Reprint
Epoxidation on Zeolites

Ethylene Epoxidation Catalyzed by Ag Nanoparticles on Ag-LSX Zeolites formed by Pressure- and Temperature-Induced Auto-Reduction


Abstract: Ag⁺-Exchanged LSX (Ag-LSX: Ag₉₆Al₅₆Si₃₈O₁₄₈·nH₂O), a large pore low silica analogue (Si/Al = 1.0) of faujasite, was prepared and post-synthetically modified using pressure and temperature in the presence of various pore-penetrating fluids. Using high-resolution synchrotron X-ray powder and single crystal diffraction we derive structural models of the as-prepared and post-synthetically modified Ag-LSX materials. In the as-prepared Ag-LSX model, we located 96 silver cations and 245 H₂O molecules distributed over seven and five distinctive sites, respectively. At 1.4(1) GPa pressure and 150 °C in ethanol the number of silver cations within the pores of Ag-LSX is reduced by ca. 47.4%, whereas the number of H₂O molecules is increased by ca. 40.8%. The formation of zero-valent silver nanoparticles deposited on Ag-LSX crystallites depends on the fluid present during pressurization. Ag-nanoparticle-Ag-zeolite hybrid materials are recovered after pressure release and shown to have different chemical reactivity when used as catalysts for ethylene epoxidation.

Zeolites are widely utilized in industry as ion-exchangers, sorbents, molecular sieves, and heterogeneous catalysts.[1] The members of the faujasite family display an unsurpassed performance in catalytic cracking of hydrocarbons and nitrogen/oxygen separation[2] and are classified into two sub-classes: X- and Y-types with a Si/Al ratio from 1 to 1.5 and Y-types with Si/Al ratios above 1.5. Amongst the X-type, low silica X (LSX) has a Si/Al ratio of 1 and the highest ion-exchange capacity.[3,4] In industrial applications, zeolites undergo thermal, compressional, and chemical changes, and understanding the systematic structural and compositional changes when exposed to temperature and pressure is crucial to modify and control properties as well as ensure prolonged performance as catalysts. A growing number of investigations using in situ temperature and pressure conditions have explored the cation-dependent dehydration and compression behavior of various X- and Y-type materials.[5]

Silver exchanged as a cation in zeolites leads to a number of materials with useful catalytic and adsorptive properties.[6] Recently a metallic silver-containing hybrid zeolite material has been successfully synthesized by annealing the small-pore zeolite natrolite at moderate pressure and temperature conditions at 1.7(1) GPa and 250 °C.[7] This reaction under pressure and temperature can be rationalized using the auto-reduction model initially proposed by Jacobs (1979)[8] in which the Ag⁺ cations in the hydrated zeolite reacts with water and the framework oxygen to form gaseous O₂ and zero-valent Ag atoms. Using only heat as pointed out by Heo et al.,[9] the auto-reduction leads to an amorphization of the zeolite framework which can be reversed by heating in O₂ or avoided by heating at elevated temperatures in flowing O₂. In Ag-Y zeolites, Tsutsumi et al. showed that nanocrystalline metallic silver particles are formed in Ag-Y zeolites after heat treatment above 600°C in the presence of hydrocarbons.[10] Beyer et al.[11] showed that the reduction mechanism of Ag in Y zeolites is dependent on temperature, and Gellens et al.[12] showed that cyclic reduction with hydrogen and oxidation with oxygen of Ag⁺ ions in Y type zeolites depends on both redox conditions and temperature. In silver exchanged chabazite, the metallic silver particles are deposited on the surfaces of chabazite crystallites after heating at 150 °C.[13] We have extended our initial high-pressure investigation on natrolites to the large-pore zeolite LSX to establish pressure-induced auto-reduction in a large pore zeolite. Our first goal was to structurally characterize the recovered Ag₉₆Ag-LSX hybrid materials containing mono- and zero-valent silver inside and outside of LSX crystallites, respectively. After distinct pressure and heat treatments of monovalent silver-exchanged low-silica X (Ag-LSX) and X (Ag-X) zeolites in the presence of different fluids used as pore-penetrating pressure-transmitting media, we derived structural models.
of the as-prepared Ag-LSX and pressure-recovered hybrid Ag/Ag-LSX materials. Using scanning electron microscopy (SEM) with energy-dispersive X-ray spectroscopy (EDX), and high-angle annular dark-filed scanning transmission electron microscopy (HAADF-STEM), we confirmed that auto-reduction of monovalent silver cations in as-prepared Ag-LSX results in zero-valent silver particles deposited on Ag-LSX crystallites. Silver and its oxides have been used as catalysts for ethylene epoxidation, and many computational studies probed the detailed reaction mechanisms of other oxidation processes. Our second goal was to investigate the catalytic reactivity and selectivity of Ag/Ag-LSX hybrids using pressure- and temperature-induced reduction for ethylene epoxidation. Our goal was not to find optimized new catalysts for this reaction, by exploring established and possibly new promoters and modifiers, but to confirm that both conversion rate and selectivity of this reaction can be significantly impacted by rather simple post-synthetic modification, which increases the parameter space of such an optimization. The pressures we use are accessible in presses used to make KBr pellets for IR measurements.

