Scalable two-step annealing method for preparing ultra-high-density single-atom catalyst libraries

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The stabilization of transition metals as isolated centres with high areal density on suitably tailored carriers is crucial for maximizing the industrial potential of single-atom heterogeneous catalysts. However, achieving single-atom dispersions at metal contents above 2 wt% remains challenging. Here we introduce a versatile approach combining impregnation and two-step annealing to synthesize ultra-high-density single-atom catalysts with metal contents up to 23 wt% for 15 metals on chemically distinct carriers. Translation to a standardized, automated protocol demonstrates the robustness of our method and provides a path to explore virtually unlimited libraries of mono- or multimetallic catalysts. At the molecular level, characterization of the synthesis mechanism through experiments and simulations shows that controlling the bonding of metal precursors with the carrier via stepwise ligand removal prevents their thermally induced aggregation into nanoparticles. The drastically enhanced reactivity with increasing metal content exemplifies the need to optimize the surface metal density for a given application. Moreover, the loading-dependent site-specific activity observed in three distinct catalytic systems reflects the well-known complexity in heterogeneous catalyst design, which now can be tackled with a library of single-atom catalysts with widely tunable metal loadings.

Improving the atom economy of chemical transformations and ensuring the maximal utilization of scarce catalytic materials are central targets for sustainable chemistry. Heterogeneous single-atom catalysts (SACs), integrating monodisperse atomic metal centres with tailored coordination environments, demonstrate the potential to fulfill both of these objectives in several energy-related transformations. A fundamental challenge in implementing this pioneering class of catalysts in many technical applications is the lack of synthetic routes enabling the preparation of these catalysts with high surface densities. Achieving the latter is particularly important to maximize the productivity per unit reactor volume or area in large-scale processes. Taking nickel SACs, which exhibit highly selective performance in the electrochemical reduction of carbon dioxide, as a representative example, a literature analysis shows that most of the materials reported to date are based on carbon-related materials and contain metal contents centred around 1 wt%, with exceptional cases up to 7 wt% (Fig. 1a). These values are significantly lower than the expected theoretical capacity of these carriers for anchoring metal atoms.

Of the distinct synthesis strategies reported for SACs, the post-synthetic introduction of metals via wet deposition routes such as impregnation is among the most amenable to scale-up. This approach also favours surface localization, unlike direct synthesis approaches such as pyrolysis, where a large fraction of the active phase may become embedded in the bulk of carrier materials. However, when introducing high metal contents, it is difficult to prevent the undesired aggregation of metal species into clusters and nanoparticles especially after applying conventional thermal treatments to remove counterions from common metal precursors (Fig. 1b,c and Supplementary Fig. 1). Strategies to circumvent this by avoiding metal salt precipitation or using equilibrium adsorption typically restrict the amount deposited on carriers. No published study has achieved ultra-high-density (UHD) SACs—defined herein as having metal contents over 10 wt% for carbon-based catalysts—by postsynthetic routes, or demonstrated their enhanced productivity in catalytic applications. Another scarcely addressed aspect is the extension to multimetallic systems, which is relevant for the development of technical catalysts since they often incorporate two or more metals, for example, as co-catalysts, promoters or stabilizers. Beyond the selection of carriers with abundant surface anchoring sites for metal atoms, the synthesis mechanism remains poorly understood at the molecular level.

