Formation of Copper Nanoparticles in LTL Nanosized Zeolite: Spectroscopic Characterization

A. Kharchenko, a V. Zholobenko, b A. Vicente, a C. Fernandez, a H. Vezin, c V. De Waele, c and S. Mintova a, *

a Laboratoire Catalyse et Spectrochimie (LCS), ENSICAEN, Université de Caen, CNRS, 14050 Caen, France
b Keele University, Keele, Staffordshire ST5 5BG, UK
c Laboratoire de Spectrochimie Infrarouge et Raman (LASIR), Université de Lille, CNRS, UMR 8516, 59000 Lille, France

Abstract: The state of copper species stabilized in nanosized LTL zeolite subjected to various post-synthesis treatments was unveiled by a range of spectroscopic techniques. FTIR and UV-Vis studies demonstrated that the reduction process of copper in the LTL nanosized zeolite leads to the formation of different species including Cu^{2+}, Cu^{+} and Cu nanoparticle (Cu NPs). The adsorption of probe molecules (NO and CO) was used to selectively monitor the copper species in the LTL nanosized zeolite upon oxidation and reduction post-synthesis treatments. Both the Cu^{2+} and Cu^{+} species were probed by NO and CO, respectively. The amount of Cu^{+} in the LTL zeolite nanocrystals was about 43% as determined by FTIR, while the amount of Cu NPs was about 55% determined by the UV-Vis spectroscopic characterization. These results were complemented by EPR, $^{29}$Si and $^{63}$Cu MAS NMR spectroscopic data. The EPR spectroscopy was further applied to monitor the effective reduction of the Cu^{2+} species and their re-oxidation, while the $^{63}$Cu MAS NMR verified the presence of Cu NPs in the LTL nanosized zeolite crystals.

Introduction

Metallic nanoparticles have attracted significant attention in catalysis due to their high activity and selectivity in many important chemical processes such as
hydrogenation/dehydrogenation, isomerization, oxidation and reduction of NO\textsubscript{x} with hydrocarbons. High catalytic activity is attributed to a high proportion of atoms located at corners and edges with low coordination numbers and have the ability to activate substrates. The number of such atoms is increasing with the decrease of their particle size and it has frequently been reported that the catalytic activity increased when the particle size decreased. Most active catalysts generally contain particles smaller than 5 nm. Substantial progress has been achieved in the synthesis of silver and gold nanomaterials with high catalytic performance. In this context, copper nanoparticles (Cu NPs) with controlled size and shape are good alternative materials for catalysis because of their electronic structure, catalytic properties and lower cost. Copper NPs that adsorb in the visible range are of particular interest due to the possibility of their application in photocatalysis and plasmonic chemistry. However, one of the main drawbacks of small metallic nanoparticles is the low thermal stability and a tendency to sinter under harsh reaction conditions, limiting their application at a large scale. These obstacles can be overcome by confining the metal NP in an inorganic matrix. Zeolites are promising hosts (matrix) for the stabilization of metal nanoparticles. The size, distribution and location of the NP are controlled by the steric constrains that originate from the zeolite topology, i.e. regular system of pores and cavities. Additionally, the zeolite structure imposes shape selectivity by limiting the diffusion of the molecules with the effective diameter bigger than the pore openings. However, the very existence of strong confinements, giving useful properties of the material, can affect their performance in catalysis and separation processes due to the slow mass transport through the micropores. Advances in the synthesis of nanosized zeolites provided materials in which the mean diffusion path is significantly lower, compared to their conventional micronsized counterparts. Indeed, the combination of superior properties on nanosized zeolites with the
exceptional reactivity of metallic NP opens a way to the design of new advanced catalytic materials.$^1$

Radiolytic and chemical approaches have been applied to control the formation of metal NPs in nanozeolites.$^9$ Nanoparticles with a mean size varying from oligomeric clusters to large nm-sized nanoparticles have been obtained depending on the type of the zeolite framework, the metal content and the reduction conditions. In the case of copper, the final oxidation state ($\text{Cu}^{II}$, $\text{Cu}^I$, $\text{Cu}^0$) is influenced by the experimental conditions and the species can coexist within the zeolite pores.$^9$

In our previous work, the formation of Cu NP in the nanosized LTL zeolite was investigated. The copper cations were introduced in zeolite nanocrystals via ion exchange after synthesis directly in the suspensions, and subsequently reduced with hydrazine monohydrate at room temperature.$^{12}$ The applied method was shown to be a promising route towards the preparation of copper NPs with desired properties due to its simplicity and reproducibility. By tailoring the synthesis conditions, it was possible to isolate supported copper nanoparticles with diverse sizes homogeneously distributed within the zeolite matrix. However, due to the nature of the synthesis process, interrupting the reduction reaction, in order to obtain the copper nanoparticles of desired size, would result in the formation of various copper species. Since the catalytic activity and selectivity can be affected by the presence of different active sites, it is important to identify both the Cu NPs and other Cu species present in the zeolite matrix. Both the targeted synthesis of copper active sites and their complementary characterization are crucial steps in the rational design of Cu NPs supported on zeolites that would allow the utilization of advantages offered by their unique properties.