In-situ high-pressure synchrotron X-ray powder diffraction (HPXRD) using a diamond anvil cell (DAC) was performed to explore the auto-reduction of Ag-LSX in the presence of different fluids (water, methanol, ethanol, acetone, benzene and toluene) (Figures 1 and S3a in the Supporting Information). The pressure measurement error is ±0.1 GPa. Figure 1 shows the normalized intensity of the metallic silver (111) peak as a function of pressure in the presence of different fluids. The normalized intensity of the metallic silver (111) peak increases the most as pressure increases in the presence of methanol and ethanol. No phase transitions of the LSX material were observed in any fluid used to ensure hydrostatic pressure. The formation of nanocrystalline metallic Ag is promoted by pressure alone, in particular in the presence of methanol and ethanol. After pressure release and exposure to ambient conditions, metallic Ag (Fm3m, a = 4.08(1) Å) is still detected in the synchrotron X-ray pattern (Figure S3a) corroborating that metallic Ag nanoparticles formed without heating under pressure do not re-oxidize to monovalent silver at ambient conditions. Furthermore, the crystallinity of the zeolite framework is preserved in pressure-induced auto-reduction—this is in marked contrast to thermally-induced Ag nanoparticle formation.

To maximize Ag migration and retain the crystallinity of Ag-LSX, optimized temperature and pressure conditions (1.0(1) GPa and 150 °C) were used to synthesize Ag-LSX-P$_{150}$ using a large volume press (see the Supporting Information, Figure S3b). After a 1 hour dwelling time for Ag migration at 1.0(1) GPa and 150 °C, the samples were recovered and subjected to further characterization and experiments.

The PXRD profile of Ag-LSX-P$_{150}$ tracked the evolution of metallic Ag$_0$ nanoparticles from Ag-LSX by the growth of the (111) and (200) diffraction peaks of the cubic Ag$_0$ phase (Figure S3b in the Supporting Information). STEM images of Ag-LSX-P$_{150}$ also confirmed the formation of Ag$_0$ particles with sizes between 5 and 25 nm at the surface of Ag-LSX crystallites (see Figure 1).

Based on our Rietveld refinements the stoichiometry of Ag-LSX-P$_{150}$ is Ag$_{51(7)}$Si$_{96}$Al$_{96}$O$_{384}$$\cdot$310(43)H$_2$O, maintaining an Si/Al ordered framework with Si/Al = 1 crystallizing in a cubic unit cell (Fd3) with $a = 24.982(2)$ (Table S2 in the Supporting Information).
The metallic Ag nanoparticles crystallizes in a cubic cell (Fm3m) with a = 4.0877(2). It is possible that the higher H₂O content in this model compared to that of Ag-LSX might be attributed to pressure-induced insertion of ethanol into the super-cages. However, due to the disordered distribution of the H₂O molecules we cannot distinguish pressure-inserted ethanol molecules from those of water using our current data. Furthermore, the charge balance of the Ag-LSX-P₂O₅ sample Ag₁₅₀Si₉₆Al₉₆O₃₈₄·₃₁₀H₂O calls for 4₅ framework protons, which due to the presence of water within the pores of the zeolite form according to: 2Ag₉₆Si₉₆Al₉₆O₃₈₄ + 22.₅H₂O ! pressure ! 11.₂₅O₂ + 9₀AgO + 2Ag₁₅₀Si₉₆Al₉₆O₃₈₄. Although this reaction is charge balanced, the error on the Ag content of Ag-LSX-P₂O₅ indicates a range of possible redox reactions.

