

Supported Cu Nanoparticles as Selective and Stable Catalysts for the Gas Phase Hydrogenation of 1,3-Butadiene in Alkene-Rich Feeds

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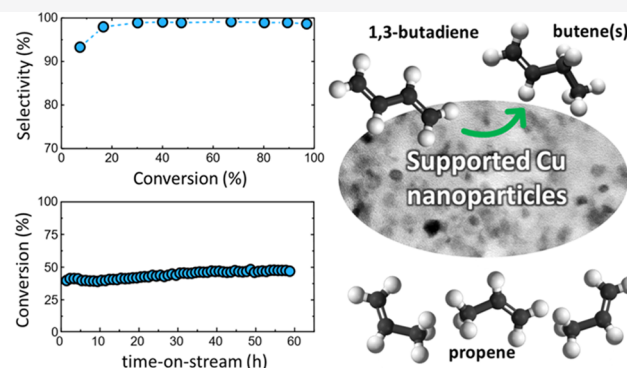
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ABSTRACT: Supported copper nanoparticles are a promising alternative to supported noble metal catalysts, in particular for the selective gas phase hydrogenation of polyunsaturated molecules. In this article, the catalytic performance of copper nanoparticles (3 and 7 nm) supported on either silica gel or graphitic carbon is discussed in the selective hydrogenation of 1,3-butadiene in the presence of a 100-fold excess of propene. We demonstrate that the routinely used temperature ramp-up method is not suitable in this case to reliably measure catalyst activity, and we present an alternative measurement method. The catalysts exhibited selectivity to butenes as high as 99% at nearly complete 1,3-butadiene conversion (95%). Kinetic analysis showed that the high selectivity can be explained by considering H₂ activation as the rate-limiting step and the occurrence of a strong adsorption of 1,3-butadiene with respect to mono-olefins on the Cu surface. The 7 nm Cu nanoparticles on SiO₂ were found to be a very stable catalyst, with almost full retention of its initial activity over 60 h of time on stream at 140 °C. This remarkable long-term stability and high selectivity toward alkenes indicate that Cu nanoparticles are a promising alternative to replace precious-metal-based catalysts in selective hydrogenation.



INTRODUCTION

Hydrogenation reactions play a crucial role in fine and bulk chemicals production. A wealth of processes is based on the addition of molecular hydrogen to an unsaturated carbon–carbon bond, often using a supported catalyst. Examples include the hydrogenation of polyunsaturated hydrocarbons to olefins,¹ hardening of vegetable oils,² and selective hydrogenation of various organic compounds such as vitamin intermediates^{3,4} as well as pharmaceutical and agrochemical active ingredients.^{3,5} The selective removal of polyunsaturated hydrocarbons from monounsaturated hydrocarbons is essential for the production of polymer- and synthesis-grade alkenes.^{1,6–8} Polyunsaturated hydrocarbons, which are often present in crude alkene streams (e.g., 1,3-butadiene, up to 1–5 wt % in C₂–C₄ steam cracking mixtures⁹), interfere with the subsequent conversion of alkenes as they can degrade polymer quality and/or poison the polymerization catalyst.^{1,10} Consequently, their concentration should be reduced to below tens of ppm.^{1,10,11} This challenge is commonly addressed by selective hydrogenation of the residual polyolefins to the corresponding mono-olefins, commonly using palladium-based

catalysts.^{12–18} Pd has a very high hydrogenation activity, which can result in a poor selectivity at high alkyne/alkadiene conversion in particular when large Pd assemblies are present. Therefore, the metal is often diluted or partially deactivated with appropriate modifiers (e.g., PdAg/Al₂O₃, PdS/CaCO₃, and PdPb/CaCO₃^{12,13,15}). However, restructuring and metal segregation (e.g., when a large excess of diluting metal is present¹⁹ or after oxidative regeneration of the catalyst²⁰) can lead to unwanted reactions such as isomerization, polymerization, and over-hydrogenation of the alkenes to alkanes which often results in limited selectivity and decreased catalyst lifetime.^{19,21}

The selective hydrogenation of alkynes and alkadienes has generated industrial interest,^{22–27} in particular regarding noble

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metals catalysis. Nonetheless, little is known about the reaction mechanism, catalyst stability, and selectivity, in particular for the case of more inexpensive and sustainable²⁸ catalysts such as those involving base metals. Copper has been reported to be an active catalyst for the selective hydrogenation of alkynes and alkadienes since the 1970s. The reactants investigated were mainly propyne,²⁹ 1,3-butadiene,^{30,31} 1-butyne,⁸ and 1-butyne-3-yne.⁸ The catalysts were prepared by reduction of precipitated Cu species⁸ and Cu²⁺ ion adsorption on silica.^{32,33} Compared to palladium-based catalysts, copper was found to be less active (3–4 orders of magnitude in terms of 1,3-butadiene turnover frequency^{5,34}), but particularly selective, showing almost full conversion to the corresponding alkenes at relatively mild conditions (temperatures below 150 °C and near-atmospheric pressure^{8,29,30}). Unfortunately, the stability of these catalysts was generally poor because of oligomer formation, often termed “green oil formation”, which limited further interest in this type of catalyst.^{6,29,35}

Renewed interest in Cu as a selective hydrogenation catalyst was spurred by a paper from Studt et al.,³⁶ who computationally predicted that the best performing metals in the selective hydrogenation of ethyne to ethene share a particularly high adsorption affinity toward the alkyne rather than the alkene. Not only Pd alloys but also base metals and related alloys such as NiZn, FeZn, and monometallic Ni and Cu were predicted to be good catalysts.³⁶ Recent experimental developments on Cu-based catalysts were reported by the group of Louis et al.³⁵ In particular, the authors showed that the addition of Zn species to TiO₂-supported Cu catalysts helped to stabilize the catalytic performance. The monometallic copper lost more than 90% activity within 4 h on stream, while for the Cu₂Zn₁ sample the decrease in activity was limited to ~30% during 20 h on stream. Partial alloying of Zn with Cu and passivation of TiO₂ surface Lewis acid sites by Zn species might play an important role in stabilizing the samples.³⁵ Further insights regarding the support effect on catalyst stability were recently reported by Masoud et al. for Au-based catalysts.³⁷ SiO₂-supported Au nanoparticles demonstrated much more stable catalytic performance than TiO₂-supported ones (10% loss of the initial activity for Au/SiO₂ after 5 days on stream at 200 °C, while Au/TiO₂ loses 90% of its initial activity after 15 h on stream, same conditions). The steady loss of activity in the case of Au/TiO₂ was due to green oil formation on the surface of the catalyst (around 2 wt % of carbonaceous deposit, as measured thermogravimetric analysis), and full recovery of the initial catalyst activity was obtained by heating in air. The same amount of carbon deposits was found on bare TiO₂ upon exposure to the reaction mixture (16 h, 200 °C),³⁷ while for SiO₂ and Au/SiO₂ the production of those species was negligible. This observation suggests that the formation of oligomers over the TiO₂ surface takes place regardless of the type of metal nanoparticles supported on it.³⁷

On the basis of these earlier findings, we decided to explore silica- and graphitic carbon-supported copper catalysts for selective hydrogenation. The supports were chosen based on their relative inertness toward oligomer formation (fouling) under reaction conditions. The samples were prepared by means of incipient wetness impregnation.^{38,39} Catalysts were tested in the selective hydrogenation of 1,3-butadiene in an alkene-rich feedstock (1,3-butadiene to propene molar ratio equal to 1 to 100),^{37,40} and we investigated in detail the catalyst selectivity and stability. Attention was also paid to the catalytic testing methodology, which was shown to be of

paramount importance to obtain reliable results for copper-based hydrogenation catalysts.

■ EXPERIMENTAL SECTION

Details on the chemicals used for the synthesis and gas feeds are reported in the [Supporting Information](#) (section 1).

Functionalization of the Carbon Support. The pristine carbon support (HSAG-500, high surface area turbostratic graphitic carbon, average pore size 250 Å, BET surface area 500 m² g⁻¹; kindly provided by Timcal Ltd.) was surface functionalized by liquid-phase oxidation using concentrated nitric acid.⁴¹ Herein, ~10 g of the pristine carbon material was suspended in HNO₃(aq) (65%, 40 mL g⁻¹) inside a 1 L round-bottom flask fitted with a reflux condenser. The pristine graphite material was heated to 80 °C in ~25 min and kept at 80 °C for 110 min. Thereafter, the reaction was quenched by diluting the suspensions with ~1.6 L of cold deionized water (25 °C). The surface-oxidized carbon was allowed to sediment for 30 min, and the supernatants were decanted. The solids were washed with deionized water until a pH of around 7 was reached. After the final washing step, the carbons were dried overnight at 120 °C and crushed into powders. Finally, the carbon support was dried for 90 min at 170 °C, under dynamic vacuum, to remove traces of adsorbed water and kept inside an Ar-filled glovebox (Mbraun LABmaster; <1 ppm of H₂O; <1 ppm of O₂).