FTIR of adsorbed probe molecules has been extensively employed to determine the state of supported metal species.$^{13-15}$ It is important to note that the choice of a suitable probe
molecule is crucial in the FTIR spectroscopy as it determines which surface property is tested.\textsuperscript{16} Often the use of different probe molecules in separate experiments, which provide complementary information, is a key to the comprehensive understanding of diverse species. CO and NO molecules can provide valuable information on the dispersion, coordination, and the oxidation state of supported metal species. In addition, the size of these test molecules allows site accessibility even in small zeolite cavities, whereas the experiments performed at liquid-nitrogen temperature afford the detection of species characterized by weak interaction with the probe, which cannot be detected at higher temperatures.\textsuperscript{16}

The focus of this work is on spectroscopic characterization of Cu species supported on LTL zeolite, particularly by FTIR using CO and NO probe molecules; the chemical reduction with hydrazine was described extensively in our previous report.\textsuperscript{12} The choice of CO and NO was determined by their sensitivity to Cu\textsuperscript{+} and Cu\textsuperscript{2+} species, respectively. Elucidating the nature of Cu species in zeolites is complex and usually demands a multi-technique approach. Therefore, the FTIR studies have been supplemented by EPR, \textsuperscript{29}Si and \textsuperscript{63}Cu MAS NMR and Diffuse Reflectance (DR) UV-Vis measurements. Indeed, EPR complements the information on the Cu\textsuperscript{2+} species, whereas the \textsuperscript{63}Cu and \textsuperscript{29}Si MAS NMR allow characterization of metallic copper species and changes in their environment. Additionally, DR UV-Vis is utilized to detect copper ions, metallic copper and different Cu-O complexes.

**Experimental Part**

Nanosized K-LTL type zeolite ((Si/Al=3) with monomodal particle size distribution was synthesized as described elsewhere.\textsuperscript{12} The as-prepared precursor suspensions were aged at room temperature for 24 hours prior to hydrothermal treatment (170 °C for 18 hours). After the crystallization process was accomplished, the nanosized zeolite crystals were recovered by multistep centrifugation (20000 rpm, 40 min) and washed with doubly distilled water reaching pH = 7-8 (sample is denoted as parent LTL). Zeolite samples were ion-exchanged
with copper nitrate (0.1 M) in colloidal zeolite suspensions using solutions containing 0.3 mmol Cu$^{2+}$/g zeolite to achieve ~10% of K$^+$ cations exchange for copper. The ion-exchange process was carried out at room temperature for 2 hours, and then the zeolite was separated by multi-step centrifugation to remove the excess of copper ions (sample is denoted as Cu$^{2+}$-LTL). Next, the incorporated copper cations were reduced in zeolite suspension using excess of hydrazine (Cu$^{2+}$/1000 N$_2$H$_4$) under N$_2$ atmosphere. The reaction was carried out at room temperature for 280 minutes achieving ~56% reduction of Cu$^{2+}$ to Cu$^0$ in agreement with in situ UV-Vis studies. Subsequently, the solid was isolated by centrifugation and freeze-dried and the final product (sample is denoted as Cu$^0$-LTL) was stored in the glove box in Ar atmosphere to prevent further Cu NPs oxidation. According to High Resolution Transmission Electron Microscopy (HRTEM) results, the copper nanoparticles are evenly distributed in the pores and on the surface of the LTL matrix with the particle size of 0.4-2.2 nm. A detailed description of this procedure along with the structural and textural characterization of the obtained Cu-LTL zeolites has been reported previously.

FTIR spectra of CO and NO adsorbed on zeolite samples were recorded using a Thermo-Nicolet Nexus instrument by accumulating of 128 scans at a spectral resolution of 4 cm$^{-1}$. Self-supported pellets (diameter (d) = 16 mm, weight (m) = 20 mg) were prepared from powdered samples and treated in situ in the transmission IR cells directly. Prior to the IR measurements, the samples were activated under oxygen or hydrogen atmosphere, or in vacuum at 673 – 723 K to study the effects of oxidation and reduction pretreatment on the state of copper species in order to identify different types of sites. After the pretreatment, increasing calibrated doses of CO and NO were introduced into the IR cell up to 1600 μmol/g and 2000 μmol/g, respectively. In the case of CO adsorption, the system was later evacuated at 100 K, 293 K and 423 K to monitor the stability of copper species. The deconvolution of the spectra has been performed with MagicPlot software using Gaussian function. The peak
positions have been determined using the second derivative and fixed for all spectra in order to fit the individual CO bands (Figure S 1).

The reduction of copper species was also studied by variable temperature EPR carried out at X-band (9.64 GHz) using a Brüker ELEXSYS E580 FT spectrometer. The state of copper was also characterized by diffuse reflectance UV-Vis spectroscopy utilizing a Varian Cary 4000 spectrophotometer. $^{63}$Cu and $^{29}$Si MAS NMR spectra were recorded on Bruker Avance III-HD 500 MHz (11.7 T) and 400 MHz (9.4T) spectrometers, respectively. $^{63}$Cu MAS NMR was done with a 90° pulse of 2 μs and a recycle of delay of 5 s. 26348 scans were made to accumulate the signal with a very broad spectral width. For $^{29}$Si MAS NMR experiments, single pulse excitation with pulse lengths of 7.0 μs (30° flip angle) and 60 s recycling delay were used. T1 measurements were performed using inversion recovery (π–tau-π/2) sequence. The pulse length was set to 6 (π/2) and 10.9 μs (π) and different tau values in a range from 50 μs to 20 s were acquired. Topspin software, from Bruker, was used to extract T1 values for each sample. Copper experiments were performed with 4-mm outer diameter zirconia rotors and a spinning speed of 12 kHz, while $^{29}$Si experiments were performed with a 7 mm zirconia rotor and a spinning speed of 4.5 kHz. Tetra-methylsilane (TMS) and CuCl were used as chemical shift reference for $^{29}$Si and $^{63}$Cu nucleus, respectively.

