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Ultrafine NiMoO_x nanoparticles confined in mesoporous carbon for the reduction of nitroarenes: effect of the composition and accessibility of the active sites†

 Shuna Li, Yipin Lv, Guolong Song, Cuncheng Li,  Daowei Gao* and Guozhu Chen *

The design of ultrafine NiMoO_x nanoparticles (NPs) confined in hierarchically porous carbon remains a great challenge due to its high calcination temperature. In addition, the composition of active sites of NiMoO_x NPs for the hydrogenation reaction is still ambiguous. Herein, we report a general approach for the synthesis of ultrafine NiMoO_x NPs confined in mesoporous carbon with different morphologies and compositions using the replication method with SBA-15 as a hard template. The pore structure of mesoporous carbon and the Ni/Mo composition valence-state were discovered to be the main factors in the reduction of nitroarenes. The NiMoO_x/mesoporous carbon-platelet (NiMoO_x/MC-PL) with short mesochannels (~350 nm) and high surface area (~995 m² g⁻¹) possessed excellent catalytic activity towards the reduction of 4-nitrophenol, whereas NiMoO_x/mesoporous carbon-hexagonal-prism (NiMoO_x/MC-HP), NiMoO_x/mesoporous carbon-long-rod (NiMoO_x/MC-LR), and NiMoO_x/mesoporous carbon-spherical (NiMoO_x/MC-SP) with long mesochannels and relatively less surface area exhibited poor catalytic performance. The bifunctional mechanism or electronic synergistic effects of Ni and Mo species enhanced their catalytic performance. A good balance between MoO_x and metallic Ni (NiMoO_x/MC-PL-450) was found to be suitable for the reduction of 4-NP.

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1. Introduction

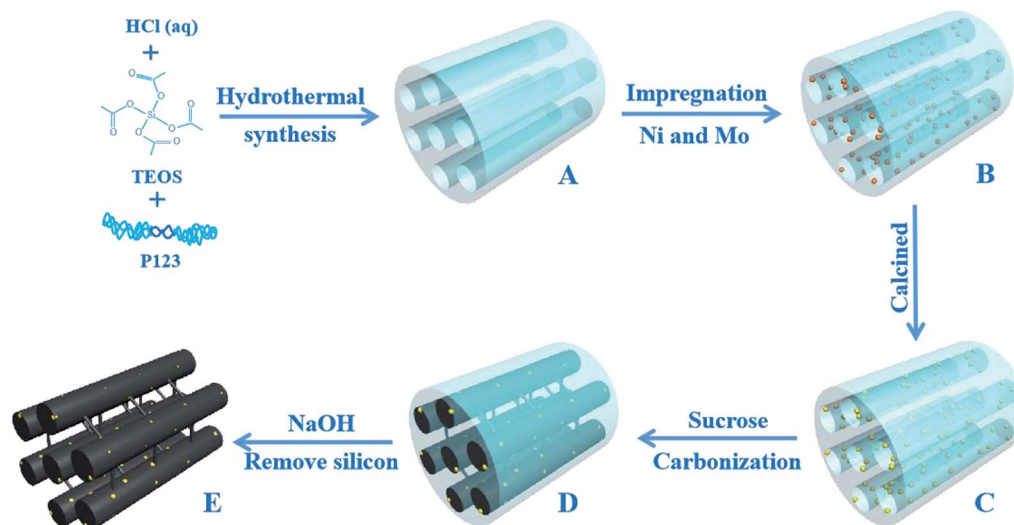
Nitroarenes are regarded as the most common pollutants from chemical, pharmaceutical, and industrial sources.^{1–4} 4-Nitrophenol (4-NP), one typical species of nitroarenes, is difficult to dispose of due to its high stability in water. At present, the most common methods for treating 4-NP include microbial degradation, electrochemical treatment, and catalytic reduction.^{5–7} Among them, the selective reduction of 4-NP to 4-aminophenol (4-AP) with NaBH₄ is of much interest because not only is this catalytic reaction green and effective, but also 4-AP is an important intermediate as a reduction product that can be widely applied in various fields such as dyes and pharmaceuticals.^{8,9} It is generally recognized that precious metals (*e.g.*, Au, Pt, and Pd) exhibit a superior performance for the reduction of 4-NP.^{10–12} However, the high cost and shortage of resources would inevitably become an obstacle to the wide application of precious metals. Therefore, transition metals with low costs and large storage capacities, especially Ni with a high selectivity and