Synthesis of Cu/SiO₂. Cu nanoparticles on silica gel were prepared via adaptation of the impregnation and drying method reported elsewhere for SiO₂-based supports.^{39,42} In particular, the impregnation solution concentration and the thermal treatment conditions were adjusted to obtain particles of ~3 and ~7 nm (final Cu weight loading equal to 5.7 wt %). Around 2 g of bare support (Davisil Grade 645, ≥99% purity, average pore size 150 Å, BET surface area 300 m² g⁻¹; Sigma-Aldrich) was dried at 230 °C for 2 h in a double-neck round-bottom flask. The material was then impregnated under static vacuum with dropwise addition of a 1 M Cu(NO₃)₂ solution acidified to pH ~ 1 with HNO₃ to incipient wetness (the volume of the solution used was equal to 90% of the total pore volume of the support). The impregnated solid was then transferred in a 4A zeolite-filled desiccator and kept there for 2 days. Subsequently, it was further dried under dynamic vacuum at room temperature (RT) for 5 days and then transferred to a tubular plug flow reactor for the final thermal decomposition treatment. Nanosized Cu⁰ particles of 3 nm (sample 3nm_Cu/SiO₂) were obtained by heating the solid to 250 °C under 20% H₂/N₂. The sample was left to cool to RT, purged with 20 vol % O₂/N₂ flow (100 mL min⁻¹ g⁻¹), and heated to 250 °C in the same gas mixture (heating ramp of 2 °C min⁻¹, isothermal hold of 2 h, 100 mL min⁻¹ gas flow for each gram of dry material). It was found that particles cannot be grown larger on silica gel by simply increasing the reduction or the nitrate decomposition temperature (see [Figure S1](#)). Larger crystallites were thus prepared by using 2% NO/N₂ as the gaseous atmosphere during the copper nitrate decomposition step, as previously described by Munnik et al.³⁹ In particular, particles of around 7 nm (sample 7nm_Cu/SiO₂) were obtained by performing the thermal treatment at 350 °C under 2% NO/N₂ (heating ramp of 2 °C min⁻¹, isothermal hold of 2 h, 300 mL min⁻¹ gas flow for each gram of dry material). The samples (supported CuO) were exposed to air and stored in closed vials at RT.

Synthesis of Cu/C. Cu/C catalysts were prepared with a similar impregnation and drying method.^{27,41} As for Cu/SiO₂, the synthesis parameters were chosen to obtain particles of ~3 and ~7 nm and guarantee homogeneous distribution of the Cu NPs over the surface of the support. For the preparation, around 2 g of dry oxidized carbon was impregnated by dropwise addition of a 2 M Cu(NO₃)₂ solution acidified to pH ~ 1 with HNO₃ to incipient wetness (the volume of the solution used was equal to 90% of the total pore volume of the support; resulting Cu weight loading equal to 6.3 wt %). The dried impregnate was heated to 230 °C (0.5 °C min⁻¹), followed by 1 h isothermal hold at 230 °C, under N₂ flow (100 mL min⁻¹ g⁻¹) to decompose the nitrate precursor. The sample was left to cool to RT, purged with 20 vol % O₂/N₂ flow (100 mL min⁻¹ g⁻¹) for 3 h at RT, and then purged with pure N₂ flow (100 mL min⁻¹ g⁻¹) for 30 min. Subsequently, the material was reduced by heating to 150 °C (2 °C min⁻¹), with 2 h isothermal hold at 150 °C, under 5 vol % H₂/N₂ flow (100 mL min⁻¹ g⁻¹). Next, the temperature was increased to either 250 or 400 °C (2 °C min⁻¹), with 1 h isothermal hold at the final temperature, to obtain the 3nm_Cu/C and 7nm_Cu/C catalysts, respectively. The final catalyst was collected after letting it cool to RT and passivating it by exposure to air (overnight) at RT and then stored in closed vials at RT. Prior to X-ray powder diffraction analysis and TEM imaging, the nanoparticles were fully oxidized under 20 vol % O₂/N₂ flow (100 mL min⁻¹ g⁻¹) at 250 °C (heating ramp of 2 °C min⁻¹, isothermal hold of 2 h).

X-ray Powder Diffraction (XRD) and TEM Imaging.

Characterization measurements were performed on XRD analysis was performed on the as-prepared samples with a Bruker D2 Phaser. Radiation source: Co K α (1.78897 Å). The diffractogram were recorded in the 2 θ interval 15°–90° with a step size of 0.05°. Rietveld refinement was performed with DIFFRAC.SUITE TOPAS software. Transmission electron micrographs of the Cu/C and Cu/SiO₂ catalysts were obtained on a Tecnai F20 apparatus, operated at 200 kV. Scanning transmission electron microscopy was performed in high-angle annular dark-field (HAADF) mode on a Talos F200X (FEI), equipped with a high-brightness field emission gun (X-FEG) and operated at 200 kV. Prior to TEM imaging, the two silica-supported samples were both ultramicrotomed to increase the contrast between the amorphous silica gel support and the Cu nanoparticles. The catalyst grains (<75 μ m) were embedded in a two-component epoxy resin (Epofix, EMS) and cured at 60 °C for 24 h. The embedded catalysts were sliced into 50 nm nominal thickness sections by means of a Diatome Ultra 35° diamond knife mounted on a Reichert-Jung Ultracut E microtome. The sections were then deposited on a TEM grid and analyzed. Graphitic carbon supported materials were finely grinded via mortar and pestle and dry deposited onto the TEM grids. Average particle sizes are defined as $d_{\text{TEM}} = \sum_i (d_{\text{NP},i}^3 / d_{\text{NP},i}^2)$, while particle size dispersion was described as the standard deviation. The number of surface copper atoms was calculated by using a copper dispersion of 1.46×10^{19} atoms per square meter of Cu exposed surface area. We calculated the latter by assuming a spherical particle shape and using the equation $S_{\text{Cu}} = 6000 / d_{\text{TEM}} \rho_{\text{Cu}}$, where S_{Cu} is the specific copper surface area (m_{Cu}² g_{Cu}⁻¹), d_{TEM} is the mean particle diameter, and ρ_{Cu} is the copper density (8.92 g cm⁻³).³⁹ The CuO and Cu particle sizes (after reduction of the precatalyst prior to catalytic tests) were assumed to be the same.

Catalytic Tests. The catalytic tests were performed with the use of a tailor-made gas-phase hydrogenation setup previously described by Masoud et al.³⁷ The experiments were performed by loading a homogeneous mixture of each individual sample (sieve fraction 75–150 μ m; total amount of copper loaded in the reactor equal to 1.28 mg) and 150 mg of SiC as thermal diluent (sieve fraction 212–425 μ m) in a U-shaped Pyrex packed bed microreactor (internal diameter of 4 mm). Prior to the test, the catalysts were reduced *in situ* under pure H₂ flow (50 mL min⁻¹) from RT to 250 °C (ramp 2 °C min⁻¹) and kept at 250 °C for 90 min. Hereafter, the catalyst was put into contact with the reaction mixture (0.3% butadiene, 30% propene, 20% hydrogen, and helium for balance with a flow rate of 50 N mL min⁻¹, 1 bar). The reactor temperature was either linearly increased from RT to 150 °C at 0.5 °C/min or held constant for 2 h long isothermal steps in the range 110–180 °C. More details on the methodology can be found later in the text (see sections [Activity Measured during a Temperature Ramp](#) and [Activity under Steady State Conditions](#)). The concentration of the products and unconverted reactants was monitored every 15 min via gas chromatography by using a flame ionization detector (GC-FID; hydrocarbons detected: C₁–C₄). The composition of the gas mixture fed to the reactor was monitored at the beginning and at the end of each catalytic run by bypassing the reactor bed. Calculated turnover frequencies (TOF, s⁻¹) were defined as the molecules of 1,3-butadiene consumed per unit time per Cu surface atom (see section 4 of the [Supporting Information](#)).

