution. In contrast, we observed XRD lines pertaining to mixed phases of CeO_2 and ZrO_2 in the XRD profile of $Ce_{0.5}Zr_{0.5}O_2$ sample indicating that Zr is not completely doped into the CeO_2 lattice. On the other hand, mesoporous CeO_2 showed XRD lines related to (111), (200), (220), (311), (331), and (400) planes, confirming the fluorite cubic structure of CeO_2 . On the similar lines, tetragonal ZrO_2 phase was identified in the XRD profile of mesoporous ZrO_2 [29]. No big change in the XRD patterns of the catalyst after catalytic runs was noted indicating that there is no change in the catalyst structure during catalytic evaluation.

To determine the specific surface area of the prepared mixed oxides, we have undertaken the BET analysis of the samples. The achieved BET results of the corresponding samples are illustrated in Table 1. Mixed oxides are expected to show larger BET surface areas than their individual single oxides, which is due to the cooperative effect of respective oxides in the final mixed oxide (Table 1). The existence of strong synergetic interaction between the foreign metal oxide and the host metal oxide is the key reason for enhancement in the surface area for mixed oxides. Interestingly, the specific surface area of the doped CeO_2 was found to enhance

Fig. 1 Powder XRD profiles of pure CeO_2 , $Ce_{0.5}Zr_{0.5}O_2$, and $Ce_{0.8}Zr_{0.2}O_2$ samples

Table 1 BET specific surface area (S) and Raman shift of various catalysts

Sample	S (m ² g ⁻¹)	Raman shift F _{2g} (cm ⁻¹)
CeO ₂	38.4 ± 3	457
Ce _{0.5} Zr _{0.5} O ₂	71.6 ± 2	465
Ce _{0.8} Zr _{0.2} O ₂	137.1 ± 2	473
Ce _{0.2} Zr _{0.8} O ₂	50.5 ± 2	–
ZrO ₂	28.2 ± 3	–

Fig. 2 Raman spectra of pure CeO₂, Ce_{0.5}Zr_{0.5}O₂, and Ce_{0.8}Zr_{0.2}O₂ samples

with the increase of Ce content. The specific surface areas of ZrO₂, CeO₂, Ce_{0.5}Zr_{0.5}O₂, and Ce_{0.8}Zr_{0.2}O₂ samples were found to be ~ 28.2, ~ 38.4, ~ 71.6, and ~ 137.1 m²/g, respectively.

Raman studies

Raman spectroscopy was employed to understand the structural properties of mesoporous Ce–Zr mixed oxides. Raman spectra of Ce_{0.5}Zr_{0.5}O₂ and Ce_{0.8}Zr_{0.2}O₂ mixed oxides along with CeO₂ are shown in Fig. 2. We observed a sharp intense Raman band located at 465 cm⁻¹ for all the samples, which indicates the presence of the Raman-active F_{2g} mode of fluorite structured CeO₂ with the space group Fm3m (Supported by XRD results) [31]. Further, we did not notice any Raman bands related to ZrO₂ phase in the investigated Raman region for both Ce–Zr mixed oxides. In particular,

F_{2g} band of Ce_{0.5}Zr_{0.5}O₂ and Ce_{0.8}Zr_{0.2}O₂ is significantly shifted and broadened with respect to pure CeO₂, which is more pronounced in the case of Ce_{0.8}Zr_{0.2}O₂ mixed oxide. The shift in F_{2g} mode is probably due to the variation in the M–O vibration frequency caused by the incorporation of Zr⁴⁺ ions into the CeO₂ lattice. Ceria-based samples typically show a Raman band in the range of ~ 580–650 cm⁻¹, corresponding to the presence of oxygen vacancy defects in the CeO₂ [32, 33]. To our delight, only high Ce content sample i.e. Ce_{0.8}Zr_{0.2}O₂ showed a prominent Raman peak at ~ 620 cm⁻¹ indicating the presence of oxygen vacancies. It is a well-established fact in the literature that oxygen vacancies play a key role in the oxidation reactions.

FE-SEM and XPS analysis

Morphological evolution of the synthesized samples was scrutinized by FE-SEM. The corresponding FE-SEM images of various catalysts are shown in Fig. 3. It can be clearly seen from Fig. 3 that pure CeO₂ and ZrO₂ show mesoporous structure. Interestingly, no apparent variation in the structure of Ce–Zr mixed oxides with different Ce/Zr ratios was noticed. This result demonstrates that the introduction of dopant into the CeO₂ lattice did not alter the original morphology of ceria. It is clear from the figure that both Ce–Zr mixed oxides, namely, Ce_{0.5}Zr_{0.5}O₂ and Ce_{0.8}Zr_{0.2}O₂ show mesoporous structure.