Ag⁺ cations and H₂O molecules are found in six and seven distinctive crystallographic sites, respectively (Figures S6c,d in the Supporting Information). Compared to the Ag-LSX model, Ag-LSX-P₂O₅ shows lower Ag occupancies at nearly all sites (Figure S6e). The number of silver ions at site I and I’ decreased from 15.₀(1) to 1₀.₄(3) and from 1₄.₆(2) to 7.₇(6), respectively (Tables S1 and S2 in the Supporting Information). The total number of silver ions at site II and II* also declined from 3₀(2) to 1₀(2). In the super-cage, the total number of silver ions at sites III’ decreased from 3₅(3) to 2₃(4), and the number of Ag sites also decreased from three to two.

For comparison, a single crystal of Ag⁺-exchanged zeolite X (Ag₈₈Al₈₈Si₃₄O₃₈₄·nH₂O) was prepared and characterized using X-ray diffraction. All diffraction peaks are indexed in the cubic space group Fd₃m, indicating a Si/Al disordered framework.

During the structural analysis, about 9₁ silver ions were found in the unit cell of zeolite X (Table S5 in the Supporting Information). These Ag⁺ ions were placed at five extra-framework sites with partial occupancies as follows: site I (9₁%), I’ (4₉%), II’ (4%), II⁺ (8₃%), III’ (17%). However, after applying high pressure (0.₇(1) GPa) and temperature (1₇₅°C) to the crystal under ethanol PTM, the number of Ag in zeolite X was considerably reduced from 9₁(3) to 3₅(1), while maintaining the same space group (Table S₅). Although there was little change in the occupancies of site I (9₁ to 9₀%) and II’ (4 to 9%), there were significant reductions of the occupancies in the other sites: site I’ (4₉ to 2₉%), II⁺ (8₃ to 2₆%). After applying high pressure and temperature to a crystal of zeolite X strong diffraction peaks of Ag⁺ were detected (Figure S5), which is in accordance with the results from the powder diffraction experiments on Ag-LSX (Figures 1 and 3 in the Supporting Information).

The PXRD pattern reveals that when heating the Ag-LSX-P₂O₅ sample above 1₇₅°C the amount of metallic silver gradually decreases (Figure 2a). This indicated that the Ag particles formed by pressure re-oxidize at higher temperatures. Some work has been done to reduce Ag⁺ ions in X zeolites in hydrogen flow and re-oxidize Ag atoms in an oxygen flow, but the previously formed silver particles on the surface of X zeolites remained unchanged with oxygen flow at 4₇₃ K.[¹³] Based on in situ synchrotron X-ray diffraction data, the changes of the unit cell volume of Ag-LSX-P₂O₅ as a function of temperature reveals three distinct regions (Figure S₈a in the Supporting Information): the unit cell volume initially contracts from ambient to 1₂₅°C, followed by an abrupt expansion between 1₂₅°C.
and 175 °C, and a successive contraction until 600 °C. The overall unit cell volume contraction seems to be related to the gradual loss of H₂O (or pressure-inserted ethanol) as corroborated by the TGA (Figure S8b). An increased number of Ag ions at site II led to larger T-O(2)-T and T-O(4)-T angles, which are associated with the S6R (Tables S2, S4 in the Supporting Information and Figure 2a). This caused the anomalous expansion of the unit-cell volume between 125 and 175 °C. Above 175 °C, the gradual reduction of the unit cell volume is attributed to the reduced Ag occupancy in the site II and the steady removal of H₂O molecules as evidenced in the Ag-LSX-P₁₀₀₀-T₃₀₀ 300 and Ag-LSX-P₁₀₀₀-T₄₀₀ 400 models (Table S2 and Figures 2 and 58).