Results and discussion

FTIR spectroscopy study of copper containing zeolite samples

The FTIR spectra (OH stretching region) of the Cu$^{2+}$-LTL and Cu$^{0}$-LTL samples after activation in vacuum at 673 K are presented in Figure 1. Both materials exhibit a band at 3747–3744 cm$^{-1}$, which is characteristic of the isolated external silanol groups.$^{16}$ High relative intensity of this band reflects the small size of the crystals that has already been reported in the literature as the spectral characteristic of the nanosized zeolites with well-
developed external surface area. The Cu$^0$-LTL sample exhibits additional bands at 3661 and 3628 cm$^{-1}$, that can be assigned to extra-framework Al-sites and bridged hydroxyl $\equiv$Si(OH)Al$\equiv$ groups, respectively.$^{15-17}$ The presence of the latter can be explained by the generation of bridging OH-groups following the chemical reduction of Cu$^{2+}$ species to Cu$^0$. Appearance of extra framework aluminium and associated Al-OH groups could be explained by the partial degradation of the formed structure during high temperature activation, which is common for zeolites with low Si/Al ratios.$^{18}$ However, these effects are not significant, as the intensity of the bands at 3661 and 3628 cm$^{-1}$ is low.

Figure 1 FTIR spectra in the OH-region of Cu-LTL samples collected at 300K after activation at 673 K and 723 K: (a) Cu$^{2+}$-LTL and (b) Cu$^0$-LTL; Inset: spectra in the range 3500-3700 cm$^{-1}$.

CO adsorption on copper containing zeolite samples: FTIR spectroscopy study
The revealing the state of copper in the Cu exchanged zeolites is of primary importance for their potential applications. It is essential to control the state of the copper species (Cu\(^0\), Cu\(^+\) and Cu\(^{2+}\) ions) and the amount of desired Cu NPs, that are necessary to realize a targeted reaction and limit the number of the undesired processes. CO adsorption on metal centres monitored by FTIR spectroscopy has been widely utilized to characterize their structure and electronic properties. The frequency shifts, as compared to the vibration of the gaseous CO at 2143 cm\(^{-1}\), can give information about the oxidation state, coordination number and the metal particle size. In the case of copper ions, CO is preferentially adsorbed on Cu\(^+\) ions since the formed carbonyl complex is stabilized by both σ bonding and π-back donation.\(^{19}\) As has been shown previously,\(^{13,19,20}\) mainly electrostatic forces bond CO to Cu\(^{2+}\) ions, while the σ component of the bond is negligible, and there is no π back-donation. The carbonyls of Cu\(^{2+}\) ions are typically detected at low temperature or at high equilibrium pressure of CO (bands higher than 2160 cm\(^{-1}\)); these bands easily vanish upon increasing of the temperature. After the cooling step to 77 K, these bands typically do not re-appear due to the reduction of Cu\(^{2+}\) with CO at elevated temperature treatment.\(^{21}\) Additionally, CO forms mainly a π-bond with Cu\(^0\) atoms, and the resulting complexes are relatively unstable. CO stretching frequencies of Cu\(^+\)-CO and Cu\(^0\)-CO complexes fall in the same spectral region, 2110-2080 cm\(^{-1}\), making their discrimination difficult. Since the Cu NPs have been identified and characterized in our previous work (e.g. using TEM and UV-vis spectroscopy),\(^{12}\) the focus of this study is to characterize the residual copper species.