Figure 4 represent the deconvoluted Ce 3d core level spectra of Ce–Zr mixed oxides including pure CeO₂. The Ce 3d XP spectra are complex, which is due to the mixing of O 2p and Ce 4f levels. The peaks represented with u indicate 3d_{3/2} spin–orbit states and those labelled by ‘v’ correspond to 3d_{5/2} contributions. As shown in Fig. 4, the peaks labelled by u’ and v’ represent the Ce³⁺ with the electronic configuration of 3d¹⁰4f¹ and other bands labelled with u, u’, u’’, v, v’’, and v’’’ represent the 3d¹⁰4f⁰ electronic state corresponding to Ce⁴⁺. The presence of both the ions i.e. Ce⁴⁺ and Ce³⁺ over the surface of the samples indicates the redox behaviour of prepared samples [34]. Interestingly, the binding energy of the mesoporous mixed oxide was markedly shifted to lower binding energy side compared to that of pure CeO₂. This interesting observation could be attributed to modification of Ce–O environment caused by dopant ions. Figure 5a displays the O 1s XP spectra of mesoporous mixed oxides along with CeO₂. Irrespective of the sample, three peaks at ~ 528.1, ~ 532.5, and ~ 534.3 eV were detected, which could be attributed to lattice oxygen, adsorbed oxygen species of hydroxyl groups, and adsorbed molecular water and/or carbonate species [35]. Two bands at 180.5–181.40 eV (3d_{3/2}) and 182.0–182.5 eV (3d_{5/2}) were observed for mesoporous Ce_{0.5}Zr_{0.5}O₂ and Ce_{0.8}Zr_{0.2}O₂

Fig. 3 FE-SEM images of **a** CeO_2 , **b** $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$, and **c** $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ samples

Fig. 4 Ce 3d XP spectra of pure CeO_2 , $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$, and $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ samples

catalysts (Fig. 5B). This observation confirms the presence of Zr^{4+} ions in the synthesized catalysts [36].

H₂-TPR studies

Figure 6 shows the H₂-TPR profiles of mesoporous $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$ and $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ mixed oxides including the pure CeO_2 for the purpose of comparison. Typically pure CeO_2 exhibits two reduction peaks at ~794 K and ~1063 K for surface and bulk reductions, respectively [37]. It is proven that the incorporation of metal ions into the ceria lattice remarkably improves its reducibility. It is obvious from the figure that the surface and bulk reduction peaks of $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$ and $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ are considerably shifted to lower temperatures compared to pure CeO_2 indicating favourable role of dopant to improve the redox ability of CeO_2 . Particularly in the case of $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$, we observed a remarkable decrease in the surface (618 K) and bulk (748 K) reduction temperatures compared to that of $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$ and CeO_2 samples. This striking observation clearly demonstrates the key role of Ce/Zr mole ratio on the reducibility of the samples. Usually redox property of the catalyst is closely associated with the formation of oxygen vacancies, which are likely to play a crucial role in the oxidation reactions. Thus, we expect more number

Fig. 5 a O 1s and b Zr 3d XP spectra of pure CeO_2 , $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$, and $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ samples

Fig. 6 H_2 -TPR profiles of pure CeO_2 , $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$, and $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ samples

of oxygen vacancies in $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ sample in line with the Raman results.

Catalytic activity

Screening of catalysts

To explore the catalytic applicability of our newly designed mesoporous mixed oxides for oxidation of olefins, styrene

Scheme 1 Catalytic oxidation of styrene

was selected as a model reaction. The catalytic oxidation of styrene produces styrene epoxide and benzaldehyde products which are identified by GC–MS (Scheme 1). Optimization of the reaction conditions is crucial to obtain high activity and selectivity of the target product. In this direction, initially we have performed the screening of all synthesized catalysts, namely, $\text{Ce}_{0.5}\text{Zr}_{0.5}\text{O}_2$, $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$, CeO_2 , and ZrO_2 for oxidation of styrene. The reaction was conducted at 353 K with isopropanol and TBHP as oxidant and solvent, respectively, for 8 h and the obtained results are illustrated in Table 2. Blank experiment gave poor conversion indicating the need of a catalyst to progress the styrene oxidation (Table 2, entry 1). Pure CeO_2 and ZrO_2 oxides showed ~36 and ~24% conversion with ~80 and ~68% selectivity to styrene epoxide, respectively (Table 2, entries 2 and 3). This result indicates that CeO_2 is relatively active and selective to styrene epoxide compared to that of ZrO_2 .