Two-step annealing approach
The two-step annealing approach introduced here comprises a versatile and scalable method for the preparation of mono- or multimetallic...
UHD-SACs of more than 15 transition metals on carriers with distinct chemical natures, such as nitrogen-doped carbon (NC), polymeric carbon nitride (PCN) and metal oxides (Supplementary Fig. 2). The approach relies on controlling ligand removal from metal precursors and the associated interactions with the carrier (Fig. 1d). The strength of the proposed method lies in effectively saturating the surface with metal and retaining a high metal coverage by a selective anchoring mechanism that maximizes the probability of bonding the metal to all available coordination sites and enables the removal of unbound species via washing, thereby preventing metal sintering in the subsequent high-temperature step to remove residual ligand and permitting the stabilization of much higher metal contents compared to conventional impregnation routes. The method uses aqueous solutions of common metal precursors (chlorides, sulfates or acetates) or mixtures thereof where multimetallic systems are targeted. To regulate ligand removal, the temperature of the first annealing step ($T_1$) needs to be lower than that of the decomposition temperature of the metal precursor; the most labile in the preparation of multimetallic SACs. For this reason, the use of metal chlorides provides greater control due to the strong intrinsic metal–chlorine bonds, which permits the use of higher temperatures in the first annealing step (Supplementary Table 1). The chemisorbed metal precursors are subsequently transformed into UHD-SACs with well-defined atomic structures by a second annealing step at a higher temperature ($T_2$) to remove the remaining ligands.

Analysis by inductively coupled plasma atomic emission spectroscopy (ICP-AES) shows that the two-step annealing method achieves metal contents higher than 10 wt%, in several cases exceeding 20 wt%, on both NC and PCN carriers for all the metals studied (Fig. 1c). Lower metal loadings of between 2 and 10 wt% result when using CeO$_2$ as the carrier, reflecting its higher relative molecular weight versus carbon-based materials. To provide a universal basis for comparison, the surface areal density in atoms per square nanometre was evaluated based on the measured Brunauer–Emmett–Teller surface area of the carrier (Supplementary Fig. 3). UHD-SACs derived from PCN evidenced the highest densities for all metals, consistent with the abundance of metal coordination

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**Fig. 1 | Synthesis of UHD-SACs.**

**a.** Literature survey on the metal contents reported in nickel SACs revealing the predominant use of carbon-based carriers. **b.** Comparison of the size of nickel species obtained on NC via a conventional impregnation strategy involving a single thermal treatment and the two-step annealing strategy introduced in this work as a function of the metal content. **c.** ADF-STEM images of 10 wt% Ni–NC samples by a single thermal treatment (top) and a two-step annealing strategy (bottom). Scale bars, 100 nm (inset, 1 nm). **d.** Strategy for the preparation of UHD-SACs. **e.** Metal loadings achieved in this study on NC, PCN and CeO$_2$ supports.
sites presented by this carrier due to the intrinsic crystalline structure based on heptazine units. Discrete ranges of values were observed for first-row (>9 atoms per nm²), second-row (~6 atoms per nm²) and third-row (<5 atoms per nm²) transition metals. The significantly higher values for first-row transition metals could be indicative of some percolation into the lattice. Values between 2.5 and 6 atoms per nm² for CeO₂ are significantly higher than previous estimates for high-loading catalysts based on this carrier. Values between 0.4 and 1 atoms per nm² are observed for NC, approaching the upper limit of 1.3 atoms per nm² calculated from the nitrogen content in the carrier.

To further confirm the generality of the approach, we extended the synthesis to other commonly used supports, including titanium dioxide and alumina (Supplementary Fig. 4). It is noteworthy that the present strategy is based on the premise that a carrier with surface functionalities able to react with the deposited metal in the investigated temperature range is used. For this reason, platinum single atoms cannot be isolated on amorphous silica with the presented method due to the weak bond between surface oxygen and platinum (Supplementary Fig. 5). Additionally, the simultaneous introduction of multiple metal precursors behaves additively and successfully yields multimetallic UHD-SACs, demonstrating the scope of the method to produce a widely tuneable materials platform (Supplementary Fig. 6).