RESULTS AND DISCUSSION

Structural Characterization of Cu/C and Cu/SiO₂. The structural properties of the prepared catalysts (oxidized form) were investigated by X-ray powder diffraction, and results are reported in [Figure 1](#). CuO crystallite sizes obtained via Rietveld refinement are shown in [Table 1](#). Both silica-supported samples (3nm_Cu/SiO₂ and 7nm_Cu/SiO₂) display two main characteristic CuO reflections peaks at 42° and 45.5° (corresponding to the [002] and [111] diffractions,

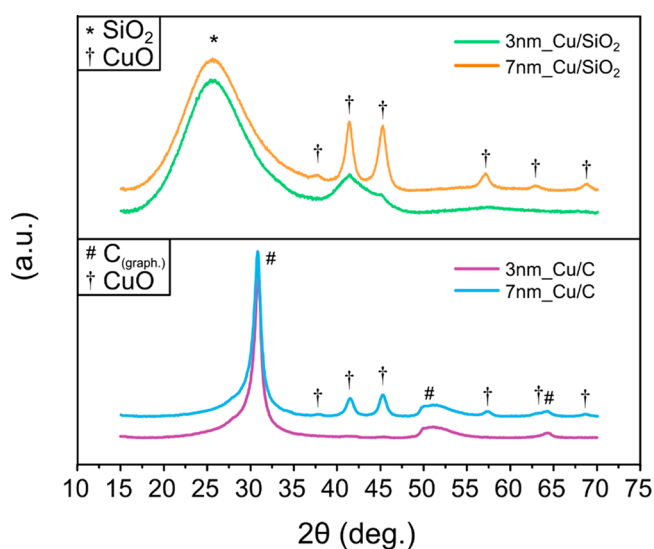


Figure 1. X-ray powder diffraction patterns of the oxidized 3nm_Cu/SiO₂, 7nm_Cu/SiO₂, 3nm_Cu/C, and 7nm_Cu/C. X-ray source: Co K α .

Table 1. Structural Properties for the As-Prepared Samples

catalysts	Cu ⁰ weight loading (%)	particle size		Cu dispersion (%)
		d_{TEM} (nm)	d_{XRD} (nm)	
3nm_Cu/SiO ₂	5.7	3.5 ± 1.0	3.1	20
7nm_Cu/SiO ₂	5.7	7.3 ± 2.4	10.4	14
3nm_Cu/C	6.3	3.1 ± 0.9	n.a.	34
7nm_Cu/C	6.3	7.3 ± 1.8	10.1	14

respectively), while the broad scattering band at 25° is ascribed to the disordered silica gel support. The peak broadening for 3nm_Cu/SiO₂ indicates the presence only small CuO crystallites in this sample. For 7nm_Cu/SiO₂ the intense and much sharper diffraction peaks demonstrate the presence of large CuO crystallites. The average CuO crystallite sizes were found to be 3.1 and 10.4 nm in 3nm_Cu/SiO₂ and 7nm_Cu/SiO₂, respectively (Table 1). The diffractogram of the 3nm_Cu/C did not show any CuO or Cu₂O reflections, implying a high copper dispersion. Lastly, the sample 7nm_Cu/C exhibited intense CuO peaks, and the average crystallite size was found to be 10.1 nm, similar to the 7nm_Cu/SiO₂ sample.

The XRD results show that the crystallite sizes on carbon can be tuned by changing the temperature at which the catalyst is treated. A reduction step performed at 400 °C led to larger and more crystalline CuO nanoparticles (sample: 7nm_Cu/C), while at 250 °C particle growth was limited (sample: 3nm_Cu/C). This was not observed in the case of CuO over SiO₂ (see Figure S1). The difference between carbon and silica supports suggests that Cu (or CuO) particles are much more strongly anchored on the surface of the silica gel rather than on the modified graphitic carbon. Moreover, reduction under H₂ at 250 °C results in visible CuO particles on SiO₂ but not on oxidized carbon by XRD, which may indicate a different interaction of the copper species with these two supports, such

as the presence of a nonhomogeneous CuO phase over the carbon support.

The CuO nanoparticle sizes and distributions on both silica and surface-oxidized graphitic carbon were investigated with TEM (Figure 2). A uniform distribution of the CuO particles over the surface of the support was observed for both the silica and the graphitic carbon. Interestingly, the 3nm_Cu/C sample contained small particles all over the surface of the carbon, which had been not detected via XRD probably due to low crystallinity. Unimodal particle size distributions were found for all the four samples, and no particles above 20 nm in diameter were present.

The particle diameters as determined by TEM are in good agreement with the crystallite sizes as determined by XRD (Table 1). In the case of 7nm_Cu/C and 7nm_Cu/SiO₂, XRD data analysis gives a slightly larger crystallite diameter (about 10 nm) in both cases. This could be explained by considering the higher degree of crystallinity and hence diffraction intensity of the larger particles present in those samples (most probably caused by slight overimpregnation of a fraction of the support), which leads to a stronger contribution to the overall signal by the larger crystallites. Hence, the effective copper dispersion used for the evaluation of the catalytic data was based on the CuO particle sizes determined from TEM data (Table 1).

Activity Measured during a Temperature Ramp.

Figure 3A depicts the 1,3-butadiene conversion as a function of the temperature measured while linearly heating the *in situ* prereduced samples from 50 to 175 °C (0.5 °C min⁻¹). All Cu-containing catalysts showed a similar conversion–temperature plot, with an onset temperature around 105 °C, followed by a sharp increase up to full hydrogenation ($T_{98\%} = 120\text{--}130$ °C). The copper catalysts were almost fully selective toward butenes, and the main product was 1-butene (selectivity ~70%, see Figure S2). Smaller amounts of *cis*-2-butene and *trans*-2-butene were also detected (10–15%), and their concentration increased along with the temperature. The selectivity to butane and propene conversion were generally

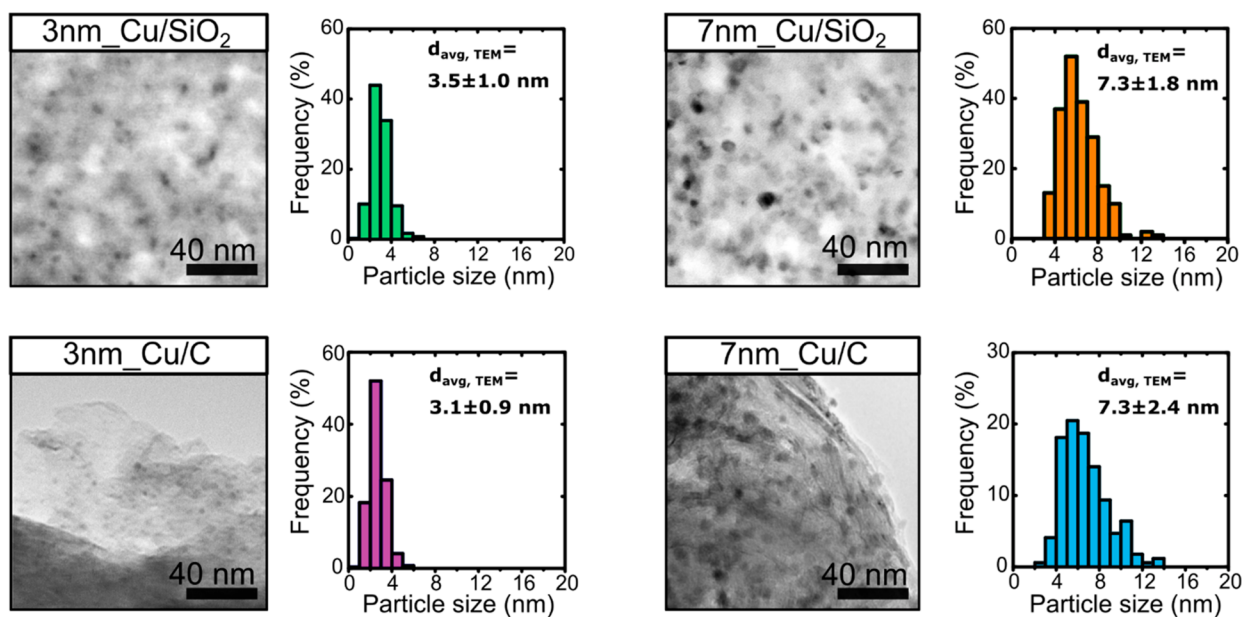


Figure 2. Representative transmission electron micrographs for the as synthesized 3nm_Cu/C, 3nm_Cu/SiO₂, 7nm_Cu/C, and 7nm_Cu/SiO₂ catalysts, with the corresponding particle size distributions. Silica-supported samples were ultramicrotomed to 50 nm slices (nominal thickness) prior to imaging.

