Highly modulated supported triazolium-based ionic liquids: direct control of the electronic environment on Cu nanoparticles†

Cristián Valdebenito, a Jose Pinto, b Michael Nazarkovsky, c Gustavo Chacón, d Oriol Martinez-Ferraté, d Kerry Wrighton-Araneda, e Diego Cortés-Arriagada, e María Belén Camarada, d, a Jesum Alves Fernandes d b and Gabriel Abarca d a

A series of new triazolium-based supported ionic liquids (SILPs), decorated with Cu NPs, were successfully prepared and applied to the N-arylation of aryl halides with anilines. The triazoles moieties were functionalised using copper-catalysed azide–alkyne cycloaddition. SILP surface characterisation showed a strong correlation between the triazolium cation volume and textural properties. STEM images showed well-dispersed Cu NPs on SILPs with a mean diameter varying from 3.6 to 4.6 nm depending on the triazolium cation used. Besides, XPS results suggest that the Cu(0)/Cu(i) ratio can be modulated by the electronic density of triazolium substituents. XPS and computational analysis gave mechanistic insights into the Cu NP stabilisation pathways, where the presence of electron-rich groups attached to a triazolium ring plays a critical role in leading to a cation adsorption pathway (\( E_{\text{ads}} = 72 \text{ kcal mol}^{-1} \)). In contrast, less electron-rich groups favour the anion adsorption pathway (\( E_{\text{ads}} = 63 \text{ kcal mol}^{-1} \)). The Cu@SILP composite with electron-rich groups showed the highest activity for the C–N Ullmann coupling reaction, which suggests that electron-rich groups might act as an electron-like reservoir to facilitate oxidative addition for N-arylation. This strategy firmly suggests the strong dependence of the nature of triazolium-based SILPs on the Cu NP surface active sites, which may provide a new environment to confine and stabilise MNPs for catalytic applications.

Introduction

Supported metal nanoparticle (MNP) systems have been extensively applied in catalytic processes, in which the metal–support interaction may change the performance, selectivity, and lifetime of the metal catalyst. 1-3 In general, the interactions between catalysts, substrates, and products are governed by the chemical environment, where the stabilisation of the MNPs plays a critical role in the outcome of the reaction. 4-6 In this regard, the functionalisation of mesoporous supports (e.g., SiO2 and Al2O3) with suitable ionic liquids (ILs) arose as a promising alternative, which combines textural properties and a tuneable chemical environment. 6 In this process, a thin IL layer is grafted over mesoporous matrices which controls the textural properties and also creates a confined space for structural and electronic stabilisation of the MNPs. 7 ILs are described as “entropic drivers” since the cation/anion functionalities can be tuned by the proper selection of each ionic moiety. 8 For instance, the use of hydrophobic IL anions can control the MNPs size and distance from the support matrix and thus modify their catalytic selectivity. 9,10 Furthermore, acidic SILPs combined with MNPs can act as multifunctional catalysts for the selective hydrodeoxygenation of aromatic substrates. 11 Despite this exciting research using imidazolium-based SILPs, there is no or very little research using different IL-based SILPs, and a fresh approach can unlock a new generation of SILPs for catalytic applications. 12-14 In this regard, the highly modulated and greener synthesis of a SILP based-triazolium family appears as an exciting novel alternative. In the case of triazoles, the incorporation of different functional groups into the triazolium ring can be easily obtained using copper-catalysed azide–alkyne cycloaddition (CuAAC), which has emerged as a central transformation of click chemistry (Fig. 1) 15,16.
Moreover, the triazoles moieties are obtained with high regioselectivity and yields (>90%) with no generation of harmful by-products, thus being considered an environmentally friendly approach. Here, we present the synthesis of novel triazolium-based SILPs decorated with Cu NPs for the N-arylation of amines with aryl halides. The morphological properties of the SILPs and Cu@SILP composites were extensively investigated by Scanning Electron Microscopy (SEM), Scanning Transmission Electron Microscopy (STEM), X-ray Photoelectron Spectroscopy (XPS) and Brunauer–Emmett–Teller (BET) measurements using the Nguyen-Do method modified by Gun’ko (MND). Furthermore, detailed quantum computational studies were carried out, revealing the IL and Cu NP interaction pathway. The catalytic properties of Cu@SILP composites were evaluated by C-N Ullmann condensation of aryl halides with aniline.