Confirming the UHD atomic dispersion
The atomic metal dispersions of the obtained UHD-SACs on NC were directly imaged by annular dark-field scanning transmission electron microscopy (ADF-STEM, Fig. 2a), revealing densely populated features ascribed to isolated atoms. Low-magnification elemental mapping by energy-dispersive X-ray spectroscopy (EDS) of NC-based UHD-SACs (Supplementary Figs. 7 and 8). The bright dots in the high-magnification ADF-STEM images of platinum on NC, PCN and CeO₂ exhibit diameters of approximately 0.2 nm, reflecting the convolution of an isolated platinum atom with the electron probe (Fig. 2b and Supplementary Fig. 9). PCN-supported UHD-SACs display high metal atom densities (Supplementary Fig. 10a) and heavy elements such as tungsten, iridium and platinum, which are distinguishable on CeO₂, also evidence atomic dispersion (Supplementary Fig. 10b). Elemental uniformity is also visible at a larger scale, as seen in low-magnification...
bands centred at 2,117 cm\(^{-1}\). This combination of metals was also confirmed the atomic dispersion of the constituent metals in NC, Pt\(_1\)/PCN and Pt\(_1\)/CeO\(_2\) reveals only one set of CO absorption platinum species are positively charged, and the valence state on f\(_{\text{Pt}}\) X-ray absorption near-edge structure (XANES) spectra and the Fourier-transformed extended X-ray absorption fine structure (EXAFS) spectra of all the UHD-SAC samples exhibit major peaks around 1.5 Å, corresponding to M–N/O bond lengths (Fig. 2d). No signatures of metal–metal coordination between 2.1 and 2.5 Å are observed in any of the samples, confirming the absence of metal nanoparticles or clusters. Consistently, infrared spectroscopy of adsorbed CO on Pt\(_1\)/NC, Pt\(_1\)/PCN and Pt\(_1\)/CeO\(_2\) reveals only one set of CO absorption bands centred at 2.117 cm\(^{-1}\) (Fig. 2d). No infrared peaks in the range of 2,070–2,090 cm\(^{-1}\) corresponding to platinum nanoparticles can be observed\(^{12}\). These results further validate the atomic-resolution ADF-STEM analyses, demonstrating the successful preparation of UHD-SACs with various metals and carriers. The Pt L\(_{\text{III}}\)-edge X-ray absorption near-edge structure (XANES) spectra and the Pt 4f X-ray photoelectron (XPS) spectra both indicate that all the platinum species are positively charged, and the valence state on NC and PCN is slightly lower than on CeO\(_2\) (Supplementary Fig. 16)\(^{11,13}\). K-edge XANES corroborates this trend for nickel, where the intensity of the white line of Ni\(_1\)/NC and Ni\(_1\)/PCN is similar and lower than that observed for Ni\(_1\)/CeO\(_2\) (Supplementary Fig. 17). ADF-STEM images of Ni and Pd and of Ni–Pd–Pt UHD-SACs also confirmed the atomic dispersion of the constituent metals in multimetallic systems (Fig. 3a–b). This combination of metals was selected to facilitate characterization, permitting the local identification of multimetallic SACs based on analysis of the intensity profiles over the different atoms, wherein heavier metal atoms exhibit higher contrast and wider peaks in terms of full width at half maximum (Supplementary Fig. 18). The spatial uniformity of individual elements in the Ni–Pd–Pt UHD-SAC is confirmed in the elemental maps acquired by energy-dispersive X-ray spectroscopy (Fig. 3c).