CO adsorption on copper containing zeolite samples has been initially studied at low temperature (100 K) by introduction of increasing calibrated doses of gas from 30 to 1600 μmol/g. The IR spectra (C-O-stretching region) of Cu-LTL samples pretreated under various conditions are shown in Figure 2, and the observed CO stretching frequencies are summarized in Table 1. Different pretreatment conditions have been used to identify the
spectral features ascribed to specific species. Careful spectral deconvolution has allowed to
differentiate them and follow their evolution with increasing CO pressure.
Adsorption of CO on the Cu$^{2+}$-LTL sample, activated under O$_2$ atmosphere, (Figure 2A)
firstly leads to the appearance of intense bands at 2140 cm$^{-1}$ and 2126 cm$^{-1}$ attributed to Cu$^+$-
CO complexes in different locations in the zeolite framework. The low intense band at 2190
cm$^{-1}$ is assigned to Cu$^{2+}$-CO species.$^{22,23}$ After the introduction of 200 μmol/g CO, two bands
at 2163 cm$^{-1}$ and 2150 cm$^{-1}$ with a shoulder at 2121 cm$^{-1}$, that rise simultaneously at higher
coverage, are also detected. These bands have been assigned to C-O-stretching of K$^+...$CO
adducts and oxygen-bonded CO.$^{13,24}$ The band 2163 cm$^{-1}$ could also be ascribed as the
interaction of CO with the silanol OH groups at low temperature.$^{13}$ The shoulder at 2177 cm$^{-1}$
indicates the interaction of CO with bridging OH groups or, which is more likely in this
system, the formation of the di-carbonyl Cu$^+$-(CO)$_2$ complexes,$^{13}$ since no OH bridging
groups are observed for the oxidized samples (Figure 1).
The frequencies of the bands in the IR spectra collected for Cu$^{2+}$-LTL sample, activated at
673 K in vacuum are similar to those for oxidized sample (Figure 2B). However, the
evolution of their intensities with the coverage level differs significantly. The bands at 2140,
2126 and 2177 cm$^{-1}$, assigned to Cu$^+$-CO and Cu$^+$-(CO)$_2$, appear first, whereas the peaks at
2190, 2163 and 2150 cm$^{-1}$ appear at higher coverage after the introduction of 200 μmol/g CO.
In both cases, a low intensity peak at 2103 cm$^{-1}$ appears at high CO coverage and is assigned
to CO adsorbed on Cu$^0$ species.$^{13}$ Figures 2A and 2B show a significant degree of oxidation
of copper to Cu$^{2+}$ in the presence of O$_2$, however, Cu$^+$ species are still preserved. The
presence of Cu$^+$-CO and Cu$^0$-CO species in the sample treated under oxygen and in vacuum
can be explained by so called “auto-reduction” of Cu$^{2+}$ at high temperature during the
dehydration of the divalent Cu$^{2+}$(H$_2$O)$_n$ complexes,$^{25,26}$ or by its reduction in the presence of
CO, that can already occur at 77 K.$^{21}$
Figure 2 Evolution of the IR spectra of copper containing LTL in the range 2300-2000 cm\(^{-1}\) upon interaction with increasing doses of CO (30-1600 µmol/g) at 100 K. Sample Cu\(^{2+}\)-LTL activated at 673 K in O\(_2\) (A), in vacuum (B), in H\(_2\) (C), and sample Cu\(^{0}\)-LTL reduced with N\(_2\)H\(_4\), activated at 723 K in vacuum (D).
Table 1 Frequencies and assignment of the C–O stretching frequencies observed in this study.

<table>
<thead>
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<th>No.</th>
<th>Species</th>
<th>Mode</th>
<th>Frequency, cm$^{-1}$</th>
<th>Notes</th>
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<td>Cu$^{2+}$-CO</td>
<td>$\nu$(CO)</td>
<td>2190</td>
<td>Unstable upon evacuation at 100 K</td>
</tr>
<tr>
<td>2</td>
<td>Cu$^+$-CO</td>
<td>$\nu$(CO)</td>
<td>2140</td>
<td>Populated first, stable upon evacuation at RT</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>2126</td>
<td>Populated first, stable upon evacuation at 423 K</td>
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<tr>
<td>3</td>
<td>Cu$^+$-(CO)$_2$</td>
<td>$\nu_s$(CO)</td>
<td>2177</td>
<td>Appear at high coverage at 100 K, not stable upon evacuation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>$\nu_{as}$(CO)</td>
<td>2150</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Cu$^{+}$-CO</td>
<td>$\nu$(CO)</td>
<td>2103</td>
<td>Unstable upon evacuation at 100 K</td>
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<tr>
<td>5</td>
<td>K$^+$-CO</td>
<td>$\nu_s$(CO)</td>
<td>2163</td>
<td>Appears at higher coverage at 100 K, unstable upon evacuation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>$\nu_{as}$(CO)</td>
<td>2150</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>2121</td>
<td>Oxygen-bonded CO</td>
</tr>
<tr>
<td>6</td>
<td>Al$^{3+}$-CO</td>
<td>$\nu$(CO)</td>
<td>2230</td>
<td>Appears at 100 K, unstable upon evacuation</td>
</tr>
<tr>
<td>7</td>
<td>OH-group</td>
<td>$\nu$(CO)</td>
<td>2163</td>
<td>Interaction with silanol group</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>2177</td>
<td>Interaction with bridging OH group</td>
</tr>
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</table>
Figures 2C and 2D present spectra of Cu$^{2+}$-LTL pretreated in H$_2$ and Cu$^0$-LTL reduced with hydrazine and activated at 723 K in vacuum, respectively. These two samples show almost identical spectra with similar distribution of band intensities and are qualitatively similar to the spectra in Figure 2A and 2B. The intensity of the band at 2140 cm$^{-1}$ is intermediate between the level of Figure 2A and 2B, and the bands at 2190 cm$^{-1}$ and 2177 cm$^{-1}$, attributed to Cu$^{2+}$-CO, Cu$^+$ di-carbonyls and possibly to bridging OH groups, exhibit a lower intensity. This is a consequence of the reduction treatments applied to these two samples and the very weak interaction of CO with the copper in (2+) oxidation state. Another important difference, when compared to Figure 2B, is the lower intensity of the bands at 2140 cm$^{-1}$ and 2126 cm$^{-1}$ (Cu$^+$-CO), and a more pronounced peak at 2103 cm$^{-1}$, assigned to Cu$^0$-CO that appears at lower pressure.