activity, have attracted widespread attention.^{13,14} Compared to pure Ni, Ni-based catalysts exhibit a much higher catalytic performance owing to the so-called synergistic effect. For example, Zhao *et al.* reported that Ni/TiO₂-C composites possessed an excellent catalytic performance owing to the synergistic effect between Ni and TiO₂.¹⁵ Xu *et al.* synthesized Ni/CeO_{2-x}/Pd by the hydrothermal method. Due to their synergistic effect, Ni/CeO_{2-x}/Pd manifested a super catalytic activity and stability for the hydrogenation of styrene and 4-NP.¹⁶ The different components in the Ni-based catalysts also had a great influence on the catalytic properties. Lu *et al.* found that Ni-MoO₃/CN@SBA-15 showed an outstanding catalytic performance and chemoselectivity towards the reduction of various substituted nitroarenes compared to those of Ni/CN@SBA-15.¹⁷ Although great efforts have been made in the synthesis and catalysis of Ni-based catalysts, there are still a few issues that we need to address. First, further stabilization of Ni-based NPs and supports is not strong enough. Hence, aggregation and leaching of Ni-based NPs would result in the significant loss of catalytic activity.^{18,19} Second, the influence of the support morphology and components of the Ni-based catalysts on the catalytic performance still needs to be explored to fully understand the mechanism for the reduction of 4-NP.

School of Chemistry and Chemical Engineering, University of Jinan, Jinan 250022, PR China. E-mail: chm_gaodw@ujn.edu.cn; chm_chengz@ujn.edu.cn

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Scheme 1 Illustration of the synthesis of $\text{NiMoO}_x/\text{MC-LR}$. (A) Ordered mesoporous silica SBA-15-LR template. (B) A after the impregnation of Ni and Mo. (C) B after being calcined at $550\text{ }^\circ\text{C}$ for 4 h. (D) C after the impregnation of sucrose and carbonization. (E) Ordered mesoporous $\text{NiMoO}_x/\text{MC-LR}$.

To enhance the stability of Ni-based catalysts, various structures were designed to fix the Ni-based NPs such as the core-shell structure ($\text{Ni}/\text{nanorod-CeO}_2/\text{SiO}_2$ and $\text{NiCo}_2\text{O}_4/\text{MnO}_2$), encapsulation into molecular sieve pores ($\text{Ni-ZrO}_2/\text{MCM-41}$ and NiO-CuO/SBA-15), and carbon nanotubes ($\text{NiCo}_2\text{O}_4/\text{CNT}$ and $\text{Ni-CeO}_2/\text{CNT}$).^{20–25} It is well-known that the molecular sieve with a porous structure can not only prevent the aggregation and loss of NPs, but the channel can also facilitate the diffusion of substrate molecules, which is beneficial to catalytic reactions.²⁶ More importantly, with the development of nanotechnology, researchers could precisely control the dimensions of molecular sieves (*e.g.*, morphology, pore size, and pore length), which was absolutely helpful to study the kinetic process of catalytic reactions that used molecular sieves as catalyst supports. In addition to mesoporous molecular sieves, mesoporous carbon (MC) was also widely used in various catalyst supports because of its good alkali resistance and electrical conductivity.²⁷ However, mesoporous carbon often has difficulty in controlling the pore size distribution and pore uniformity. When the mesoporous carbon was derived from mesoporous molecular sieves that acted as templates, the as-prepared carbon inherited the unique structural characteristics of molecular sieves as well as combined efficiently the structural advantages of mesoporous molecular sieves and carbon.

Herein, we report a general synthesis procedure for the ultrafine NiMoO_x NPs confined in MC with a tunable pore structure and NiMo composition valence-state using SBA-15 as a hard template. First, the $\text{NiMoO}_x/\text{SBA-15}$ catalysts were synthesized by co-impregnating Ni and Mo precursors into the mesochannels of SBA-15 (Scheme 1A–C). Second, NiMoO_x/MC catalysts were prepared by the replication method using $\text{NiMoO}_x/\text{SBA-15}$ as a hard template (Scheme 1C–E). The ultrafine NiMoO_x NPs with diameters less than 2 nm were uniformly distributed on the mesochannel structure of the mesoporous

carbon and had a high stability attributed to the confinement of the mesochannel. The bifunctional mechanism or electronic synergistic effects of Ni and Mo species for the reduction of 4-NP and other nitroarenes with different substituent groups were systematically investigated. A possible mechanism for the reduction of nitroarenes catalyzed by the NiMoO_x bimetallic catalysts was also proposed.

2. Experimental

2.1 Materials

All chemicals were analytical grade reagents and used without further purification. Sucrose ($\text{C}_{12}\text{H}_{22}\text{O}_{11}$), sodium hydroxide (NaOH), sodium borohydride (NaBH_4), and ethanol ($\text{C}_2\text{H}_6\text{O}$) were obtained from Sinopharm Chemical Reagent Co. Ltd. Carbon disulfide (CS_2) and *p*-nitrophenol ($\text{C}_6\text{H}_5\text{NO}_3$) were purchased from Shanghai Macklin Biochemical Co. Ltd. Sulfuric acid (H_2SO_4) was purchased from Beijing Chemical Reagent Company.