**Mechanistic insights into the controlled ligand removal**

To understand the greater effectiveness of the two-step annealing strategy, we followed the evolution of the metal and anchor site during the synthesis by EXAFS, XANES, XPS and density functional theory (DFT) modelling. Starting from nickel chloride (NiCl\(_2\)) and NC, we identify adsorbed NiCl\(_2\) (NiCl\(_2\)/NC), partially chlorinated (NiCl\(_2\)/NC) and the fully bound Ni (Ni\(_1\)/NC) single atoms as key intermediates or products, respectively (Supplementary Fig. 20a). Whereas X-ray diffraction analysis of the materials before and after the respective annealing steps confirms the disappearance of characteristic NiCl\(_2\) reflections, it provides no information on changes in the local environment (Supplementary Fig. 20b). Comparison of both XANES and EXAFS spectra evidences appreciable differences between the intermediates. A single peak at around 1.97 Å for NiCl\(_2\)/NC corresponds to the Ni–Cl bond from pure NiCl\(_2\). After low-temperature annealing, a strong signal appears at around 1.40 Å for NiCl\(_2\)/NC, consistent with the newly formed Ni–N bonds\(^{11,13}\) (Fig. 5a). Concurrently, the peak position of the Ni–Cl bond moved from 1.97 to 1.75 Å due to the transformation of NiCl\(_2\)/NC to a NiN\(_2\)Cl structure. From the fitting results (Supplementary Fig. 21 and Supplementary Table 2), the coordination numbers of the nitrogen and chlorine atoms in the first coordination sphere of nickel are around 4.8 and 0.9, respectively, with a square-pyramidal configuration for Ni–N bonding. Following the high-temperature annealing forming Ni\(_1\)/NC, only the Ni–N peak at around 1.40 Å is seen with no evidence of Ni–Cl bonds, indicating that the anchored nickel of shaking instead of sonication for mixing the carrier with the aqueous solution of the metal precursor (Fig. 4b). Accordingly, the procedure could readily obtain nickel SACs over a broad range of metal content (Fig. 4c). Slightly lower values observed in the samples prepared by the automated compared to the laboratory protocol originate from operating under ambient rather than reduced pressure during the mixing step, which decreases surface wetting of the micropores. Given the virtually limitless potential combinations of metals, carriers and stoichiometry, the possibility to automate the synthesis protocol enabling the reproducible preparation of well-defined UHD-SAC libraries is highly attractive for the derivation of more precise synthesis–property–performance relationships to guide their design in targeted applications. Besides versatility, another important feature of our approach is the scalability. Because the two-step annealing approach employs simple unit operations (impregnation, thermal treatment, washing and filtration), it readily translates to the kilogramme-scale production of UHD-SACs, with no appreciable differences in the properties of the resulting material (Supplementary Fig. 19).

**Amenability to automation and scale-up**

Motivated by growing efforts across catalysis research towards automation and standardization of procedures, which are most advanced in computational chemistry and catalyst characterization and testing, we translated the two-step annealing approach to an automated synthesis protocol. The use of a robotic platform (Fig. 4a) permits precise control over the quantity and speed of metal addition and over the extent of sample washing, and requires only minor adaptions of the laboratory-based protocol such as the use of shaking instead of sonication for mixing the carrier with the aqueous solution of the metal precursor (Fig. 4b). Accordingly, the procedure could readily obtain nickel SACs over a broad range of metal content (Fig. 4c). Slightly lower values observed in the samples prepared by the automated compared to the laboratory protocol originate from operating under ambient rather than reduced pressure during the mixing step, which decreases surface wetting of the micropores. Given the virtually limitless potential combinations of metals, carriers and stoichiometry, the possibility to automate the synthesis protocol enabling the reproducible preparation of well-defined UHD-SAC libraries is highly attractive for the derivation of more precise synthesis–property–performance relationships to guide their design in targeted applications. Besides versatility, another important feature of our approach is the scalability. Because the two-step annealing approach employs simple unit operations (impregnation, thermal treatment, washing and filtration), it readily translates to the kilogramme-scale production of UHD-SACs, with no appreciable differences in the properties of the resulting material (Supplementary Fig. 19).