The stability of copper carbonyls was tested by progressive evacuation at 100 K (Figures S2 and Figure 3) at higher temperatures. Only the bands at 2140 cm$^{-1}$ and 2126 cm$^{-1}$, attributed to Cu$^+$-CO complexes (Figure 3), remain after evacuation at ambient temperature. After evacuation at 423 K, a prominent peak at 2126 cm$^{-1}$ still remains while the intensity of the band at 2140 cm$^{-1}$ decreases drastically.\textsuperscript{13}

The results show that the main differences between the above four samples are the intensities of the bands at 2140 cm$^{-1}$ and 2126 cm$^{-1}$, both assigned to Cu$^+$-CO interaction. Considering that the amount of copper is the same in all samples, the spectral changes can be assigned to the increased number of CO molecules coordinated to Cu NPs. This agrees with the literature data, showing that high temperature reduction of copper exchanged zeolites leads to the formation of Cu NPs. This is consistent with our previous findings demonstrating the generation of Cu NPs in Cu$^{2+}$-LTL sample reduced with hydrazine in aqueous media.\textsuperscript{12} It should be noted that all samples before and after reduction contain Cu$^+$-species that firstly interact under introducing CO.
FTIR spectroscopy of adsorbed CO can be used to estimate the concentration of Cu\(^{+}\) sites in the samples, and consequently the Cu\(^{+}\)/Cu\(_{\text{total}}\) ratio. The integrated areas of the ν(CO) band at 2140 cm\(^{-1}\) and 2126 cm\(^{-1}\) after deconvolution of the recorded spectra have been determined. Assuming the CO molar absorption coefficient for the band at 2140 cm\(^{-1}\) as 13.5 cm/μmol and for the band at 2126 cm\(^{-1}\) as 10 cm/μmol (Ref \(^{13}\)), and the surface of the pellet as 2 cm\(^2\), the concentration of Cu\(^{+}\) species is calculated by the following equation:

\[
n = \frac{\text{Area} \times S_{\text{pellet}}}{\varepsilon \times m_{\text{pellet}}},
\]

where \(n\) is the concentration of Cu\(^{+}\) (μmol/g); \(\text{Area}\) is the integrated area of the CO adsorption bands at 2140 cm\(^{-1}\) and 2126 cm\(^{-1}\) obtained after the deconvolution (cm\(^{-1}\)); \(\varepsilon\) is the...
molar absorption coefficient of CO on Cu\(^+\) (cm/\(\mu\)mol); \(S_{\text{pellet}}\) is the surface of the pellet (2 cm\(^2\)); \(m_{\text{pellet}}\) is the mass of the pellet (g).

The evolution of the integrated areas of the v(CO) bands at 2140 and 2126 cm\(^{-1}\) as a function of the introduced CO display an incremental increase followed by a plateau as shown in Figure 4A-B. Therefore, considering the Cu\(^+\) : CO stoichiometry to be 1 : 1,\(^2\) the amount of CO adsorbed on Cu\(^+\) at "saturation" can be determined from the intersection of two trend lines built from the slope and the plateau. The results reported in Table S1 confirm the high stability of Cu\(^+\) in the LTL zeolite since Cu\(^+\) is presented in all the samples in significant amounts. The quantity of Cu\(^+\) species in the Cu\(^0\)-LTL sample is almost 44% of the total copper present in the sample, which is in agreement with the UV-Vis measurements\(^1\) showing that the relative amount of Cu\(^0\) species is about 55% with respect to the total copper loading.
Figure 4. (A) Integrated area of the ν(CO) band at 2140 cm$^{-1}$ and (B) integrated area of the ν(CO) band at 2126 cm$^{-1}$ bound to copper as a function of CO concentration introduced for a set of samples: (a) Cu$^{2+}$-LTL activated at 673 K in O$_2$, (b) Cu$^{2+}$-LTL activated at 673 K in vacuum, (c) Cu$^{2+}$-LTL activated at 673 K in H$_2$ and (d) Cu$^0$-LTL activated at 723 K in vacuum.
NO adsorption on copper containing zeolite samples: FTIR spectroscopy study

The Cu$^{2+}$ cations form stronger bonds with NO than with CO. As a result, surface Cu$^{2+}$-NO species are easily formed at room temperature and detected in the IR region 1964 – 1845 cm$^{-1}$. NO bonds predominantly by a σ-bond to the cations and the higher the N-O stretching frequency is the stronger is the bond. However, NO molecule has one more electron than CO which makes π-back donation more difficult. As a result, the Cu$^{+}$-NO species are not stabilized by a σ-π synergistic effect and are considered unstable.$^{19}$ It should also be mentioned that to the best of our knowledge there is no literature data published on the FTIR spectra of the interaction of nitric oxide with Cu NPs.