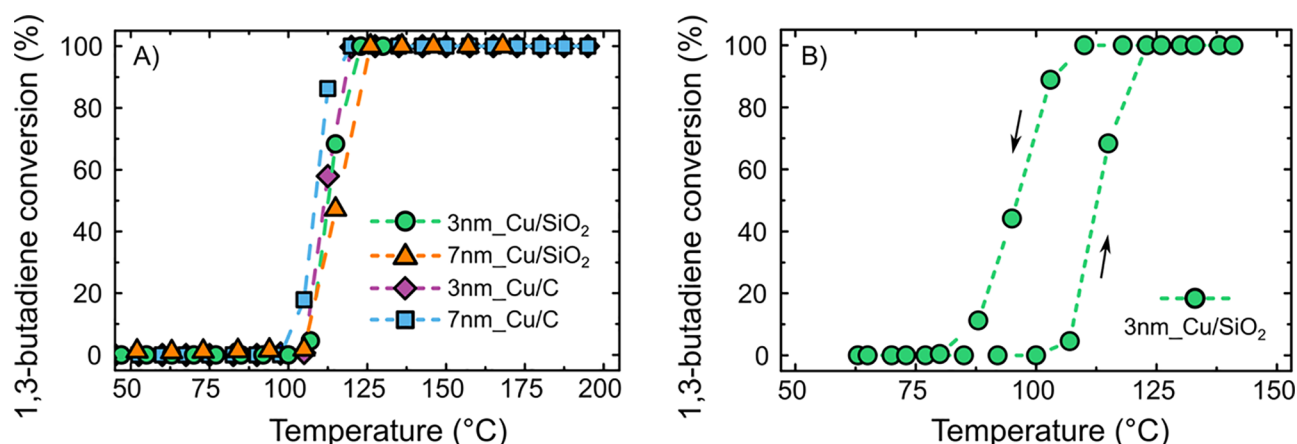


Figure 3. (A) 1,3-Butadiene conversion as a function of temperature for the 3nm_Cu/SiO₂, 7nm_Cu/SiO₂, 3nm_Cu/C, and 7nm_Cu/C samples. The data points were collected while heating the prereduced catalyst from 50 to 175 °C (heating ramp 0.5 °C min⁻¹). (B) 1,3-Butadiene conversion as a function of the temperature for the sample 3nm_Cu/SiO₂. The data points were collected while heating the *in situ* prereduced catalyst from 50 to 175 °C and then cooling the used catalyst back to RT (temperature ramp: 0.5 °C min⁻¹). Reaction conditions: butadiene/propene/H₂/He = 0.15/15/10/24.85 mL min⁻¹, 1 bar absolute pressure and 1.28 mg of Cu loaded. GHSV: 35000 h⁻¹.

below 1.5% and 1%, respectively, at almost full (>95%) 1,3-butadiene conversion. The bare supports did not show any hydrogenation activity when exposed to the reactant mixture at temperatures up to 200 °C.

All samples seem to have the same activity (per unit Cu mass) within error at temperatures between 105 and 125 °C. This is surprising as differences due to the different particles sizes and supports might be expected. Hence, we started a more detailed study of the activity by temperature cycling experiments, with as an example results for 3nm_Cu/SiO₂ reported in Figure 3B. A pronounced hysteresis was observed; the high 1,3-butadiene conversion was retained to temperatures below 105 °C during cooling. A possible explanation might be the formation of local hot spots in the catalyst bed due to the exothermicity of the hydrogenation reaction, although this is not expected under the given conditions (see section 2 of the Supporting Information). Another possibility is that the low activity between 85 and 105 °C during the first heating ramp is due to deactivation of the surface of the copper nanoparticles by oxidation. Oxidation of copper is favored at low temperatures (e.g., during cooling steps after *in situ* activation or at the beginning of the catalytic test heating ramp) and can be caused by trace amounts of oxygen and/or water that are always present in commercial gas mixtures. The onset of activity might correspond to the reduction of Cu oxides by the H₂ present in the reaction mixture. This is supported by the fact that indeed the onset of hydrogenation activity shown in Figure 3 is very close to the onset of reduction of Cu oxides (see Figure S3). Such deactivation by formation of surface oxides, which are reduced at a certain temperature, would also explain the similar activity onset temperatures observed for all catalysts (Figure 3A).

Interestingly, a sudden increase in conversion between 100 and 120 °C was previously reported also in the case of hydrogenation of various alkynes using Cu-based catalysts.⁸ In fact, using a single heating ramp is a common way to assess the activity of catalysts. Although this might be a valid method when measuring catalytic performance of more noble metals such as Pd,⁴³ Pt,⁴⁴ or Au,^{37,45} clearly for Cu-based catalysts the intrinsic properties might be obscured by oxidation. Therefore, another type of testing protocol had to be developed, as described in the next section.

Activity under Steady-State Conditions. The new measurements protocol involved a first step of catalyst preconditioning for 15 h under reaction mixture at $T = 110\text{--}130$ °C. Subsequently, the catalyst was heated (or cooled) to the desired temperature, and kinetic data were recorded for 2 h in isothermal conditions. The data points used for performance evaluation were collected at the end of a 2 h hold to reach a steady state in terms of solid-to-gas interface composition and thermal equilibrium. The process was iterated at each temperature level, and multiple points were collected at one chosen temperature (e.g., 120 °C for the sample 3nm_Cu/SiO₂, as reported in Figure S4) to detect any deactivation or activation phenomena of the catalyst during the entire duration of the test. As an example, the 1,3-butadiene conversions as a function of the temperature obtained with the described protocol are given in Figure 4A,B for three selected catalysts: 3nm_Cu/SiO₂, 7nm_Cu/SiO₂, and 7nm_Cu/C. With this new measurement protocol, the activity plot looked very different from the one obtained by linearly heating the catalyst (Figure 3): the new protocol yielded a well-defined onset temperature of activity, no hysteresis, and clear differences between the various catalysts. Both 7nm_Cu/C and 7nm_Cu/SiO₂ showed very similar conversion profiles (Figure 4A). Hydrogenation activity was measurable starting at around 115 °C with almost full 1,3-butadiene conversion reached at around 170 °C. However, the sample 3nm_Cu/SiO₂ (Figure 4B) displayed a higher activity than the 7nm_Cu/SiO₂, with full hydrogenation at 135 °C.

The turnover frequencies (normalized to the number of exposed copper atoms) measured at 125 °C were $(6 \pm 2) \times 10^{-3}$, $(5 \pm 2) \times 10^{-3}$, and $(22 \pm 6) \times 10^{-3} \text{ s}^{-1}$ respectively for 7nm_Cu/C, 7nm_Cu/SiO₂, and 3nm_Cu/SiO₂. The TOFs measured for Cu nanoparticles in this work can be compared for those previously reported in the literature. Wang et al. measured a turnover frequency of $32 \times 10^{-3} \text{ s}^{-1}$ at 105 °C for 5 nm Cu on TiO₂ (same reaction conditions),⁴⁶ comparable but appreciably higher activity than the catalyst hereby reported. The authors, in fact, stated that the catalyst was able to partially hydrogenate 1,3-butadiene at temperatures as low as 60 °C.⁴⁶ The lower Cu-based activity for the catalysts supported on SiO₂ and carbon, compared to TiO₂, may be caused by the presence of support effects for small nano-

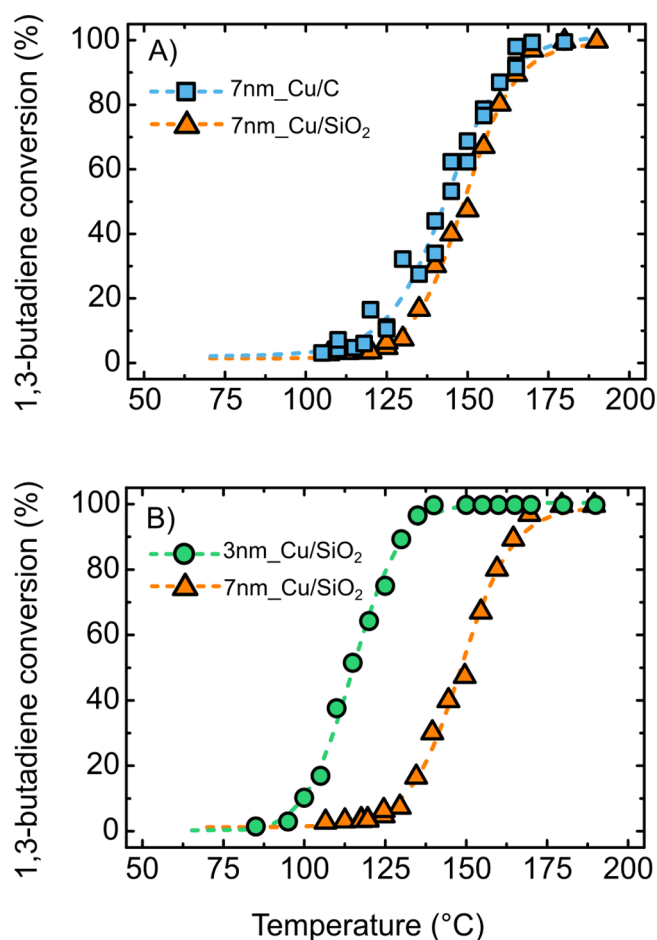


Figure 4. 1,3-Butadiene conversion as a function of temperature for (A) 7nm_Cu/SiO₂ and 7nm_Cu/C and (B) 3nm_Cu/SiO₂ and 7nm_Cu/SiO₂ samples. The catalysts were preconditioned for at least 15 h at 110–130 °C prior to measurements (see Figure S4). The catalytic data were collected after allowing at least 2 h at a given *T* for reaching stable state conditions. Conditions: 1,3-butadiene/propene/H₂/He = 0.15/15/10/24.85 mL min⁻¹, 1 bar absolute pressure and 1.28 mg of Cu loaded for each test. GHSV: 35000 h⁻¹.

particles on reducible metal oxide supports. For example, Masoud et al. reported a 4-fold higher activity for <5 nm Au nanoparticles supported on TiO₂ compared to Au/SiO₂.³⁷ The activity may also be strongly influenced by the specific catalyst pretreatment and reaction conditions. The Cu-based activities for 1,3-butadiene hydrogenation investigated in our work were in every case higher than for Au/SiO₂ catalysts (4×10^{-3} s⁻¹ at 120 °C)³⁷ as well as for Ag/SiO₂ systems (0.2×10^{-3} s⁻¹ at 120 °C)⁴⁰ yet much lower than for Pd/Al₂O₃ (10.5 s⁻¹ at 40 °C) as measured under similar reaction conditions.³⁴ Hence, our results confirm that Cu catalysts are reasonably active catalysts for 1,3-butadiene hydrogenation. However, at least as important as the activity is the catalyst stability, which is discussed in the next section.