To confirm the reversible redox reaction of metallic Ag and Ag-LSX, pressure- and temperature-treatments performed on the Ag-LSX-P₁₀₀₀-T hybrid material after cooling from 600 °C to room temperature of Ag-LSX-P₁₀₀₀-T in a DAC at 0.8(1) GPa and 150 °C, indicated that the metallic Ag particles are formed again (Figure S9 in the Supporting Information). Ag-LSX-P₁₀₀₀-T₃₀₀ (Ag-LSX-P₁₀₀₀-T after pressure- and heat-treatment) was then heated inside a furnace (WiseTherm FH) at 300 °C and 500 °C for 1 hour. Ag-LSX-P₁₀₀₀-T₃₀₀ (Ag-LSX-P₁₀₀₀-T with heat-treatment) showed the reduction of the intensities of the diffraction peaks of metallic Ag particles (Figure S9).

Figure 1 shows the morphological changes as seen in SEM and HAADF-STEM images and the corresponding X-ray diffraction pattern of different Ag-LSX materials using different post-synthetic pressure- and heat-treatments. The XRD pattern and SEM image shows no indications for metallic silver in the as-prepared Ag-LSX. A heat treatment up to 600 °C results in a collapsed and largely amorphous Ag-LSX phase and no formation of metallic silver nanoparticles. Ag-LSX-P₁₀₀₀ hybrid materials made by pressure- and heat-treatments at 1.4(1) GPa and 150 °C, respectively, contain silver nanoparticles observed in both SEM and HAADF-STEM images. The HAADF-STEM images and X-ray diffraction experiments reveal that a fraction of Ag⁺ remains in the pores of Ag-LSX. Additional heating of the Ag-LSX-P₁₀₀₀ sample up to 600 °C reduces the amount silver nanoparticles on the Ag-LSX crystallites as corroborated by both XRD pattern and SEM images. Pressurizing Ag-LSX up to 13.0(1) GPa in the presence of ethanol results in a highly disordered zeolite material and Ag nanoparticles. Higher pressures thus trigger a pressure-induced amorphization. Amorphization is also observed at temperatures above 300 °C.

The catalytic activity of the hybrid materials depends on (1) the post-synthetic conditions used, such as heat treatment at 600 °C, and pressure treatment in the presence of ethanol as well as (2) the specific reaction conditions studied, such as reaction temperature and pretreatment conditions in the reactor. For all catalysts, C₄H₄ conversion increased with increasing temperature and remained under 5 % at temperatures below 220 °C, as shown in Figure 1. Spline line fitting was used between experimental points on Figure 1 to guide the eye. Ag-LSX-P₁₀₀₀ exhibited the highest C₂H₄ conversion, with 23.3 ± 0.6 % conversion at 275 °C, whereas Ag-LSX showed the lowest conversion with just 3.9 ± 0.2 % at 303 °C. Conversely, Ag-LSX displayed the highest conversion among the four catalysts when the furnace set points were decreased to 150 °C. Ag-LSX-600 °C was the most selective catalyst studied, achieving 29.2 ± 4.9 % ETO selectivity at 146 °C at a 1.2 ± 0.1 % conversion rate, as shown in Figure 1. With the exception of the reaction data collected at 200 °C, Ag-LSX-600 °C consistently showed higher ETO selectivity than the standard 15 % Ag/α-Al₂O₃ catalyst. Although Ag-LSX and Ag-LSX-600 °C were found to exceed the selectivity of the 15 % Ag/α-Al₂O₃ catalyst under certain conditions, Ag-LSX-P₁₀₀₀ and Ag-LSX-P₁₀₀₀-T₄₀₀ were on average less selective than all other catalysts studied, with Ag-LSX-P₁₀₀₀-T₄₀₀ achieving a maximum ETO selectivity of just 10.9 ± 4.8 % and a 3.0 ± 0.5 % C₂H₄ conversion rate at a bed temperature of 209 °C. However, catalytic selectivity also varies with the conversion rate, and a direct comparison of selectivity is only meaningful at similar conversion rates. Furthermore, the industrial standard in ethylene oxide selectivity is close to 90%. As mentioned above, our catalyst does not contain alkali, rhenium, and no chlorine compounds were added to the feed as in industrial processes.