**Experimental**

**Materials**

Aniline, iodobenzene, bromobenzene, chlorobenzene, 3-iodopropyltrimethoxysilane, benzyl bromide, phenylacetylene, NaN₃, CuCl₂, 2H₂O, CuI, NaBH₄, MgSO₄, and BMI-BF₄ were purchased from Sigma-Aldrich. All chemicals were used without further purification, except for DCM, toluene, MeOH, acetonitrile, Et₂O, and EtOAc, which were purified by standard procedures.

**Methods**

¹H NMR measurements of the ILs were performed on a Varian 400 MHz. ²⁹Si and ¹³C cross-polarisation magic angle spinning (CP-MAS) solid-state measurements of the supports were carried out using a Varian 500 MHz spectrometer. The CP MAS spectra were recorded using a 2-channel 4 mm Narrow Bore probe and zirconium rotors. Fourier-transform infrared spectra (FT-IR) were recorded using a 2-channel 4 mm Narrow Bore probe and 1325 cm⁻¹ resolution of zirconium rotors. Fourier-transform infrared spectra (FT-IR) were recorded using a 2-channel 4 mm Narrow Bore probe and 1325 cm⁻¹ resolution of zirconium rotors. Thermoanalytical analyses (TGA) were performed using a Q50 TA TGA operating under a nitrogen gas atmosphere (10 mL min⁻¹). Furthermore, detailed quantum computational studies were carried out, revealing the IL and Cu NP interaction pathway. The catalytic properties of Cu@SILP composites were evaluated by C-N Ullmann condensation of aryl halides with aniline.

**Synthesis of triazole1–3**

Alkyne (5 mmol, 1.0 eq.), alkyl halide (1.0 eq.), and NaN₃ (1.3 eq.) were loaded into a 25 mL round-bottom flask. Then [Cu(PPh₃)₃] (0.05 mol%) was added and dissolved in water (5 mL). The reaction was stirred for 4 h at room temperature, and the progress of the reaction was monitored using thin-layer chromatography (TLC). After its completion, the reaction mixture was filtered, and the residue was dissolved in DCM. The combined organic layer was later concentrated in a vacuum to yield the corresponding triazoles. The products were characterised by spectroscopic analysis as described before.

**Synthesis of SiO₂–1**

Initially, 200 g mesh silica gel was soaked in 30% HCl overnight to hydrolyse its surface. Next, activated silica gel (100 g) was suspended in dry toluene (30 mL) in a round-bottom flask equipped with a reflux condenser under nitrogen. While being stirred, 3-iodopropyltrimethoxysilane (0.05 M) was added dropwise. The suspension was refluxed for 72 h. After cooling, the solid was collected by filtration and exhaustively washed by Soxhlet extraction with ethanol and water, and then dried under reduced pressure to yield iodopropyl–silica gel (SiO₂–1).

**Synthesis of SILP1–3**

SiO₂–1 (5 g) was suspended in dry acetonitrile (30 mL), with stoichiometric amounts of triazoles1–3 (see Scheme 1 for details) and refluxed during 24 h. Then, the particles were collected by filtration and repeatedly washed with Et₂O and water, and then dried off under vacuum at 110 °C for 4 h to obtain SILP1–3 powder.