Stability. Measurements performed at steady-state conditions revealed that the catalysts, after proper preconditioning, can display stable and reproducible activity (see Figure 5). However, TEM analysis of the used catalysts (see Figure S5) revealed that a carbon supported catalyst, 7nm_Cu/C, suffered from particle growth (from ~7 to ~23 nm) while the silica-supported samples retained their initial particle size. Hence, we focused in more detail on the Cu/SiO₂ system (see Figure S6

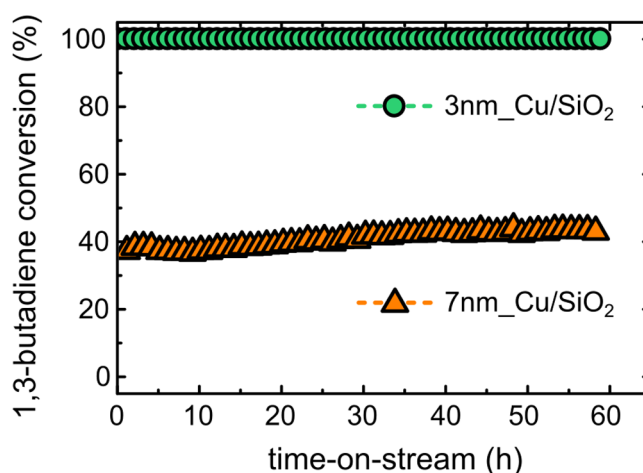


Figure 5. 1,3-Butadiene conversion as a function of the time-on-stream for the 3nm_Cu/SiO₂ (test performed at 130 °C) and 7nm_Cu/SiO₂ (140 °C) samples. Conditions: 1,3-butadiene/propene/H₂/He = 0.15/15/10/24.85 mL min⁻¹, 1 bar absolute pressure and 1.28 mg of Cu loaded for each test. GHSV: 35000 h⁻¹.

for more details on the carbon supported catalysts) and more specifically on the activated (reduced) 3nm_Cu/SiO₂ and 7nm_Cu/SiO₂ catalysts.

The stability of these two selected catalysts was investigated by means of isothermal runs. Selectivity to butenes and butenes composition are reported in Figure S7. The 3nm_Cu/SiO₂ sample tested at 130 °C exhibited full conversion for the entire 60 h of catalytic test. At full conversion, it was not possible to observe small changes in activity. Long-term tests on the sample 7nm_Cu/SiO₂ (140 °C, otherwise same conditions) revealed a high stability, with the 1,3-butadiene conversion equal to ~40% from the beginning to the end of the run. It is interesting to note a good agreement between these data and the one collected at steady-state conditions (Figure 4) for both the samples, which underlines the remarkable stability of these catalysts (in particular for the 7 nm Cu on SiO₂). Lastly, *ex situ* FT-IR investigations of the fresh and used samples revealed no significant oligomers/green oil formation (see Figure S8).

The catalysts described in this work were significantly more stable than Cu catalysts immobilized on other oxidic supports reported in the literature. A direct comparison between our systems and 5 nm Cu/TiO₂ described by Wang et al. tested using the same reaction mixture, yet *T* = 90 °C, shows that the 5 nm Cu/TiO₂ catalyst lost all more than 90% activity after <10 h on stream.³⁵ Deactivation was mainly ascribed to significant “green oil” or coke poisoning (formation of C₄₊ oligomers or carbonaceous deposit).^{6,29,35} No results for Cu catalysts on SiO₂ have been reported previously; however, differences in stability between SiO₂- and TiO₂- supported catalysts were reported by our group for Au-based systems used in selective hydrogenation for the same reaction mixture.³⁷ Higher stability was observed for Au nanoparticles immobilized on the silica support than on titania P25 (almost full retention of the activity for Au/SiO₂ catalyst after 16 h at 200 °C, versus 88% loss in activity for Au/TiO₂ tested in the same conditions). Also in this case, used catalyst characterization revealed important coke formation for the sample Au/TiO₂.³⁷ An important role of the support was reported for the hydrogenation of alkynes as well. Sárkány,⁴⁷ for instance, observed that the amount of carbon deposited over Pd catalyst

supported on TiO_2 was twice the one on $\alpha\text{-Al}_2\text{O}_3$ during pulse-flow hydrogenation experiment of ethyne/ethene mixtures (3.9 carbon atoms per Pd surface atom for Pd/ TiO_2 versus 1.66 for Pd/ Al_2O_3). Combination of adjacent vinyl species and/or insertion of ethyne into Pd–C bond of adsorbed vinyl intermediates to form a diene were considered to be the initiator of the formation of green oil in ethyne hydrogenation. The authors postulated that the TiO_2 support, due to the presence of Ti^{3+} sites decorating the Pd particle, enhanced the concentration of retained hydrocarbons over the surface of the catalyst, leading to a greater formation of C_{6+} oligomers.⁴⁷ It is clear that using more inert supports, as we show here for Cu-based catalysts, is a successful strategy to produce stable catalysts for the hydrogenation of alkadienes.

Selectivity. In this section, the selectivities of 3nm_Cu/ SiO_2 and 7nm_Cu/ SiO_2 are discussed in more detail, based on the data from Figure 4. Selectivities are reported in terms of both the evolution of the propene conversion (Figure 6A) and

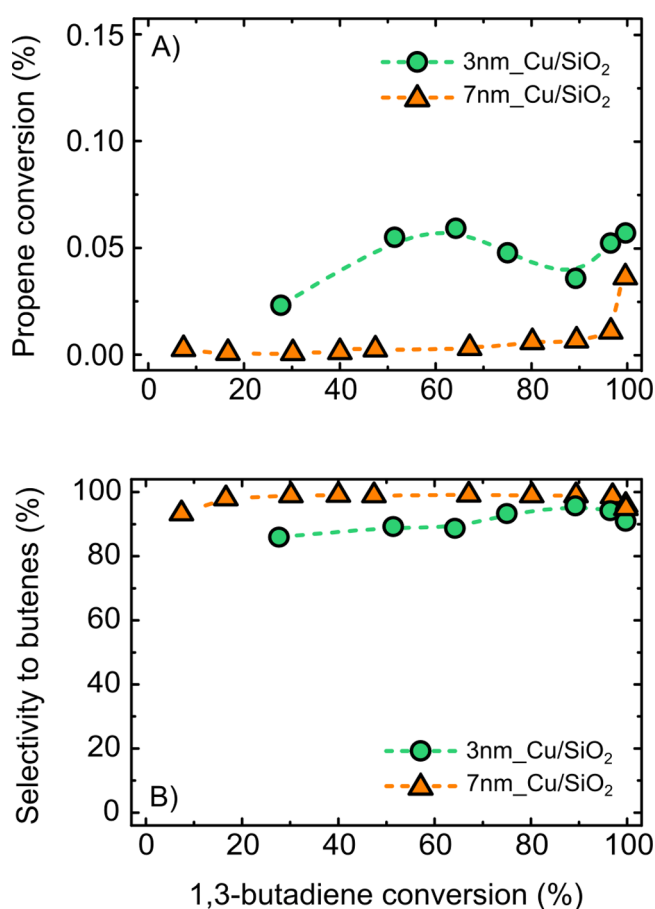


Figure 6. (A) Propene conversion and (B) selectivity to butenes data (defined as the ratio between the butenes productivity over the total hydrogen consumption) as a function of the 1,3-butadiene conversion for the reduced 3nm_Cu/ SiO_2 and 7nm_Cu/ SiO_2 . Pretreatment, reaction conditions, and testing methodology are as in Figure 4.

the selectivity to butenes (defined as the ratio between the C_4 alkenes productivity over the total hydrogen consumption; see section 4 of the Supporting Information) (Figure 6B) as a function of the 1,3-butadiene conversion. Hydrogenation of alkenes to alkanes was limited, with a propene conversion below 0.1% in the entire 1,3-butadiene conversion range (Figure 6A). Sample 3nm_Cu/ SiO_2 exhibited a maximal

alkene conversion of 0.06%, while for 7nm_Cu/ SiO_2 the hydrogenation of propene was equal to 0.013% at almost full (97%) 1,3-butadiene conversion (corresponding to roughly 40 ppm of propane present in the reactor outlet stream).