Table 2 Oxidation of styrene to styrene epoxide (SO) and benzaldehyde (BA) over Ce–Zr mixed oxide catalyst with different Ce/Zr mole ratios and other catalysts

Entry	Catalyst	Conversion (%)	Selectivity to SO (%)	Selectivity to BA (%)	Others (%)
1	Blank	4.5	–	–	–
2	CeO ₂	36	80	11	9
3	ZrO ₂	24	68	9	23
4 ^a	CeO ₂	28	70	18	12
5 ^b	ZrO ₂	20	65	10	25
6	Ce _{0.5} Zr _{0.5} O ₂	76	83	12	5
7	Ce _{0.8} Zr _{0.2} O ₂	98	90	8	2
8 ^c	Ce _{0.8} Zr _{0.2} O ₂	75	70	15	15

Reaction conditions: Styrene (4 mmol), catalyst (100 mg), TBHP oxidant, isopropanol (5 ml), time (8 h), and temperature (353 K)

^{a,b}Nanocrystalline oxides prepared by co-precipitation method

^cCe–Zr mixed oxide prepared by co-precipitation method

To know the significance of the morphology of the samples, we have conducted the reaction using nanocrystalline CeO₂ and ZrO₂ catalysts under identical conditions that provided ~28 and ~20% styrene conversion, respectively, (Table 2, entries 4 and 5), confirming the key role of catalyst morphology for styrene oxidation.

Later, we studied the influence of Ce/Zr mole ratio towards oxidation of styrene. For this purpose, Ce–Zr mixed oxides with different Ce/Zr mole ratios were tested under same reaction conditions and the obtained results are compiled in Table 2. As can be noted from the Table 2, both conversion and selectivity are significantly varied with the change of Ce/Zr mole ratio (Table 2). The Ce_{0.5}Zr_{0.5}O₂ catalyst provided ~76% conversion with ~83% and 12% selectivity to styrene epoxide and benzaldehyde, respectively (Table 2, entry 6). As the concentration of Ce increases, the conversion of styrene increased from ~83 to ~98% with ~90 and ~8% selectivity of styrene epoxide and benzaldehyde, respectively (Table 2, entry 7). A high catalytic efficacy of Ce_{0.8}Zr_{0.2}O₂ mixed oxide could be attributed to larger BET surface area, more defect sites (oxygen vacancies), and high reducibility supported by XRD, BET, FE-SEM, and H₂-TPR studies. Meanwhile, to confirm the influence of morphology, nanocrystalline Ce_{0.5}Zr_{0.5}O₂ prepared by coprecipitation method was also tested for styrene oxidation under same reaction conditions and it showed relatively less activity than the mesoporous Ce_{0.8}Zr_{0.2}O₂ mixed oxide (Table 2, entry 8).

Fig. 7 Effect of solvent on the oxidation of styrene to styrene epoxide (SO) and benzaldehyde (BA) over Ce_{0.8}Zr_{0.2}O₂ catalyst. Reaction conditions: Styrene (4 mmol), time (8 h), temperature (353 K), catalyst (50 mg), TBHP, and isopropanol (3 mL)

Effect of solvent

To understand the role of polarity of solvent on styrene oxidation, variety of solvents were explored at 353 K for 8 h using TBHP as the oxidant with mesoporous Ce_{0.8}Zr_{0.2}O₂ mixed oxide catalyst and the achieved results are shown in Fig. 7. From the figure it is believed that polarity of solvent highly influences the activity and selectivity of the catalysts. Use of non-polar solvents like hexane shows ~62% styrene conversion, and this is due to the low solubility of the styrene in hexane solvent. In the case of polar solvents, four different solvents were tested and the results reveal an enhancement in the conversion of styrene with the increase of solvent polarity. Interestingly, we observed ~73, ~79, 86, and ~98% conversion for 1,4-dioxane, acetone, acetonitrile, and isopropanol solvents, respectively. A high activity and selectivity of styrene epoxide (~90%) was achieved with isopropanol solvent. High oxygen solubility in isopropanol solvent is likely to be the probable reason for this result. Therefore, we have considered isopropanol as the suitable solvent to optimize the other reaction parameters for this reaction.

Effect of reaction temperature

The effect of reaction temperature on the oxidation of styrene was examined in the range of 323–363 K over the highly active Ce_{0.8}Zr_{0.2}O₂ catalyst using TBHP as the oxidant in isopropanol solvent. The obtained results are presented in Fig. 8. As expected, conversion of styrene is increased with the raise of reaction temperature. Obviously, reaction temperature had notable effect on styrene

Fig. 8 Effect of reaction temperature on the oxidation of styrene to styrene epoxide (SO) and benzaldehyde (BA) over $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ catalyst. Reaction conditions: Styrene (4 mmol), time (8 h), catalyst (50 mg), TBHP, and isopropanol (3 mL)

oxidation in terms of conversion and selectivity of the target product. It can be clearly seen that the conversion of styrene gradually increases from ~34 to ~98% with the increase of reaction temperature from 323 to 353 K. Particularly, there was no significant variation in the selectivity of styrene epoxide (~88–92%) with the reaction temperature. Further increasing the temperature from 353 to 363 K, no considerable variation in the conversion of styrene (~98.5%) was observed but the selectivity of styrene epoxide was slightly dropped to ~88%. Therefore, the 353 K is considered as an optimum temperature for further studies.