**Synthesis of Cu@SILP1–3**

A solution of CuCl₂.2H₂O (0.25 mmol) and MeOH (20 mL) was added to SILP1–3 (100 mg) under constant stirring at room temperature for 30 min. A solution of NaBH₄ (5 mmol) dissolved in MeOH (3 mL) was added to the reaction mixture dropwise. The reaction mixture turned black due to the formation of Cu NPs that were washed with MeOH (3 × 10 mL) and Et₂O (3 × 10
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Results and discussion

Preparation and characterisation of SILP1–3

The SILPs were prepared by covalent attachment of triazolium moieties on a silica matrix. The 1,4-disubstituted 1,2,3-triazole moieties were prepared with high regioselectivity and yields (>90%) by click chemistry of an alkyne and in situ generated azide. The prepared triazole was further alkylated to afford 1,3,4-trisubstituted 1,2,3-triazolium compounds (Scheme 1, see the ESI† for details). In this work, we prepared supports with high electronic density (SILP1) by the addition of phenyl and benzyl groups to the triazolium ring and supports with lower electronic density (SILP2–3) replacing the aromatic groups by an alkyl chain. Solid-state 29Si NMR and CP-MAS 13C NMR analyses confirmed the presence of silanol groups and the incorporation of alkyl silanes on the silica surface (Fig. 2a and S1†). The chemical environments of Si atoms are associated with unreacted Si–C bonds and the T2 and T3 species that confirm the presence of unchanged triazolium ILs in SILP1–3 (Tables S1 and S2†). FT-IR spectra show two characteristic peaks, at 1458 cm\(^{-1}\) and 1570 cm\(^{-1}\), associated with the stretching frequencies of the triazolium ring,\(^{43}\) and a sharp peak at 1190 cm\(^{-1}\) related to \(v(\text{Si–O–Si})\), which confirms the silica matrix functionalisation (Fig. S2†). TGA measurements were carried out to verify and quantify the incorporation of alkyl iodosilane and triazolium moieties over the silica surface and to determine their thermal properties. SILP1–3 showed similar weight loss behaviour with increasing temperature. The first weight loss region (ca. 100 °C) is associated with adsorbed water added to a round-bottom flask. The reaction mixture was stirred at 120 °C for 4 h. After the completion of the reaction, the mixture was cooled and extracted with water/EtOAc. The combined organic extracts were dried with anhydrous MgSO\(_4\), and the solvent was removed by rotary evaporation. The conversion and selectivity were determined by GC analysis.

Scheme 1 Synthetic route of SILP materials.

N-Arylation of aryl halides with aniline

In a typical reaction, amine (1.2 mmol), aryl halide (1 mmol), base (2 mmol), Cu@SILP1–3 (1 mol%), and solvent (3 mL) were...
confined into the porous silica matrix, whereas the second weight loss observed at 200–400 °C is ascribed to IL moieties (SILP1–3).\textsuperscript{43} The loading of ILs into the silica matrix was found to be ca. 25–28%, as expected (Fig. S3\textsuperscript{†}). The chemical surface composition of the SILPs was evaluated by XPS (Fig. S4\textsuperscript{†}). The ratio of N 1s to Si 2p corresponds approximately to the expected ratio between triazolium and the silica matrix (Table S3,\textsuperscript{†} entries 1–3). The C 1s and N 1s signals displayed a positive shift for SILP1 against SILP2–3 (Fig. 1b and c), attributed to the electronic effect promoted by the aromatic groups (SILP1), in comparison to ILs with alkyl chains at the triazolium ring (SILP2–3).\textsuperscript{44}