Catalytic performance at 250 °C was found to depend on whether the catalysts were exposed to an oxidative or reductive environment prior to testing, as shown in Figure 2. Figure 2b shows that the conversion increased significantly for all catalysts after an 18 hour reductive treatment at 250 °C with the lowest conversions were obtained when no pretreatment was used. Contrary to the results for the untreated and pre-calcined catalysts, in which Ag-LSX-P₁₀₀₀ achieved the highest conversion at 250 °C, Ag-LSX-600 °C converted the most C₂H₄ after the reductive pretreatment, with 41.0 ± 2.3 %. Ag-LSX showed 10.8 ± 4.1 % selectivity after the oxidative pretreatment, but this value dropped to 4.4 ± 1.1 % after the reductive pretreatment. Ag-LSX-600 °C followed a similar trend, reaching 9.6 ± 2.2 % conversion after calcination and decreasing to 7.7 ± 0.6 % after reduction. All other catalysts reached their maximum selectivity after a reductive pretreatment. The highest selectivity achieved throughout all pretreatment experiments at 250 °C was 12.2 ± 0.4 %, which was achieved by Ag-LSX-P₁₀₀₀-T₄₀₀ after reductive treatment.

We have shown that post-synthetic modifications of Ag-LSX using temperature and pressure in the presence of various fluids results in the formation of a wide variety of Ag₅/Ag-LSX hybrid materials. Our structural work establishes the different Ag-LSX structures present in, that is, Ag-LSX-P₁₀₀₀ and confirms that Ag⁺ ions in the zeolite migrate during the post-synthetic treatment to form Ag₅ nanoparticles on the surfaces of the zeolite crystallites. The degree of crystallinity of the Ag-LSX material and the amount of Ag nanoparticles on their surfaces can be controlled. We also note that the type of fluid plays a role in the pressure treatment. This provides an alternative route for a non-thermal in situ formation of Ag nanoparticles on zeolite crystallite surfaces. Pressure-driven in situ synthesis of metallic nanoparticles on the surface of zeolite crystallites should be explored in other systems as it provides an alternative route to thermal and/or physical deposition and is highly reproducible.

The catalytic reactivity and selectivity of various Ag₅/Ag-LSX hybrids during ethylene epoxidation (ETO) clearly establishes the importance of pressure- and temperature-induced post-
synthetic modifications. As expected, the different morphologies depicted in Figure 1 impact the chemical reactivity. At temperatures above 225 °C, the pressure-driven post-synthetic modification results in the highest C\textsubscript{2}H\textsubscript{4} conversion comparable to those of a non-optimized 15% Ag/\textalpha-Al\textsubscript{2}O\textsubscript{3} catalyst. The subsequent loss of crystallinity found in hybrid materials which were heated up to 600 °C reduces the C\textsubscript{2}H\textsubscript{4} conversion to the level of the 15% Ag/\textalpha-Al\textsubscript{2}O\textsubscript{3} catalyst and other post-synthetically modified hybrids. The observed changes in C\textsubscript{2}H\textsubscript{4} conversion and ETO selectivity make the case that taking an appropriately cation substituted zeolite and exploring post-synthetic modifications using both pressure and temperature is a promising strategy to create new heterogeneous metallic nanoparticles on zeolites as support. This should further intensify the use of post-synthetic modifications in high-throughput studies of heterogeneous catalysts. Additional in situ treatment of the catalysts reveals that a prolonged reduction over 18 hours at 250 °C results in significant changes of the conversion rate. In particular, the heat-treated Ag-LSX-600 °C increases its C\textsubscript{2}H\textsubscript{4} conversion rate, as do the pressure-treated Ag-LSX-\textit{P}_{\text{ISOV}} and pressure- and heat-treated Ag-LSX-\textit{P}_{\text{ISOV}}-600 °C while maintaining their selectivity.

We show that under hydrostatic pressure in the presence of pore-penetrating methanol and ethanol auto-reduction of Ag-LSX to Ag-nanoparticle-Ag\textsuperscript{2+}-zeolite hybrid materials occurs at moderate temperatures, while maintaining the framework crystallinity and morphology. Intriguing future possibilities are the formation of bi-metallic nanoparticles by appropriate cation substitution in the zeolites and the exploration of reactions requiring metals in zero and higher oxidation states.

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**Conflict of interest**

The authors declare no conflict of interest.