2.2 Synthesis of NiMoO_x/MC catalysts with different morphologies

The mesoporous SBA-15 silica used as the template was synthesized according to the literature.²⁸ The $\text{NiMoO}_x/\text{SBA-15}$ catalysts with different morphologies were synthesized by the impregnation method. After impregnation, the samples were dried at $100\text{ }^\circ\text{C}$ for 4 h and then calcined at $550\text{ }^\circ\text{C}$ for 4 h. The procedure is shown in Scheme 1. For MC, a typical synthesis method was followed: 1.0 g of $\text{NiMoO}_x/\text{SBA-15}$ was dissolved in 5 mL of water containing 1.25 g of sucrose and 0.14 g of H_2SO_4 . The mixture was stirred for 1 h and dried at $100\text{ }^\circ\text{C}$ for 5 h. The mixture was then dried at $160\text{ }^\circ\text{C}$ for 5 h. The obtained powder was mixed with a solution that included 0.8 g of sucrose and 0.14 g of H_2SO_4 . The mixture was then heated at $100\text{ }^\circ\text{C}$ for 5 h



and dried at 160 °C for 5 h. The black powder was calcined under N₂ atmosphere at 800 °C for 2 h. The resulting composite was added into a 2 M NaOH solution at room temperature to remove the silica. After filtration and washing, all samples were dried in an oven. The obtained catalysts were denoted as NiMoO_x/MC-PL (NiMoO_x/mesoporous carbon-platelet), NiMoO_x/MC-SP (NiMoO_x/mesoporous carbon-spherical), NiMoO_x/MC-LR (NiMoO_x/mesoporous carbon-long-rod), and NiMoO_x/MC-HP (NiMoO_x/mesoporous carbon-hexagonal-prism). For comparison, catalysts with different compositions were prepared by the same method, and were designated as NiO/MC-PL, MoO₃/MC-PL, NiMoO_x, and MC-PL. For further comparison, catalysts with different Ni/Mo valence-states were prepared by hydrogen reduction at different temperatures (450 °C and 750 °C) and CS₂ vulcanization (referred to as NiMoO_x/MC-PL-450, NiMoO_x/MC-PL-750, and NiMoS/MC-PL, respectively).

2.3 Catalyst characterization

The X-ray diffraction (XRD) patterns of the catalysts were characterized with a powder X-ray diffractometer (Shimadzu XRD 6000). The surface morphologies of the catalysts were obtained

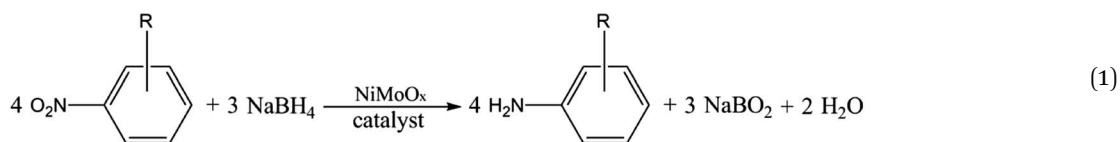
oxidation states of the chemical species were characterized using a X-ray photoelectron spectrometer (XPS) on the PHI5000 Versaprobe system. The contents of Ni and Mo were obtained from an inductively coupled plasma mass spectrometer (ICP-MS, Thermo Scientific XSeries-2).

The hydrogen temperature-programmed reduction (H₂-TPR) was performed on a PCA-1200 instrument. The NiMoO_x/MC-PL catalyst was pretreated under an Ar atmosphere (30 mL min⁻¹) at 100 °C for 30 min. The samples were then reduced with 7% H₂/Ar mixed gas (30 mL min⁻¹) from 25 °C to 700 °C.

2.4 Catalyst measurements

The catalytic reduction of 4-NP over catalysts was carried out at room temperature, which was analyzed by a UV-Visible spectroscopy instrument (TU-1901). Freshly prepared NaBH₄ solution (1.25 mL, 0.02 M), a certain amount of catalyst solution (1.25 mL, 0.4 mg mL⁻¹), and 4-NP (25 μL, 0.012 M) were mixed together. Subsequently, the absorbance of the solution in the range of 200 to 500 nm was measured using an UV-Vis absorption spectrometer. The corresponding reduction equations are as follows:

The conversion of 4-nitrophenol is expressed by



using an emission scanning electron microscope (SEM-SU8010). High-resolution transmission electron microscopy (HRTEM) and elemental mapping were performed with a Tecnai F20 instrument at 200 kV. The surface area and pore size were analyzed on a Micromeritics TriStar II 3020 analyzer. The

$$\text{Conversion(4-NP)} = \left(1 - \frac{A_t}{A_0}\right) \times 100\% \quad (2)$$

where A_0 and A_t are the absorbance of 4-NP at time 0 and t , respectively.