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**Fig. 3** | Visualization of multimetallic UHD-SACs. a,b, Atomic-resolution ADF-STEM images of Ni and Pd (a) and Ni–Pd–Pt multimetallic UHD-SACs (b) on NC. c, EDS map of Ni, Pd and Pt in Ni\(_1\)Pd\(_1\)Pt\(_1\)/NC. Scale bars, 1 nm (a and b) and 50 nm (c).
one HCl molecule is exothermic and occurs spontaneously, forming the first transition state (TS1) with a barrier of 1.39 eV and endothermic reaction energy of 0.14 eV. The following rearrangement releasing a second HCl molecule drives the transformation of NiCl/NC to Ni/NC (Int2). This process requires a barrier of 0.72 eV with endothermic reaction energy of 0.28 eV. Hence, the formation of Ni/NC proceeds via two endothermic and one spontaneous step, which is in line with our experimental procedure, where two annealing steps are needed. A similar landscape is obtained for the formation of Fe1/NC, Cu1/NC and Ni1/Co3 using metal chloride precursors and of Ni1/NC using NiSO4 as a metal precursor (Supplementary Figs. 22 and 23).

**Enhanced productivity of UHD-SACs**

As a proof of concept, we evaluated the catalytic performance of representative UHD-SACs in applications where supported single atoms demonstrate promising technological potential. Nickel SACs excel in the electrochemical reduction of carbon dioxide to carbon monoxide (Supplementary Fig. 24). For the Ni1/NC catalyst, the only products detected are gas-phase CO and H2, and no liquids in the voltage range from 0 to −1.15 V versus a reversible hydrogen electrode (RHE). The linear sweep voltammetry (LSV) curves reveal significantly higher current density for the UHD Ni1/NC than the low-density Ni/NC (Fig. 6a). In terms of Faradaic efficiency towards CO, the UHD and low-density Ni/NC catalysts exhibit a maximum of 97% at around −0.95 and −0.75 V versus RHE, respectively (Fig. 6b). Although the nickel content in UHD Ni1/NC catalyst is ten times higher than in the low-density Ni/NC catalyst, it conserves the high selectivity towards CO. Notably, the UHD Ni1/NC catalyst maintains greater than 86% Faradaic efficiency even at a negative potential of −1.15 V versus RHE. The activity scaled with the nickel content, showing a gradual increase in the current density of CO for metal contents ranging from 1.6 to 16.3 wt% (Supplementary Fig. 25). The site-specific activity is similar across the range of investigated loadings with a slightly lower (~30%) activity per site for the highest metal contents in the voltage range from −0.55 to −1.05 V (Supplementary Fig. 25). An approximately 30% loss in activity per site is probably due to the percolation of a small portion of nickel metal into the inner part of the carbon carrier. These nickel SACs were buried inside would not be directly accessible by CO2, leading to a decrease in site-specific reactivity because the total amount of nickel SACs is counted in the calculation. It is expected that for metal species with small atomic radius and high thermal mobility, an increase of metal loading may also result in a slightly higher ratio of single atoms embedded in the bulk of host materials, consistent with our experimental observations (Supplementary Fig. 25). DFT modelling reveals the favourable energetics of the CO2 reduction reaction over parasitic hydrogen formation at the Ni1/NC site (Supplementary Fig. 26). The main advantage of the UHD Ni1/NC catalyst for CO2 reduction is that it delivers a much higher CO partial current density due to the high abundance of nickel active sites.
compared to the conventional low-density counterpart. Furthermore, it preserves stable CO\textsubscript{2} reduction at constant current density for CO (20 mA cm\textsuperscript{-2}) during 48 h of continuous operation (Fig. 6c); the current density and Faradaic efficiency profiles during two consecutive linear voltammetry sweeps are identical (Supplementary Fig. 27). ADF-STEM images measured after the electrochemical stability test show identical well-dispersed nickel atoms as seen in the fresh Ni\textsubscript{1}/NC catalyst (Supplementary Fig. 28).