The textural properties of the SILPs were evaluated by BET measurements and analysed by MND that suggests the presence of porous-like cylinders for all SILPs. From BET measurements, the specific surface area of SILP1, SILP2, and SILP3 increases congruently with the total pore volume and average pore radius moving towards the nanopore region (Fig. 2d). The MND analysis shows similar values of $S_{\text{meso}}/S_{\text{BET}}$ (higher than 99%) and an increase of the $S_{\text{meso}}$ absolute value in the order SILP1 < SILP2 < SILP3 (Table S4\textsuperscript{†}). Also, PSD$_V$ (pore size distribution by volume) and PSD$_A$ (pore size distribution by specific surface area) show a shift to smaller radii in SILP1 < SILP2 < SILP3 within the mesoporous range (Table S5\textsuperscript{†}). These results are in good agreement with Nonlocal density functional theory (NLDFT) and Barrett–Joyner–Halenda (BJH) analyses (Fig. S5, Table S5\textsuperscript{†}).\textsuperscript{45} Additionally, the surface roughness degree and porous structure of the SILPs were determined by fractal dimension ($D_f$) analysis employing the Frenkel–Halsey–Hill plot, where for a “flat” surface, $D_f = 2$, whereas for an irregular (real) surface, $D_f$ varies between 2 and 3.\textsuperscript{46} The apparent decreases of roughness SILP1 < SILP2 < SILP3 (Fig. 2e) follows the same trend as the specific surface area and PSD$_V$–PSD$_A$ analysis (Fig. 2d, Tables S4 and S5\textsuperscript{†}).\textsuperscript{47} These results suggest that different substituents over the triazolium ring can directly modify the textural properties of the silica matrix. The addition of large substituents, such as SILP1 (e.g., phenyl-benzyl groups), led to a decrease in the surface area, roughness, and pore volume when compared to SILP2–3. This behavior coincides with observed nitrogen adsorption energy distribution which follows the trend SILP1 < SILP2 < SILP3 (Fig. S6\textsuperscript{†}). FE-SEM showed substantial changes for silica surface roughness after the incorporation of the triazolium-based ILs (Fig. S7\textsuperscript{†}). In addition, the FE-SEM images suggest that the SILP1–3 surface exhibited wavy and rope-like mesoporous domains corroborating the MND analysis.

**Characterisation of Cu@SILP1–3**

STEM measurements showed that Cu NPs were homogeneously dispersed on the surface of SILPs displaying a mean diameter of 3.6 ± 1.0 nm, 4.0 ± 1.3 nm, and 4.6 ± 1.2 nm for Cu@SILP1, Cu@SILP2, and Cu@SILP3, respectively (Fig. 3, S8, S9 and S10\textsuperscript{†}). It can be related to lower roughness and porous size (SILP1 < SILP2 < SILP3), as smaller pores can act as cavities for the stabilisation of smaller NPs. Further insights into Cu NP electronic stabilisation by the ILs were obtained by XPS (Fig. 4a and S11\textsuperscript{†}), XRD (Fig. 4b), and computation analysis (Fig. 5). XPS measurements of Cu@SILP1–3 showed a broad peak at 932.2 eV (Cu 2p$_{3/2}$) and 951.9 eV (Cu 2p$_{1/2}$) that can be assigned to a mixture of valence states Cu(0) and Cu(I).\textsuperscript{48,49} The XRD analysis also confirms the presence of metallic Cu and Cu$_2$O, in agreement with XPS results, as well as STEM analysis that shows Cu NP lattice fringes (111) associated with the FCC crystalline phase of Cu$_2$O (Fig. 4b–inset).\textsuperscript{38} In order to qualitatively estimate the Cu(0) and Cu(I) ratio, Cu LMM analysis was performed indicating a majority presence of Cu(I) valence states for
Stabilisation/interaction between SILP1 and Cu NPs.

Computational studies were performed to understand the stabilisation/interaction between SILP1 and Cu NPs. Cu55 clusters with six Cu(100) and eight Cu(111) faces were chosen in order to represent the Cu NPs synthesised in this work (Fig. 5a and b). Each Cu(100) and Cu(111) pair of faces shares three Cu atoms at the edges [Cu(edge) atoms] (Fig. 5c). Cu55 shows Cu–Cu bond lengths in the range of 2.4–2.5 Å in agreement with crystallographic data for metallic Cu (2.55 Å).32