Fig. 9 Effect of reaction time on the selective oxidation of styrene to styrene epoxide (SO) and benzaldehyde (BA) over $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ catalyst. Reaction conditions: Styrene (4 mmol), temperature (353 K), catalyst (50 mg), TBHP, and isopropanol (3 mL)

Effect of reaction time

The effect of reaction time on the activity and distribution of products for styrene oxidation was studied over the $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ catalyst at 353 K using TBHP as the oxidant in isopropanol solvent and the obtained results are compiled in Fig. 9. As can be seen from the figure, conversion of styrene increases with the increase of reaction time. The conversion as well as selectivity of styrene epoxide was found to increase with the increasing of reaction time from 2 to 8 h. The achieved styrene conversions were ~18, ~44, ~68, and ~98% for 2, 4, 6, and 8 h of reaction times, respectively. Further increase of reaction time from 8 to 10 h, no variation in the conversion of styrene was observed. Thus, we consider 8 h as the optimum reaction time for the oxidation of styrene.

Effect of styrene/TBHP molar ratio

The influence of styrene/TBHP molar ratio was examined for oxidation of styrene at 353 K for 8 h and the obtained results are shown in Fig. 10. The styrene conversion was found to be low at low styrene/TBHP mole ratio due to less availability of reactive oxygen to interact with the substrate. Conversion of styrene considerably increases with the increase of styrene/TBHP molar ratio, which is due to the fact that more number of TBHP molecules adsorb on the active sites, thereby leading to higher activity by interacting with more number of styrene molecules. The achieved styrene conversions are ~45, ~66, ~98, and ~99% for 1:0.50, 1:1, 1:2, and 1:4 mol ratios, respectively. Interestingly, we found high selectivity to styrene epoxide at all ratios. Therefore, these promising results clearly illustrated the remarkable role of

Fig. 10 Effect of styrene/TBHP mole ratio on styrene oxidation to styrene epoxide (SO) and benzaldehyde (BA) over $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ catalyst. Reaction conditions: Styrene (4 mmol), time (8 h), temperature (353 K), catalyst (50 mg), TBHP, and isopropanol (3 mL)

Fig. 11 Recyclability of $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ catalyst for styrene oxidation to styrene epoxide (SO) and benzaldehyde (BA). Reaction conditions: Styrene (4 mmol), time (8 h), temperature (353 K), catalyst (50 mg), TBHP, and isopropanol (3 mL)

cyclohexene/TBHP molar ratio for this reaction in terms of conversion and distribution of products.

Reusability

Finally, we have studied the efficiency of $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ catalyst up to 6 cycles to assess its reusability for styrene oxidation using TBHP as the oxidant in isopropanol solvent. The obtained results are presented in Fig. 11. The fixed reaction parameters for these studies were as follows: styrene (4 mmol), catalyst amount (50 mg), isopropanol (3 mL), reaction time (8 h), and temperature (353 K). After completion of each cycle, the solid catalyst was recovered from the reaction mixture by means of centrifugation, washed several times with ethanol to remove reaction gradients and then dried at 423 K for 2 h. Remarkably no variation in the catalytic activity of $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ was found up to 6 cycles for the styrene oxidation. The achieved styrene conversions were ~98, ~97.5, ~97.1, ~96.8, ~96.5 and ~95% for 1st, 2nd, 3rd, 4th, 5th and 6th cycles, respectively. Also, no considerable variation in the selectivity of styrene epoxide product was observed with the repeated use of catalyst.

Conclusion

In this work, mesoporous Ce–Zr mixed oxides with different Ce/Zr mole ratios along with CeO_2 and ZrO_2 were synthesized by inverse micelle template self-assembly method, characterized by various techniques and evaluated for styrene oxidation. Characterization studies reveal that the Ce/Zr mole ratio precisely modifies the physicochemical properties

of the Ce–Zr metal oxide catalysts. Among the investigated catalysts, the $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ combination exhibited high styrene conversion (~98%) followed by $\text{Ce}_{0.8}\text{Zr}_{0.2}\text{O}_2$ (76%), CeO_2 (36%) and ZrO_2 (24%) after 8 h of reaction at 353 K. It was proved that Ce/Zr mole ratio and morphology had a remarkable influence on the catalytic performance and selectivity of the products. A larger BET surface area, more defect sites, and strong synergetic interaction were identified as the key factors in achieving high activity and superior selectivity to styrene epoxide for the Ce–Zr mixed oxide catalyst. Further, optimization of various reaction parameters was accomplished using highly active catalyst and its recyclability was studied up to six cycles with consistent results.

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