In addition to electrochemical applications, SACs are the preferred nanostructure in several thermally driven reactions. Due to their structural similarities, they have attracted significant interest for the replacement of homogeneous catalysts in organic transformations\textsuperscript{14}. Evaluation of Cu\textsubscript{1}/PCN catalysts in the azide–alkyne cycloaddition reaction of 4-ethynylanisole with benzyl azide demonstrated a remarkable enhancement in the performance with increasing metal content, from virtually negligible activity in the conventional SAC to yielding the desired product with 92% isolated yield over ten consecutive runs (Fig. 6d,e and Supplementary Fig. 29). As a result, the site-specific activity for Cu\textsubscript{1}/PCN with the highest metal content is observed to be seven times higher than that for Cu\textsubscript{1}/PCN with the low metal content (Supplementary Fig. 30). The enhanced productivity not only arises from the increased number of active sites per unit reactor volume or area but also receives a significant contribution from the dramatically increased site-specific activity with increasing copper loading for cross-coupling reactions. This experimental result indicates that well-isolated single-atom sites present in low-loading SACs may not be able to co-adsorb and activate the multiple reactants required for organic cross-coupling reactions, while the synergistic effect derived from the presence of neighbouring single-atom sites promotes a dramatically enhanced site-specific activity for UHD-Cu\textsubscript{1}/PCN SACs. In a more classical gas-phase application, platinum single-atom catalysts were recently shown to comprise stable and active catalysts as alternatives to currently applied mercury-based catalysts in the production of vinyl chloride monomer (VCM) via acetylene hydrochlorination\textsuperscript{40}. The presence of halogen species in the reaction mixture induces mobility of supported metal atoms but increasing the strength of interaction with the carrier too much renders them inactive. Here the initial yield of VCM over Pt\textsubscript{1}/NC catalysts directly correlated with the metal content of the catalysts applied, but an optimal metal content of 4.4 wt% was observed for stable performance, above which a gradual drop in VCM yield was observed with time-on-stream due to sintering (Fig. 6f). The initial yield of VCM over Pt\textsubscript{1}/NC SAC scales with platinum content, which

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**Fig. 5 | Mechanistic investigation of the synthesis of UHD-SACs.**

**a**, Fourier-transformed EXAFS (a) and N 1s XPS (b) spectra of NiCl\textsubscript{2}/NC, NiCl/NC and Ni\textsubscript{1}/NC. R is the radial distance. **c**, DFT-calculated energy (E) pathway for the formation of Ni–N\textsubscript{4} from the NiCl\textsubscript{2} precursor with representative transition states and intermediates depicted inset. \(\Delta G\) is the change in Gibbs free energy and \(\Delta H\) is the change in enthalpy.
also suggests that the reactivity of Pt1/NC SACs indeed scales with metal loading. The larger atomic radius of platinum disfavors its thermal diffusion and percolation into inner bulk. Nearly all the platinum atoms anchored on the surface of NC are readily accessible by the reactants, leading to nearly zero loss of activity per site with increasing platinum loading (Supplementary Fig. 31). The enhanced performance evidenced by these results demonstrates the exciting scope of UHD-SACs to improve both the productivity and understanding of mono- or multimetallic single atom systems in a broad variety of catalytic applications in the near future.

Discussion

The scalable wet-chemistry method established here to prepare UHD-SACs across a broad range of transition metals on chemically distinct carriers comprises a breakthrough towards their practical implementation. Our experimental and theoretical analysis showed that the success of the two-step annealing approach at stabilizing high surface densities of isolated metal atoms derives from the selective bonding of metal precursors to the carrier. This controlled interaction enables the removal of weakly interacting species that would otherwise aggregate upon complete decomposition. Besides monometallic systems, a library of multimetallic systems can be readily accessed, evidencing the general applicability of the two-step annealing method. The reproducibility of the synthesis upon automation and at large scale demonstrates the robustness of the approach, paving the way for the widespread application of UHD-SACs in sustainable chemical and energy transformations.

Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41565-021-01022-y.