With the computational method, we first examined the interactions presents on Cu55@SILP1 (Fig. 5d). The triazolium ring interacts with Cu(111) and Cu(edge) atoms at a 2.4 Å bond length, whereas phenyl and benzyl show intermolecular bond lengths of 2.1 and 2.2 Å on Cu(100) and Cu(edge) atoms, respectively. In Cu55@SILP2, the cation interacts through the benzyl hydrogen and the triazole group with Cu(edge) atoms at bond lengths of 2.1 to 2.2 Å, respectively (Fig. 5e). In the case of Cu55@SILP3, the triazole ring interacts with Cu(edge) atoms at a bond length of 2.1 Å, and the butyl groups of the triazolium interact with Cu55 at a hydrogen bond length of 2.0 Å (Fig. 5f). The iodide anion, SILP1–3, interacts with Cu atoms with bond lengths of up to ~2.9 Å. Note that for Cu55@SILP2–3 the triazolium cation tends to interact with two faces of Cu55, whereas the triazolium cation of SILP1 interacts with at least three faces of Cu55. These results demonstrate that SILP1 covers a larger surface of Cu55 compared with SILP2–3, thus providing a high contact between the IL (SILP1) and Cu atoms.

The adsorption energy \( E_{\text{ads}} \) was calculated to evaluate the strength of the interaction between SILP1–3 and Cu55. A competitive model based on co-adsorption of both IL cations and anions on the Cu55 surface was employed considering the free states of each fragment as a starting point (Fig. 6).33 For Cu55@SILP1, the cation pathway was found to be preferred due to its higher \( E_{\text{ads}} \) (72 kcal mol\(^{-1}\)) compared to the anion pathway (\( E_{\text{ads}} = 63 \) kcal mol\(^{-1}\)). The subsequent addition of iodide anions to the system gives an overall \( E_{\text{ads}} \) of 172 kcal mol\(^{-1}\). For Cu55@SILP2–3, the anion pathway is more favoured as the \( E_{\text{ads}} \) is higher (\( E_{\text{ads}} = 63 \) kcal mol\(^{-1}\)) compared with the cation pathway (\( E_{\text{ads}} = 35 \) kcal mol\(^{-1}\)). The complete \( E_{\text{ads}} \) for Cu55@SILP2–3 including their respective cations is 152 kcal mol\(^{-1}\). The distinct pathways of IL adsorption on the Cu55 surface of SILP1 compared with SILP2–3 are found to be related to the minimum energy arrangement of the IL cations on the Cu55 surface (Fig. 5). As discussed previously, SILP1 has two electron-rich groups that interact with three faces of the Cu55 cluster, thus favouring the cation pathway, whereas the relatively weaker interaction of SILP2–3 cations with Cu55 surfaces leads to a more favourable anion pathway. The total comparative \( E_{\text{ads}} \) of Cu55@SILP1 (172 kcal mol\(^{-1}\)) and \( E_{\text{ads}} \) of Cu55@SILP2–3 (152 kcal mol\(^{-1}\)) suggest a weaker interaction of SILP2–3 with the surface of Cu NPs when compared to SILP1. Furthermore, the overall electron transfer between SILP1–3 and Cu55 clusters (\( \Delta q_{\text{Cu55@SILP}} \)) was calculated using the \( \Delta q \) descriptor for each fragment (Table 1). The iodide anion acts as an electron-donating fragment, whereas the electron-acceptor fragments are both Cu55 and triazolium cations. The \( \Delta q_{\text{Cu55@SILP}} \) shows a higher charge transfer from iodide anions to both Cu55 and the triazolium cations according to the following trend SILP1 > SILP2 = SILP3. The charge transferred from iodide is distributed between triazolium and Cu55, which can be divided into two fragmental contributions \( \Delta q_{\text{Cu55}} \) and \( \Delta q_{\text{triazolium}} \). The \( \Delta q_{\text{Cu55}} \) shows values going from 0.19 to 0.49|e|, with Cu55@SILP1 showing...
a higher $|e|$ transfer value. The same trend was observed for $\Delta q_{\text{triazolium}}$, with SILP1 showing higher $|e|$ values than SILP2–3. These results reinforced the major role of the electronic density of triazolium substituents in the stabilisation of Cu NPs.