Although the propene conversion is commonly used in the literature to assess the performance of this class of hydrogenation catalysts, it is valuable to additionally look at the total selectivity to butenes (Figure 6B). In this way both undesired reactions, hydrogenation of propene and over-hydrogenation of 1,3-butadiene, can be taken into consideration. Both silica supported catalysts presented high selectivity. For instance, at 97% 1,3-butadiene conversion, the selectivities to butenes were 94% for the 3 nm Cu and an astonishing 99% to butenes for 7nm_Cu/ SiO_2 . It is interesting to appreciate the substantial difference in selectivity between the two catalysts. The higher overall activity of 3nm_Cu/ SiO_2 in terms of 1,3-butadiene hydrogenation to butenes (Figure 4B) and alkenes hydrogenation with respect to 7nm_Cu/ SiO_2 (Figure 6) suggests a possible particle size effect. This aspect will be covered in further studies.

To better understand the catalyst selectivity, sample 3nm_Cu/ SiO_2 was also tested at temperatures well above 140 °C, the T required to achieve full hydrogenation of 1,3-butadiene. Under these conditions (with top-down flow) at the bottom of the catalytic bed, the 1,3-butadiene concentration is negligible, and the reaction atmosphere consists mainly of propene, butenes, and H_2 . The fraction of the catalytic bed that experiences these conditions increases with the reaction temperature. The 1,3-butadiene conversion, propene conversion, and C_4 composition are reported in Figure 7. The

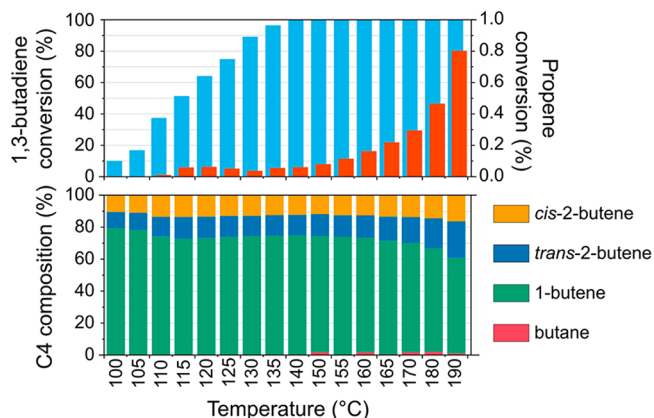


Figure 7. 1,3-Butadiene conversion (blue bars), propene conversion (orange bars), and C_4 (butenes) distribution as a function of temperature for 3nm_Cu/ SiO_2 . Pretreatment, reaction conditions, and testing methodology are as in Figure 4.

propene conversion is low and constant from 110 °C up to 150 °C (temperatures at which the 1,3-butadiene consumption is below 98%). At temperatures higher than 140 °C, i.e., at full conversion, when the concentration of 1,3-butadiene is depleted in the bottom of the catalytic bed, the propene conversion increases (even though remains lower than 1% at 50 °C higher than the temperature required for full 1,3-butadiene hydrogenation).

A similar trend was observed for the C_4 products, with the *trans*-2-butene concentration increasing with temperature only above 150 °C, at the expense of the less stable 1-butene, a sign that isomerization becomes significant only when 100%

conversion is reached and when part of the catalytic bed is depleted in 1,3-butadiene (see Figure S9 for butenes thermodynamic data). Concomitantly, *n*-butane appears as a product of the hydrogenation of C_4 species from 150 °C onward. This clearly suggests that competitive adsorption, hence the strong adsorption of 1,3-butadiene on Cu, is a key factor in the catalyst high selectivity for alkadiene conversion, as side reactions such as isomerization or further hydrogenation of alkenes only take place once the 1,3-butadiene concentration is severely depleted in the reactor and hence also low in concentration at the surface of the Cu particles. This can be compared to what was observed for the palladium-based catalysts for which the preferential adsorption of alkynes and dienes makes it possible to maintain high alkenes selectivity up to relatively high conversion, although the hydrogenation rate of adsorbed ethene is for instance about 100 times higher than that of adsorbed ethyne.^{48,49}

Further evidence is provided by analyzing the reaction kinetics data. From the study of the 1,3-butadiene hydrogenation rate as a function of the partial pressure of the reactive species (details in Figure S10), we observed a first-order dependency in H_2 pressure, a fractional positive order in 1,3-butadiene (0.25) and a nearly zeroth order in propene (0.08). The first-order dependency in hydrogen pressure suggests a low hydrogen surface coverage (linear adsorption regime),^{1,31} in line with the general observation that H_2 adsorbs relatively weakly on Cu.^{1,30} The only slightly positive order in 1,3-butadiene can be ascribed to its particularly strong affinity to bind the copper nanoparticles, since almost complete surface saturation is reached. The low reaction order in 1,3-butadiene is in line with literature reports for experiments involving Ni, Au, Pd, and Co.^{1,18,31,50} In addition, this kinetic behavior partially resembles what was observed in the case of Pd model catalysts (Pd/ Al_2O_3 /NiAl(110), Pd > 4 nm),¹⁸ for which the 1,3-butadiene adsorption can be so strong that the hydrocarbon can even limit hydrogen penetrating the adsorption layer and adsorbing onto Pd active sites.¹⁸ Lastly, the near-zeroth reaction order in propene partial pressure shows that the competition of adsorption between the butadiene and propene is strongly in favor of the former.

We hence explain the almost full selectivity to butenes of our Cu/SiO₂ system (Figure 6) to the strong preferential bonding of polyunsaturated molecules over monounsaturated ones. As long as there is an appreciable 1,3-butadiene concentration, the diene almost fully covers the surface of the Cu catalyst, hindering the hydrogenation and/or isomerization of alkenes (Figure 7). Interestingly, by assuming similar intrinsic hydrogenation rates for all the adsorbed hydrocarbons, the selectivity measured for 7nm_Cu/SiO₂ at 130 °C would correspond to a 1,3-butadiene to propene surface coverage ratio of 50:1. Conversion of butenes (mainly 1-butene) was also very limited, which indicates that they (mainly 1-butene), once formed, then rapidly desorb from the surface of the copper nanoparticles. An adsorption/desorption-driven selectivity was already predicted computationally in the case of hydrogenation of alkynes/alkenes mixtures, such as ethyne/ethene,³⁶ propyne/propene over Cu and Ni,⁵¹ and the hydrogenation of enynes over Cu.⁵² In a recent study of the selective hydrogenation of 1-phenyl-1-propyne in a batch reactor, the authors found that Cu/SiO₂ has no intrinsic selectivity toward alkenes formation at full conversion since the apparent rate of hydrogenation of the alkene to the alkane is faster than that of hydrogenation of the alkyne to the alkane.⁵³

They rather assigned the high selectivity to alkenes to a competitive substrate adsorption on the Cu nanoparticles with the adsorption constant of the alkyne on Cu/SiO₂ being orders of magnitude higher than that of the alkene.

Lastly, a comparison with the literature reveals that the propene conversion values here obtained (Figure 6A) are drastically lower than that for other Cu-based catalysts in the literature (0.1–1%), as measured under similar reaction conditions (Cu/TiO₂,⁴⁶ Cu–Zn/TiO₂,³⁵ Cu–Au/TiO₂, and Au/TiO₂⁶). Our Cu/SiO₂ systems also outperform Pd-based NPs (generally >1% propene conversion, same conditions),⁵⁴ single-atom Pd and Pt/Cu (~0.1 and ~1% propene conversion; corresponding gas feed: 1.9/2% butadiene, 70/20% propene, 4.7/16% hydrogen and balance He, respectively),^{55,56} Au/SiO₂ catalysts (for which the conversion of propene was already particularly low, <0.1%, same conditions),^{37,40} or PdAu bimetallic nanorods (selectivity to butenes below 90%, same conditions).⁵⁴ To the best of our knowledge the highest butenes selectivity (>98%) under similar conditions was reported for Cu-based catalysts obtained via carbonization of MOFs (HKUST-1, 50 wt % Cu).⁵⁷

CONCLUSIONS

Copper nanoparticles (~3 and ~7 nm) supported either on silica gel or on surface-oxidized graphitic carbon were synthesized and used as catalysts for the selective hydrogenation of 1,3-butadiene in the presence of a 100-fold excess of propene. A new protocol for the measurement of catalytic data for Cu-based hydrogenation catalysts was developed, which is a more reliable alternative to the ramp-up methodology. The catalysts exhibited full conversion of the alkadiene at mild temperatures (130–170 °C). The conversion of propene at almost full 1,3-butadiene consumption was generally less than 0.2% (0.01% for the 7 nm Cu on SiO₂ system), and the selectivity to butenes was above 94% for all the catalysts (up to 99% for the same catalyst 7nm_Cu/SiO₂). Reaction order and detailed product analysis proved that the exceptionally high selectivity can be ascribed to the strong preferential adsorption on the Cu surface of alkadienes rather than alkenes. Furthermore, the 7 nm Cu/SiO₂ retained its initial activity for the full 60 h time on stream test at 140 °C.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.jpcc.0c08077>.