The stepwise adsorption of calibrated doses of NO has been carried out at low temperature on the same series of samples as for CO adsorption. The IR spectra contain the groups of bands in the 1980 – 1850 cm$^{-1}$ and 1850 – 1650 cm$^{-1}$ regions (Figure 5A-D). The band positions and their assignment$^{14}$ are summarized in Table 2. The bands observed at 1953, 1938, 1905, 1886 cm$^{-1}$ are characteristic of Cu$^{2+}$-NO species and probably correspond to four different Cu$^{2+}$ sites. According to the literature,$^{14,28}$ the intense bands at lower frequencies can be assigned to isolated Cu$^{2+}$ ions, while the bands at higher frequencies belong to nitrosyls formed with associated Cu$^{2+}$-O-Cu$^{2+}$ sites or isolated Cu$^{2+}$ cations in the square planar coordination. As shown in Figure 5, the intensity of the bands with lower frequency (1886 cm$^{-1}$) are less affected by the sample treatment in vacuum or in H$_2$. Indeed, the isolated Cu$^{2+}$ sites are expected to be more difficult to reduce.$^{29-31}$ The second set of bands includes peaks at 1825, 1810, 1797 and 1730 cm$^{-1}$, that generally represent the interaction of NO with Cu$^{+}$-sites.$^{14}$ They can be assigned to Cu$^{+}$-NO mono-nitrosyls (peaks at 1810 and 1797 cm$^{-1}$) which have been previously observed in zeolite Y, and to Cu$^{+}$-(NO)$_2$ di-nitrosyls characterized by a doublet with $v_s$ and $v_{as}$ at 1825 and 1730 cm$^{-1}$, respectively which evolve with increasing NO pressure.$^{14,31}$
It should also be mentioned that other bands have been observed in the spectra. The peaks at 1770 cm\(^{-1}\) and 1863 cm\(^{-1}\) stay for \textit{cis}-(N\textsubscript{2}O\textsubscript{2}) weakly adsorbed in the zeolite channels, that appears due to isomerization of NO at low temperatures\textsuperscript{17,29,32}. The peaks centred at 1886 cm\(^{-1}\) and 1874 cm\(^{-1}\) become visible in the spectra of Cu\textsuperscript{0}-LTL sample, these are assigned to K\textsuperscript{+}···NO monomers (Figure 5C).\textsuperscript{17} They are masked in the case of Cu\textsuperscript{2+}-LTL pretreated in O\textsubscript{2} or under vacuum by Cu\textsuperscript{2+}-NO species (Figure 5A,B). All four samples contain the peak at 2245 cm\(^{-1}\), which has been previously described as weakly bonded N\textsubscript{2}O with the zeolite.\textsuperscript{17}

The peaks centred at 1683, 1712 and 1749 cm\(^{-1}\) can be attributed to N\textsubscript{2}O\textsubscript{4} or NO\textsubscript{2}. The band at 1749 cm\(^{-1}\) is typical of N\textsubscript{2}O\textsubscript{4} adsorbed on zeolite acidic hydroxyls, as can be seen this band is more pronounced in case of the reduced and auto-reduced samples, as the bridging OH-groups can form after the reduction of Cu\textsuperscript{2+} ions to Cu\textsuperscript{0}. The presence of various N\textsubscript{x}O\textsubscript{y} in the spectra could be explained by NO disproportion. It is known that the following equilibrium exists for NO: 3NO ↔ NO\textsubscript{2} + N\textsubscript{2}O, which is shifted to the right at higher pressures. Also, NO\textsubscript{2} can easily dimerize to form N\textsubscript{2}O\textsubscript{4}.\textsuperscript{14}

In general, the NO adsorption on Cu\textsuperscript{+} in the LTL zeolite is similar to the CO adsorption. There is a significant increase in the amount of Cu\textsuperscript{+} species for the sample prepared in vacuum compared to the one pretreated in O\textsubscript{2}. For the sample reduced under H\textsubscript{2} and Cu\textsuperscript{0}-LTL sample pretreated in vacuum, there is an evident decrease in the intensity of the bands associated with Cu\textsuperscript{+} and Cu\textsuperscript{2+} species. This can be attributed to the formation of metallic nanoparticles (Cu NPs) that do not interact with NO molecules. However, spectral lines associated to both (1+) and (2+) copper oxidation states are still present.

The adsorption of NO on the sample Cu\textsuperscript{2+}-LTL treated under oxidative conditions initially results in the interaction of NO with Cu\textsuperscript{2+} in different sites. It is followed by the appearance of a vibrational band associated with Cu\textsuperscript{+}-NO interactions at higher NO dose (Figure 5A). For the sample treated under vacuum (Figure 5B) the global intensity of the Cu\textsuperscript{2+}-NO
decreases except for the band at 1886 cm$^{-1}$ that is associated with the less accessible Cu$^{2+}$ in planar coordination. In addition, a strong increase of the contributions assigned to the Cu$^{+}$-NO species can be observed indicating the reduction of Cu$^{2+}$ to Cu$^{+}$ copper species. For the sample reduced under H$_2$ or hydrazine, the vibrational spectra contains only a weak contribution from the Cu$^{2+}$ site and two weak contributions associated with Cu$^{+}$. This demonstrates that significant amount of the Cu$^{2+}$ initially present has been reduced to Cu$^{0}$ metal species.
Figure 5 IR spectra (2400-1500 cm\(^{-1}\)) of copper containing LTL sample upon interaction with increasing doses of NO (60-2000 µmol/g) at 100 K. Sample Cu\(^{2+}\)-LTL activated at 673 K in O\(_2\) (A), in vacuum (B), in H\(_2\) (C), and sample Cu\(^{0}\)-LTL reduced with N\(_2\)H\(_4\) activated at 723 K in vacuum (D). Inserts in (C) and (D): visualization of the peaks by rescaling of the y-axis.
Table 2 Frequencies and assignment of the N–O stretching frequencies observed in this study.