Received: 5 April 2021; Accepted: 29 September 2021; Published online: 25 November 2021

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Methods

Synthesis of NC. A two-dimensional zeolitic imidazolate framework (2D-ZIF-8) was synthesized as the NC precursor. Typically, Zn(NO3)2·6H2O (12.75 g) and 2-methylimidazole (29.15 g) were dissolved separately in deionized water (1,000 ml). Then the two aqueous solutions were rapidly mixed and vigorously stirred for 2 h. The resulting white precipitate was left undisturbed for 12 h. The product was collected by centrifugation, then washed with water and ethanol, and subsequently dried at 80 °C overnight. To prepare NC, the 2D-ZIF-8 (5 g) was mixed with KCl (100 g) in deionized water (400 ml) and then dried under rotary evaporation. After drying (120 °C overnight), the KCl intercalated powder was heated to 700 °C (heating rate, 2 °C min−1) for 5 h in a nitrogen flow. The product was washed with hydrochloric acid (2 M), deionized water, and ethanol, then dried at 80 °C overnight.

Synthesis of Ni1/CeO2. To achieve 30 wt% metal feeding, NiCl2·6H2O (174 mg) and CeO2 (100 mg) were dispersed in 20 ml ethanol solution, sonicated for 10 min, and dried first by rotary evaporation and then in an oven at 80 °C. For low-temperature annealing, the powder was heated to 300 °C (heating rate, 5 °C min−1) for 5 h in a nitrogen flow. After thorough washing using a water–ethanol mixture, the obtained powder was washed thoroughly and then dried at 80 °C overnight. To prepare NC, the 2D-ZIF-8 (5 g) was mixed with KCl (100 g) in deionized water (400 ml) and then dried under rotary evaporation. After drying (120 °C overnight), the KCl intercalated powder was heated to 700 °C (heating rate, 2 °C min−1) for 5 h in a nitrogen flow. The product was washed with hydrochloric acid (2 M), deionized water, and ethanol, then dried at 80 °C overnight.

Synthesis of Ni1Pd1Pt1/NC. The nickel, palladium and platinum precursors were, respectively, NiCl2·6H2O (24 mg), PdCl2 (20 mg) and NC (100 mg) dispersed in 0.5 M HCl solution (20 ml), sonicated for 10 min and dried first by rotary evaporation and then in an oven at 80 °C. For the low-temperature annealing, the powder was heated to 200 °C (heating rate, 5 °C min−1) for 5 h in a nitrogen flow. After thorough washing using dimethylsulfoxide (DMSO) and water, the dried powder (80 °C) was subjected to a high-temperature annealing at 500 °C (heating rate, 2 °C min−1) for 5 h in a nitrogen flow.

Electrochemical tests using Ni1/NC. All CO reduction experiments were performed using a three-electrode system connected to an electrochemical workstation (CHI 760E). Ag/AgCl with saturated KCl solution and platinum mesh served as the reference electrode and the counterelectrode, respectively. The electrocatalyst was dispersed in 0.1 M HCl solution and sonicated for 10 min under static air.

Azide–alkyne cycloaddition using Cu/PCN. Azide–alkyne cycloaddition was carried out in a 20 ml glass tube with Cu/PCN catalyst (2 mol% Cu, 0.5 mmol of 4-ethynylbenzene, 1.5 mmol of benzyl azide and 8 ml of 1:1 water/tert-butanol mixture and stirred at 60 °C for 12 h. H NMR and 13C NMR spectra were recorded on Bruker 500 MHz spectrometer using residue solvent peaks as internal standards (DMSO-d6). GC yield was determined by using decane as an internal standard.