Materials, numerical check for mass and heat transport limitations, C_4 products distribution measured during temperature ramp experiments, H_2 -TPR of the prepared samples, catalyst stability during steady-state experiments, transmission electron micrographs of fresh and used catalysts, stability measurements for Cu/C samples, selectivity to butenes during stability tests, *ex situ* FT-IR collected for fresh and used 3 nm Cu on SiO₂, C_4 alkenes equilibrium composition, reaction rate order analysis, TOF, and selectivity to butenes calculations (PDF)

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Author Contributions

G.T. and R.B. contributed equally to this work. G.T., R.B., P.E.d.J., and L.D. designed the experiments. R.B. synthesized and characterized the functionalized carbon support and the C-supported CuO nanoparticles. G.T. synthesized and characterized the SiO₂-supported CuO nanoparticles. G.T. and R.B. performed the catalytic experiments and analyzed the data. G.T. and P.E.d.J. wrote the paper. P.E.d.J. conceived and supervised the project. All authors discussed the results and edited the manuscript.

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Notes

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REFERENCES

- (1) Bond, G. C. *Metal-Catalysed Reactions of Hydrocarbons*; Springer: New York, 2005; Vol. 1.
- (2) Veldsink, J. W.; Bouma, M. J.; Schöön, N. H.; Beenackers, A. A. C. M. Heterogeneous Hydrogenation of Vegetable Oils: A Literature Review. *Catal. Rev.: Sci. Eng.* **1997**, *39*, 253–318.
- (3) Chen, B.; Dingerdisen, U.; Krauter, J. G. E.; Lansink Rotgerink, H. G. J.; Möbus, K.; Ostgard, D. J.; Panster, P.; Riermeier, T. H.; Seebald, S.; Tacke, T.; et al. New Developments in Hydrogenation Catalysis Particularly in Synthesis of Fine and Intermediate Chemicals. *Appl. Catal., A* **2005**, *280*, 17–46.
- (4) Bonrath, W.; Medlock, J.; Schutz, J.; Wustenberg, B.; Netscher, T. In *Hydrogenation*; Karamé, I., Ed.; IntechOpen: Rijeka, Croatia, 2012; Vol. 1, pp 69–90.
- (5) Blaser, H.-U.; Malan, C.; Pugin, B.; Spindler, F.; Steiner, H.; Studer, M. Selective Hydrogenation for Fine Chemicals: Recent

Trends and New Developments. *Adv. Synth. Catal.* **2003**, *345*, 103–151.

(6) Delannoy, L.; Thrimurthulu, G.; Reddy, P. S.; Méthivier, C.; Nelayah, J.; Reddy, B. M.; Ricolleau, C.; Louis, C. Selective Hydrogenation of Butadiene over TiO₂ Supported Copper, Gold and Gold-Copper Catalysts Prepared by Deposition-Precipitation. *Phys. Chem. Chem. Phys.* **2014**, *16*, 26514–26527.

(7) Nikolaev, S. A.; Zhanavskiy, L. N.; Smirnov, V. V.; Averyanov, V. A.; Zhanavskiy, K. L. Catalytic Hydrogenation of Alkyne and Alkadiene Impurities from Alkenes. Practical and Theoretical Aspects. *Russ. Chem. Rev.* **2009**, *78*, 231–247.

(8) Koeppel, R. A.; Wehrli, J. T.; Wainwright, M. S.; Trimma, D. L.; Cant, N. W. Selective Hydrogenation of C₄-Alkynes over a Copper on Silica Catalyst. *Appl. Catal., A* **1994**, *120*, 163–177.

(9) Ren, T.; Patel, M.; Blok, K. Olefins from Conventional and Heavy Feedstocks: Energy Use in Steam Cracking and Alternative Processes. *Energy* **2006**, *31*, 425–451.

(10) Červený, L. *Catalytic Hydrogenation*; Elsevier: Amsterdam, 1986.

(11) Bender, M. An Overview of Industrial Processes for the Production of Olefins - C₄ Hydrocarbons. *ChemBioEng. Rev.* **2014**, *1*, 136–147.

(12) Che, C.; Liang, Y.; Qian, Y.; Han, W.; Zhang, F.; Gou, G.; Jing, X.; Chang, X.; Gui, Q.; Gu, L.; et al. Palladium-Based Supported Hydrogenation Catalyst, and Preparation Method and Application Thereof. U.S. Patent US 2018/0290949 A1, 2018.

(13) Borodziński, A.; Bond, G. C. Selective Hydrogenation of Ethyne in Ethene-Rich Streams on Palladium Catalysts, Part 2: Steady-State Kinetics and Effects of Palladium Particle Size, Carbon Monoxide, and Promoters. *Catal. Rev.: Sci. Eng.* **2008**, *50*, 379–469.

(14) Chinayon, S.; Mekasuwandumrong, O.; Praserttham, P.; Panpranot, J. Selective Hydrogenation of Acetylene over Pd Catalysts Supported on Nanocrystalline α -Al₂O₃ and Zn-Modified α -Al₂O₃. *Catal. Commun.* **2008**, *9*, 2297–2302.

(15) Ravanchi, M. T.; Sahebdehfar, S.; Fard, M. R.; Fadaerayeni, S.; Bigdeli, P. Pd-Ag/ α -Al₂O₃ Catalyst Deactivation in Acetylene Selective Hydrogenation Process. *Chem. Eng. Technol.* **2016**, *39*, 301–310.

(16) Piccolo, L.; Piednoir, A.; Bertolini, J. C. Pd-Au Single-Crystal Surfaces: Segregation Properties and Catalytic Activity in the Selective Hydrogenation of 1,3-Butadiene. *Surf. Sci.* **2005**, *592*, 169–181.

(17) Piccolo, L.; Valcarcel, A.; Bausach, M.; Thomazeau, C.; Uzio, D.; Berhault, G. Tuning the Shape of Nanoparticles to Control Their Catalytic Properties: Selective Hydrogenation of 1,3-Butadiene on Pd/Al₂O₃. *Phys. Chem. Chem. Phys.* **2008**, *10*, 5504–5506.

(18) Silvestre-Albero, J.; Rupprechter, G.; Freund, H. J. Atmospheric Pressure Studies of Selective 1,3-Butadiene Hydrogenation on Well-Defined Pd/Al₂O₃/NiAl(110) Model Catalysts: Effect of Pd Particle Size. *J. Catal.* **2006**, *240*, 58–65.

(19) Vignola, E.; Steinmann, S. N.; Vandegehuchte, B. D.; Curulla, D.; Sautet, P. C₂H₂-Induced Surface Restructuring of Pd-Ag Catalysts: Insights from Theoretical Modeling. *J. Phys. Chem. C* **2016**, *120*, 26320–26327.

(20) Jin, Y.; Datye, A. K.; Rightor, E.; Gulotty, R.; Waterman, W.; Smith, M.; Holbrook, M.; Maj, J.; Blackson, J. The Influence of Catalyst Restructuring on the Selective Hydrogenation of Acetylene to Ethylene. *J. Catal.* **2001**, *203*, 292–306.

(21) Furlong, B. K.; Hightower, J. W.; Chan, T. Y.-L.; Sarkany, A.; Gucci, L. 1,3-Butadiene Selective Hydrogenation Over Pd/Alumina and CuPd/Alumina Catalysts. *Appl. Catal., A* **1994**, *117*, 41–51.

(22) Hill, T.; Haake, M.; Schwab, E.; Frenzel, A.; Worz, H. Selective Catalytic Gas-Phase Hydrogenation of Alkynes, Dienes, Alkenynes and/or Polyenes. U.S. Patent US 2003/0069458 A1, 2003.

(23) Arganbright, R. P. Selective Hydrogenation of Dienes and Acetylenes in C₃ Streams. WIPO Patent WO 94/04477, 1994.

(24) Putman, H. M.; Adams, J. R. Process and Catalyst for Selective Hydrogenation of Dienes and Acetylenes. U.S. Patent US 8,227,650 B2, 2012.