**Catalytic activity of Cu@SILP1–3**

The catalytic activity of the Cu@SILP1–3 composites was evaluated in the C–N Ullmann coupling reaction to generate diphenylamine selectively from aryl halides (iodo, bromo, and chlorobenzene) and aniline. This model reaction was proposed to investigate the IL steric and electronic environment effect on the Cu NP surface during the catalytic cycle. Catalytic experiments using KOH, K$_2$CO$_3$, Cs$_2$CO$_3$ and N(Et)$_3$ were performed (Table S6†). KOH has shown the highest catalytic conversion compared to other bases used, which we ascribe to its stronger basicity that may enhance the formation of the amido-complex, and thus KOH was used in all reactions. All SILPs displayed high selectivity towards the N-arylated product, with traces of triphenylamine and biphenyl as side-products (Table 2). The catalytic activity of Cu@SILP1–3 was observed in the following order Cu@SILP1 > Cu@SILP2 > Cu@SILP3 (entries 1–3). It is important to mention that no significant difference in the catalytic activity between Cu@SILP1 and Cu@SILP2 was observed for iodobenzene, whereas for bromo- and chloro-derivatives a significant difference was observed (entries 4, 5, 7 and 8). This behaviour can be related to the faster C–X activation for iodo-derivatives compared to bromo- and chloro-derivatives. These results are in agreement with previous reports where the coupling product can be modulated by the reactivity of the aryl halide compound ($R$–I $\gg$ R–Br $>$ R–Cl). Additionally, neat Cu$_2$O NPs were tested in the absence of SILPs (Table 2, entry 10). No significant diphenylamine production was observed, which might be due to a weak stabilisation of Cu(i)/Cu(III) moieties during the catalysis cycle, in the absence of ILS (SILPs).

Based on these results, two possible behaviours can be suggested: (i) the presence of an electron-rich IL layer (SILP1 > SILP2 > SILP3) may favour the oxidative addition that involves Cu(i)/Cu(III) as a rate-determining step (RDS) of this reaction under these conditions. Therefore, it would be reasonable to conclude that a preferential Cu(i) species over the surface might facilitate the oxidative addition. (ii) Reduced yields were achieved for different halogen sources, because of the low reactivity.

### Table 1

Fragmentary electron transfer ($\Delta q|e|$) between the fragments Cu$_{55}$, IL anion, and cation for Cu$_{55}$@SILP1–3

| Samples          | $\Delta q_{\text{Cu@SILP}}$ | $\Delta q_{\text{Cu}_{55}}$ | $\Delta q_{\text{triazolium}}$ |
|------------------|-----------------------------|-----------------------------|-------------------------------|
| Cu$_{55}$@SILP1  | 0.88                        | 0.49                        | 0.39                          |
| Cu$_{55}$@SILP2  | 0.52                        | 0.19                        | 0.34                          |
| Cu$_{55}$@SILP3  | 0.58                        | 0.28                        | 0.30                          |

### Table 2

Catalytic properties of Cu@SILP1–3 for C–N Ullmann coupling reactions

| Entry | Cat     | X   | Conv (%) | Sel (%) | TOF$^a$ (h$^{-1}$) |
|-------|---------|-----|----------|---------|--------------------|
| 1     | Cu@SILP1 | I   | 88       | 95      | 374                |
| 2     | Cu@SILP2 | I   | 83       | 94      | 311                |
| 3     | Cu@SILP3 | I   | 77       | 94      | 301                |
| 4     | Cu@SILP1 | Br  | 61       | 92      | 184                |
| 5     | Cu@SILP2 | Br  | 48       | 91      | 150                |
| 6     | Cu@SILP3 | Br  | 44       | 93      | 164                |
| 7     | Cu@SILP1 | Cl  | 44       | 90      | 132                |
| 8     | Cu@SILP2 | Cl  | 31       | 86      | 103                |
| 9     | Cu@SILP3 | Cl  | 30       | 86      | 92                 |
| 10    | Cu$_2$O@SiO$_2$ | I | 35       | 81      | 161                |