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<th>Frequency, cm⁻¹</th>
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<tr>
<td>1</td>
<td>Cu²⁺-NO</td>
<td>ν(NO)</td>
<td>1955</td>
<td>Corresponds to associated Cu²⁺ sites, unstable upon evacuation at 100 K</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1938</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1905</td>
<td>Cu²⁺ sites in square pyramidal configuration</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1886</td>
<td>Cu²⁺ sites in square planar configuration</td>
</tr>
<tr>
<td>2</td>
<td>Cu⁺-NO</td>
<td>ν(NO)</td>
<td>1797</td>
<td>Converts to dinitrosyls, not stable upon evacuation</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1810</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Cu⁺-(NO)₂</td>
<td>ν₆(NO)</td>
<td>1825</td>
<td>Appear at high coverages at 100 K, not stable upon evacuation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>ν₉(NO)</td>
<td>1730</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>cis-(N₂O₂)</td>
<td>ν(NO)</td>
<td>1863</td>
<td>Weakly adsorbed in the zeolite channels, unstable upon evacuation at 100 K</td>
</tr>
<tr>
<td></td>
<td></td>
<td>ν₃(NO)</td>
<td>1770</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>N₂O</td>
<td>ν(N-N)</td>
<td>2245</td>
<td>Weakly bonded in zeolite channels, unstable upon evacuation at 100 K</td>
</tr>
<tr>
<td>6</td>
<td>N₂O₄/NO₂</td>
<td>ν₃(NO₂)</td>
<td>1683</td>
<td>Weakly bonded in zeolite channels, unstable upon evacuation at 100 K</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1712</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1749</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>K⁺-NO</td>
<td>ν(NO)</td>
<td>1886</td>
<td>Unstable upon evacuation at 100 K</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>1874</td>
<td></td>
</tr>
</tbody>
</table>
UV-Vis characterization of copper containing zeolite samples

The self-supported pellets of Cu$^{2+}$-LTL pretreated in O$_2$ and Cu$^{0}$-LTL pretreated in vacuum at same temperatures as for the FTIR experiments have been examined using diffuse reflectance UV-Vis spectroscopy (Figure 6). The Cu$^{2+}$-LTL sample contains absorption bands in the regions 240 – 450 nm and 600 – 800 nm, that are described in the literature as Cu$^{2+}$ ← O$^{2-}$ charge-transfer transitions and d – d transitions of isolated distorted octahedral Cu$^{2+}$ ions, respectively.$^{23,33}$ The presence of metallic copper species in Cu$^{0}$-LTL sample was revealed by an absorption peak at about 550 nm, which is attributed to surface plasmon (SPR) resonance of the metallic Cu.$^{33,34}$ This sample also shows band at 240 nm that is already assigned to Cu$^{2+}$ species. A broad feature between 360 and 430 nm could be assigned to O-Cu-O and Cu-O-Cu complexes.$^{35}$

![Figure 6 DR UV-Vis spectra of self-supported pellets of (a) Cu$^{0}$-LTL reduced with N$_2$H$_4$ activated at 723 K in vacuum and (b) Cu$^{2+}$-LTL activated under oxygen atmosphere at 673K.](image)
EPR characterization of copper containing zeolites

EPR spectroscopy can be used for monitoring paramagnetic Cu$^{2+}$ cations and changes in their local environment, while other copper oxidation states (Cu$^+$, Cu$^0$) are EPR silent.\textsuperscript{36} Figure 7 shows a drastic decrease in the amount of Cu$^{2+}$ species upon reduction and partial re-oxidation under calcination in air. Rasia et al. have shown the correlation between the hyperfine coupling constant A$_{\parallel}$ and g-factor (g$_{\parallel}$) in the EPR spectra of Cu(II) complexes.\textsuperscript{37} The changing values of these parameters can be used to determine the type of ligands coordinated to Cu$^{2+}$ ions at different stages. According to the correlation diagram established by Rasia et al. the starting Cu$^{2+}$-LTL sample contains copper coordinated to four oxygen atoms, while after the hydrazine treatment, the unreduced copper species are coordinated to both nitrogen from hydrazine and oxygen atoms from the zeolite framework (Table 3). These observations are in good agreement with previous reports, which describe Cu$^{2+}$ closely interacting with four framework oxygen atoms in the 6MR’s ring.\textsuperscript{38}

Table 3 The values of hyperfine coupling constant A$_{\parallel}$ and g-factor (g$_{\parallel}$) obtained from EPR spectra of copper containing zeolite samples.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Spin Hamiltonian Parameters</th>
<th>g$_{\parallel}$</th>
<th>A$_{\parallel}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu$^{2+}$-LTL</td>
<td></td>
<td>2.4</td>
<td>132</td>
</tr>
<tr>
<td>Cu$^0$-LTL</td>
<td></td>
<td>2.27</td>
<td>177</td>
</tr>
<tr>
<td>Cu$^0$-LTL-reox</td>
<td></td>
<td>2.39</td>
<td>130</td>
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</table>
29Si and 63Cu MAS NMR characterization of copper containing zeolites

Owing to the low natural abundance of 4.7% and a long spin-lattice relaxation time (T1) of 29Si nuclei, the 29Si MAS NMR measurements of porous silicates require long measuring time. While T1 varies from a few microseconds to around 30 s for most of the zeolites, it can reach 5000 seconds in layered silicate nacrite. However, it has been shown that the introduction of paramagnetic centres into mesoporous and microporous silicate crystals significantly shortens the relaxation time due to a more effective nuclear relaxation by direct interaction between the 29Si nucleus and the electron spin. Hence, the measurement of the relaxation rate changes in 29Si MAS NMR signal of copper-doped LTL zeolite should allow monitoring the degree of reduction of Cu2+ cations.