- (25) Sugeta, M.; Fukada, H. Catalyst for Selective Hydrogenation of Acetylene Compounds in 1,3- Butadiene, Method for Producing the Same and Method of Using the Same. E.P. Patent EP 2 329 879 A1, 2011.
- (26) Ryu, Y. J. Selective Hydrogenation Process and Catalyst. WIPO Patent WO 2006/044005 A2, 2006.
- (27) Beerthuis, R.; Sunley, J. G.; De Jong, K.; De Jongh, P. Selective Hydrogenation of Polyunsaturates. WIPO Patent WO2019/233961, 2019.
- (28) Nuss, P.; Eckelman, M. J. Life Cycle Assessment of Metals: A Scientific Synthesis. *PLoS One* **2014**, *9*, No. e101298.
- (29) Wehrli, J. T.; Thomas, D. J.; Wainwright, M. S.; Trimm, D. L.; Cant, N. W. Selective Hydrogenation of Propyne over Supported Copper Catalysts: Influence of Support. *Appl. Catal.* **1991**, *70*, 253–262.
- (30) Nishimura, E.; Inoue, Y.; Yasumori, I. The Mechanism of the Selective Hydrogenation of 1,3-Butadiene on Copper Surfaces. *Bull. Chem. Soc. Jpn.* **1975**, *48*, 803–807.
- (31) Phillipson, J. J.; Wells, P. B.; Wilson, G. R. The Hydrogenation of Alkadienes. Part III. The Hydrogenation of Buta-1,3-Diene Catalysed by Iron, Cobalt, Nickel, and Copper. *J. Chem. Soc. A* **1969**, 1351–1363.
- (32) Ossipoff, N. J.; Cant, N. W. The Hydrogenation and Oligomerization of Propyne over an Ion-Exchanged Copper on Silica Catalyst. *J. Catal.* **1994**, *148*, 125–133.
- (33) Wehrli, J. T.; Thomas, D. J.; Wainwright, M. S.; Trimm, D. L.; Cant, N. W. Selective Hydrogenation of Propyne over an Ion-Exchanged Copper on Silica Catalyst. *Appl. Catal.* **1990**, *66*, 199–208.
- (34) Garcia Cervantes, G.; Cadete Santos Aires, F. J.; Bertolini, J. C. Compared Properties of Pd on Thermo-Conductor Supports (SiC , Si_3N_4) and Pd on Oxide Supports (Al_2O_3 , SiO_2) for the 1,3-Butadiene Hydrogenation Reaction. *J. Catal.* **2003**, *214*, 26–32.
- (35) Wang, Z.; Wang, G.; Louis, C.; Delannoy, L. Novel Non-Noble Bimetallic Cu-Zn/TiO₂ Catalysts for Selective Hydrogenation of Butadiene. *J. Catal.* **2017**, *347*, 185–196.
- (36) Studt, F.; Abild-Pedersen, F.; Bligaard, T.; Sørensen, R. Z.; Christensen, C. H.; Nørskov, J. K. Identification of Non-Precious Metal Alloy Catalysts for Selective Hydrogenation of Acetylene. *Science* **2008**, *320*, 1320–1322.
- (37) Masoud, N.; Delannoy, L.; Schaink, H.; van der Eerden, A.; de Rijk, J. W.; Silva, T. A. G.; Banerjee, D.; Meeldijk, J. D.; de Jong, K. P.; Louis, C.; et al. Superior Stability of Au/SiO₂ Compared to Au/TiO₂ Catalysts for the Selective Hydrogenation of Butadiene. *ACS Catal.* **2017**, *7*, 5594–5603.
- (38) Munnik, P.; de Jongh, P. E.; de Jong, K. P. Recent Developments in the Synthesis of Supported Catalysts. *Chem. Rev.* **2015**, *115*, 6687–6718.
- (39) Munnik, P.; Wolters, M.; Gabrielson, A.; Pollington, S. D.; Headdock, G.; Bitter, J. H.; de Jongh, P. E.; de Jong, K. P. Copper Nitrate Redispersion to Arrive at Highly Active Silica-Supported Copper Catalysts. *J. Phys. Chem. C* **2011**, *115*, 14698–14706.
- (40) Masoud, N.; Delannoy, L.; Calers, C.; Gallet, J. J.; Bournel, F.; de Jong, K. P.; Louis, C.; de Jongh, P. E. Silica-Supported Au–Ag Catalysts for the Selective Hydrogenation of Butadiene. *ChemCatChem* **2017**, *9*, 2418–2425.
- (41) Beerthuis, R.; de Rijk, J. W.; Deeley, J. M. S.; Sunley, G. J.; de Jong, K. P.; de Jongh, P. E. Particle Size Effects in Copper-Catalyzed Hydrogenation of Ethyl Acetate. *J. Catal.* **2020**, *388*, 30–37.
- (42) Wang, G.; van den Berg, R.; de Mello Donega, C.; de Jong, K. P.; de Jongh, P. E. Silica-Supported Cu₂O Nanoparticles with Tunable Size for Sustainable Hydrogen Generation. *Appl. Catal., B* **2016**, *192*, 199–207.
- (43) Briot, P.; Primet, M. Catalytic Oxidation of Methane over Palladium Supported on Alumina. Effect of Aging under Reactants. *Appl. Catal.* **1991**, *68*, 301–314.
- (44) Yadav, O. P.; Palmqvist, A.; Cruise, N.; Holmberg, K. Synthesis of Platinum Nanoparticles in Microemulsions and Their Catalytic Activity for the Oxidation of Carbon Monoxide. *Colloids Surf., A* **2003**, *221*, 131–134.
- (45) Hugon, A.; Delannoy, L.; Louis, C. Supported Gold Catalysts for Selective Hydrogenation of 1,3-Butadiene in the Presence of an Excess of Alkenes. *Gold Bull.* **2008**, *41*, 127–138.
- (46) Wang, Z.; Brouri, D.; Casale, S.; Delannoy, L.; Louis, C. Exploration of the Preparation of Cu/TiO₂ Catalysts by Deposition-Precipitation with Urea for Selective Hydrogenation of Unsaturated Hydrocarbons. *J. Catal.* **2016**, *340*, 95–106.
- (47) Sárkány, A. Formation of C₄ Oligomers in Hydrogenation of Acetylene over Pd/Al₂O₃ Catalysts. *React. Kinet. Catal. Lett.* **2001**, *74*, 299–307.
- (48) Bond, G. C.; Wells, P. B. The Mechanism of the Hydrogenation of Unsaturated Hydrocarbons on Transition Metal Catalysts. *Adv. Catal.* **1965**, *15*, 91–226.
- (49) Silvestre-Albero, J.; Rupprechter, G.; Freund, H. J. From Pd Nanoparticles to Single Crystals: 1,3-Butadiene Hydrogenation on Well-Defined Model Catalysts. *Chem. Commun.* **2006**, *1*, 80–82.
- (50) Hugon, A.; Delannoy, L.; Louis, C. Influence of the Reactant Concentration in Selective Hydrogenation of 1,3-Butadiene over Supported Gold Catalysts under Alkene Rich Conditions: A Consideration of Reaction Mechanism. *Gold Bull.* **2009**, *42*, 310–320.
- (51) Bridier, B.; López, N.; Pérez-Ramírez, J. Partial Hydrogenation of Propyne over Copper-Based Catalysts and Comparison with Nickel-Based Analogues. *J. Catal.* **2010**, *269*, 80–92.
- (52) Bridier, B.; Karhánek, D.; Pérez-Ramírez, J.; López, N. Molecular Understanding of Enyne Hydrogenation over Palladium and Copper Catalysts. *ChemCatChem* **2012**, *4*, 1420–1427.
- (53) Kaeffer, N.; Larmier, K.; Fedorov, A.; Copéret, C. Origin of Ligand-Driven Selectivity in Alkyne Semihydrogenation over Silica-Supported Copper Nanoparticles. *J. Catal.* **2018**, *364*, 437–445.
- (54) van der Hoeven, J. E. S.; Jelic, J.; Olthof, L. A.; Totarella, G.; Studt, F.; van Blaaderen, A.; de Jongh, P. E. Unlocking Synergy in Bimetallic Catalysts by Core-Shell Design. *ChemRxiv* DOI: 10.26434/chemrxiv.13218155.v1 (accessed November 17, 2020).
- (55) Yan, H.; Cheng, H.; Yi, H.; Lin, Y.; Yao, T.; Wang, C.; Li, J.; Wei, S.; Lu, J. Single-Atom Pd₁/Graphene Catalyst Achieved by Atomic Layer Deposition: Remarkable Performance in Selective Hydrogenation of 1,3-Butadiene. *J. Am. Chem. Soc.* **2015**, *137*, 10484–10487.
- (56) Lucci, F. R.; Liu, J.; Marcinkowski, M. D.; Yang, M.; Allard, L. F.; Flytzani-Stephanopoulos, M.; Sykes, E. C. H. Selective Hydrogenation of 1,3-Butadiene on Platinum-Copper Alloys at the Single-Atom Limit. *Nat. Commun.* **2015**, *6*, 1–8.
- (57) Hu, N.; Li, X.-Y.; Liu, S.-M.; Wang, Z.; He, X.-K.; Hou, Y.-X.; Wang, Y.-X.; Deng, Z.; Chen, L.-H.; Su, B.-L. Enhanced Stability of Highly-Dispersed Copper Catalyst Supported by Hierarchically Porous Carbon for Long Term Selective Hydrogenation. *Chin. J. Catal.* **2020**, *41*, 1081–1090.

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