*a Reaction conditions: Cu@SILP1–3 (1 mol%), aniline (1.2 mmol), aryl halide (1 mmol), KOH (2 mmol), DMSO (3 mL), 120 °C, and 4 h. *a Conversion determined by GC. TOF = mol diphenylamine converted/(mol Cu surface $\times$ time). Calculated from the slope of plots of $\text{TON} = t$ at low substrate conversions. *Neat Cu$_2$O NPs supported on SiO$_2$ without an ionic liquid (mean diameter 9.5 ± 2.0 nm).
observed for deactivated aryl halides on nucleophilic substitutions (C–X: I > Br > Cl).37
In this regard, the molecular electrostatic potential (MEP) was calculated for Cu@SILP1–3. The MEP results show that charge polarisation takes place on Cu NPs after the charge transfer between SILPs and Cu NPs (Fig. 7). This process occurs due to a surface charge imbalance between the iodide anion and neighbouring Cu(edge) atoms that present higher positive polarisation.39 Therefore, as Cu@SILP1 presented a higher overall electron transfer (between the IL and Cu), it can increase its catalytic activity for Ullmann coupling, considering the oxidative addition step as the RDS.54 In this regard, we suggest that the IL cation (SILP1) might work like an “electron-like reservoir” that improves the rates of Cu(I)/Cu(III) redox reactions, and thus, its catalytic activity compared to SILP2–3 (Tables 1 and 2). These results are supported by the Brønsted–Evans–Polanyi (BEP) relationship59 that links the MNP stabilisation with catalytic activity. It employs the E_ads and BEP relationship to explain the catalytic activity of MNPs.59 A recent study based on Cu clusters of different sizes (from Cu13 to Cu77, including Cu55 clusters) compared the E_ads and the catalytic activity in the CO2-hydrogenation reaction to methanol showing similar results to this work.54

Conclusions
A new series of triazolium-based SILPs were successfully synthesised and decorated with Cu NPs. The triazolium ring was modified with aromatic and alkyl groups in order to study the influence on the steric and electronic properties of the SILPs. The textural analysis demonstrated that the surface area, porous volume, and roughness decrease according to the volume of substituents in the triazolium cation, thus showing the high tunability of triazolium-based SILPs. Moreover, the stabilisation of Cu NPs seems to be related to the surface properties of the obtained SILPs as the mean diameter of the Cu NPs decreases according to the SILP porous size.

Furthermore, XPS and computational investigations showed that the electronic density of the triazolium cation substituents plays a vital role in the stabilisation pathway of the Cu NPs. In the case of SILP1, two aromatic groups (electron-rich) were incorporated into the triazolium ring leading to a preferential stabilisation of Cu NPs through the cation pathway. In opposition, when alkyl groups are added, the anion pathway is more favourable. The increase of overall E_ads calculated for Cu@SILPs follows the order Cu@SILP1 > Cu@SILP2 ≈ Cu@SILP3. The computational studies suggest a stronger interaction between SILP1 and Cu NPs occurs due to the higher charge transfer between the IL and Cu atoms. The catalytic results revealed that Cu@SILP1 displayed the best catalytic activity compared to Cu@SILP2–3 for the C–N Ullmann coupling reaction. We suggest that the IL cation of SILP1 may act as an electron-like reservoir adjusting the Cu atom electronic environment (increasing the Cu(I)/Cu(III) ratio). This behaviour may favour the oxidation state as a RDS, leading to a higher activity compared to Cu@SILP2–3.

Indeed, the triazolium-based SILP approach is easily transferable for other catalytic platforms, and thus this work will open new avenues to design supported metal catalysts for many other catalytic applications.

Conflicts of interest
The authors declare no conflict of interest.

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