The 29Si MAS spectra of parent LTL, Cu2+-LTL and Cu0-LTL samples are shown in Figure 8. Five peaks at -88, -90, -96, -101, and -106 ppm are assigned to Si(0Si4Al), Si(1Si3Al), Si(2Si2Al), Si(3Si1Al), Si(4Si0Al), respectively. The T1 values for different sites in the
parent LTL and copper doped zeolites are summarized in Table 4. As can be seen, $^{29}$Si T1 values of different silica sites in the parent zeolite are in the range of 9-12 s, while after the introduction of 1 paramagnetic Cu$^{2+}$ ion per unit cell, the relaxation rate decreases drastically.

The analysis of the T1 values for different Si sites can be utilized to indicate the preferred coordination sites for Cu$^{2+}$ cations. The enhancement of relaxation (which corresponds to lower values of T1) follows the order Si(4Al) > Si(3Al) > Si(2Al) ≥ Si(Al) ≥ Si(0Al), suggesting that the most favourable coordination sites for Cu$^{2+}$ species are Si(4Al) and Si(3Al). For the zeolites containing reduced copper, the relaxation time has the values of the same order of magnitude as for the parent LTL sample. These results suggest a significant decrease in the amount of paramagnetic Cu$^{2+}$ species in the reduced sample.

Table 4 Relaxation time (T1) of different Q_n sites in the LTL and Cu-LTL zeolite samples.

<table>
<thead>
<tr>
<th>Sites</th>
<th>Parent LTL</th>
<th>Cu$^{2+}$-LTL</th>
<th>Cu$^{0}$-LTL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Q_4(4Al)</td>
<td>11.6</td>
<td>2.9 x10^{-2}</td>
<td>23.3</td>
</tr>
<tr>
<td>Q_4(3Al)</td>
<td>10.0</td>
<td>5.5 x10^{-2}</td>
<td>19.0</td>
</tr>
<tr>
<td>Q_4(2Al)</td>
<td>9.2</td>
<td>8.4 x10^{-2}</td>
<td>13.9</td>
</tr>
<tr>
<td>Q_4(1Al)</td>
<td>9.2</td>
<td>9.2 x10^{-2}</td>
<td>12.3</td>
</tr>
<tr>
<td>Q_4(0Al)</td>
<td>9.1</td>
<td>8.5 x10^{-2}</td>
<td>12.0</td>
</tr>
</tbody>
</table>
The reduced Cu\textsuperscript{0}-LTL sample has been also examined by \textsuperscript{63}Cu MAS NMR spectroscopy. The anisotropic resonance at 2330 ppm prevails in the \textsuperscript{63}Cu NMR spectra of the parent LTL and Cu\textsuperscript{2+}-LTL samples, which arises due to the Cu radio frequency coil in the probe (Figure 9). No other signals are observed in these materials. Two extra resonances at 2340 ppm and 360
ppm are detected in the spectrum of Cu\textsuperscript{0}-LTL sample. The peak at 2340 ppm that is resolved from the peak of the Cu coil (at 2330 ppm) has been previously assigned to metal Cu particles.\textsuperscript{42} It has been shown by FTIR that the Cu\textsuperscript{0}-LTL sample contains Cu\textsuperscript{+} species, and therefore, the peak at 360 ppm can be assigned to Cu\textsuperscript{+}. 
Conclusions

The examination of oxidation states of supported copper species in LTL nanosized zeolites utilizing a range of characterization techniques demonstrates that the reduction of copper is a complex process, involving different species such as Cu$^{2+}$, Cu$^+$ and Cu NPs in different locations. Adsorption of NO and CO allows quantitative monitoring of the modifications in the copper containing LTL zeolite upon oxidation and reduction, as both the Cu$^{2+}$ and Cu$^+$ can be selectively probed. The FTIR results indicate the heterogeneity of Cu$^{2+}$ and Cu$^+$ cations in the Cu-LTL zeolite prepared by ion-exchange procedure. Furthermore, quantitative analysis of the data obtained from the CO adsorption followed by FTIR spectroscopy shows that the relative amount of copper in (1+) state is about 43%. This agrees with the amount of copper nanoparticles (Cu$^0$) determined by UV-Vis measurement (about 55%). The reduction of Cu$^{2+}$ species and the presence of copper cations and Cu$^0$ have been also confirmed by adsorption of NO, $^{29}$Si MAS NMR and EPR. The $^{29}$Si MAS spectra showed that the most favourable coordination sites for Cu$^{2+}$ species are Si(4Al) and Si(3Al), while the $^{63}$Cu MAS NMR verified the presence of copper nanoparticles (Cu$^0$).
Acknowledgements

We acknowledge the financial support provided by TAR-G-ED ANR project.

Notes and references


Oxidation state and distribution of Cu-species in LTL zeolite upon reduction? Spectroscopy gives an answer!