Aqueous hydrochlorination using Pt/NC. Aqueous hydrochlorination was performed on a Bruker D8 Focus powder X-ray diffractometer using Cu Kα radiation (40 kV, 40 mA) at room temperature. Transmission electron microscopy (TEM) images were obtained with a JEOL Titan 80-300/STEM (TEM) column operated at 300 kV. X-ray photoelectron spectroscopy (XPS) measurements were carried out in a custom-designed ultrahigh-vacuum system with a base pressure lower than 2 × 10−10 mbar. Al Kr (photoelectron energy 1.4867 eV) was used as the excitation source for XPS. The metal loadings in all the samples were measured by ICP-AES. N2 isotherms were measured at −196 °C using a Quantachrome Instruments AutosorbQ (Boynton Beach, FL). Fourier transform infrared spectra for CO adsorption were obtained at 25 °C on a Bruker Equinox 55 spectrometer equipped with a mercury cadmium telluride detector. The XANES and EXAFS measurements were carried out at the XAFCA beamline of the Singapore Synchrotron Light Source (SLSL) and Shanghai Synchrotron Radiation Facility (SSRF), Shanghai Institute of Applied Physics (SINAP). A Si (111) double-crystal monochromator was used to filter the X-ray beam. Metal foils were ultrasonically cleaned in acetone, methanol, and deionized water, and then dried under filtered nitrogen. All calculations were performed using DFT with Vienna Ab initio Simulation Package (VASP)44,45. The projector augmented-wave method46 was used to represent the core electron states. The generalized gradient approximation with Perdew–Burke–Ernzerhof exchange-correlation functional47 and a plane wave representation for the wave function with a cut-off energy of 500 eV were used. Lattice parameters and all atoms were fully relaxed for total energy optimization. The Monkhorst–Pack k-point grid was used to model the graphene surface and the Brillouin zones were sampled by a Monkhorst–Pack k-point mesh with a 3 × 3 × 1 k-point grid. A 15 Å vacuum space was set in the z-direction, to avoid periodic image interactions. The transition states were searched using climbing image nudged elastic band and dimer methods48 and further confirmed by having only one imaginary vibrational frequency.

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**Acknowledgements**

J. Lu acknowledges support from MOE grant (R-143-000-B47-114), the Ministry of Education (Singapore) through the Research Centre of Excellence program (Award EDUN C-33-18-279-V12, Institute for Functional Intelligent Materials) and the National University of Singapore Flagship Green Energy Program (R-143-000-A55-646). X.Z. acknowledges support from a Presidential Postdoctoral Fellowship, Nanyang Technological University, Singapore via grant 03H1S000973C150. S.M., D.F.A., and J.P.-R. acknowledge funding from the NCCR Catalysis, a National Centre of Competence in Research funded by the Swiss National Science Foundation. Jun Li acknowledges financial support by the National Natural Science Foundation of China (grant number 22033005) and the Guangdong Provincial Key Laboratory of Catalysis (2020B121201002). Computational resources were supported by the Center for Computational Science and Engineering (SUSTech) and Tsinghua National Laboratory for Information Science and Technology. We would like to acknowledge the Facility for Analysis, Characterization, Testing and Simulation, Nanyang Technological University, Singapore, for use of their electron microscopy facilities.

**Author contributions**

X.H. and J. Lu conceived and designed the experiments. J.P.-R. conceived the automated synthesis protocol. J. Lu and J.P.-R. supervised the project and organized the collaboration. X.H. performed materials synthesis. S.M. and D.F.A performed automated synthesis. X.H. and T.S. performed the activity test. S.X. performed the XAFS measurement. X.Z., H.X. and D.K. performed the electron microscopy experiments and data analysis. C.S. helped to perform the CO-DRIFTS measurements. Jing Li performed the XPS measurements. Z.L. performed the nitrogen sorption measurements. H.Y. and Y.C. performed the X-ray diffraction measurements. K.H. and Jun Li carried out theoretical calculations. X.H., S.M., J.P.-R., Jun Li and J. Lu co-wrote the manuscript. All authors discussed and commented on the manuscript.

**Competing interests**

The authors declare no competing interests.

**Additional information**

Supplementary information The online version contains supplementary material available at https://doi.org/10.1038/s41565-021-01022-y.

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Peer review information Nature Nanotechnology thanks Abhaya Datye and the other, anonymous, reviewer(s) for their contribution to the peer review of this work.

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