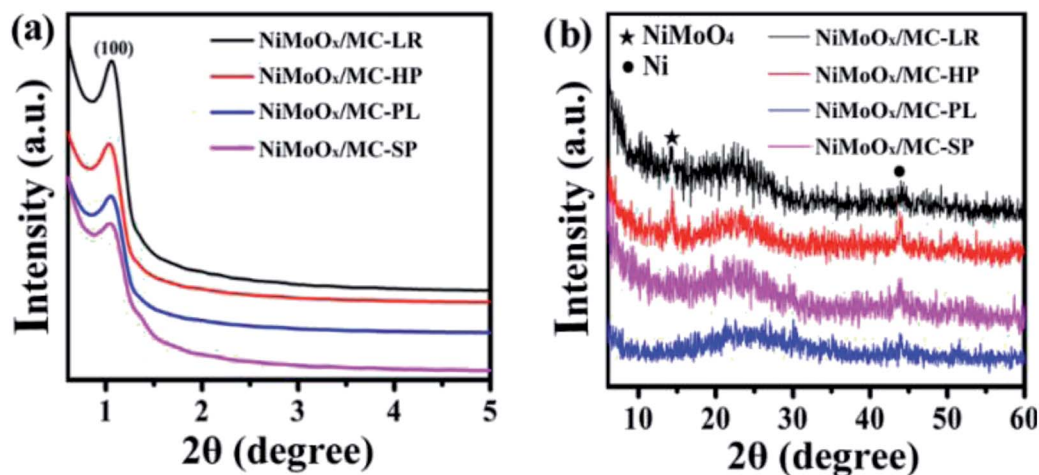


Fig. 1 XRD patterns for various morphologies of NiMoO_x/MC (a and b) in the small-angle and wide-angle.



$$k = \frac{-\ln(C_t/C_0)}{t} \quad (3)$$

$$K = -\frac{k}{m} \quad (4)$$

where k is the rate constant, C_0 and C_t are 4-NP concentrations at time 0 and t , respectively. Also, m represents the mass of the catalyst.

Typically, in a glass reactor, a solution of the catalyst (0.5 g L⁻¹, 1 mL), NaBH₄ (0.4 mol L⁻¹, 0.75 mL) and nitroarenes (1 mmol L⁻¹, 8 mL) were stirred. The reaction was then quenched by adding NH₄Cl solution and the reaction mixture was extracted with ethyl acetate. Subsequently, the mixture was dried using MgSO₄. The product obtained was analyzed using gas chromatography (GC-2010 Plus, capillary column, KB-1) and gas chromatography-mass spectrometry (GC-MS, Thermo-Finnigan Trace DSQ coupled with a HP-5MS column).

3. Results and discussion

3.1 Characterization of the as-synthesized catalysts

The XRD patterns of different morphologies of the catalysts are displayed in Fig. S1† and 1. Fig. S1(a)† and 1(a) show the small-angle XRD patterns of NiMoO_x/SBA-15 and the corresponding NiMoO_x/MC. All samples displayed a sharp diffraction peak below 1.1° and two other peaks between 1.1° and 2.0° that can be indexed as (100), (110) and (200) diffraction signals associated with an ordered hexagonal mesostructure. The unchanged ordered structure was a good indication that the NiMoO_x/MC material could be replicated from NiMoO_x/SBA-15. In addition, compared with the XRD patterns of NiMoO_x/SBA-15, the characteristic diffraction peaks attributed to the (100) in the XRD patterns of NiMoO_x/MC shifted to the wide-angle direction, thus indicating that the pore size of NiMoO_x/MC was smaller than that of NiMoO_x/SBA-15.²⁹ Fig. S1(b)† reveals the wide-angle XRD patterns of NiMoO_x/SBA-15 catalysts. There was one broad peak between 20° and 30° for all the catalysts, which could be indexed to amorphous silica. One peak centered at 27.5° was attributed to the (011) crystal plane of MoO₃ (JCPDS card 47-1320).^{30,31} Interestingly, the dispersion of MoO₃ was different with respect to the support morphology changes. Among the four samples, the (011) peak intensity of MoO₃ in NiMoO_x/SBA-15-PL was higher than those of the other catalysts, which indicated that MoO₃ had poor dispersibility in NiMoO_x/SBA-15-PL. The wide-angle XRD patterns for NiMoO_x/MC with different morphologies exhibited three main diffraction peaks, as illustrated in Fig. 1(b). One wide diffraction peak between 20° and 30° was attributed to carbon. The diffraction pattern at 14.3° originated from the NiMoO₄ phase (JCPDS card 33-0948), which was ascribed to (110). Another diffraction pattern at 44.5° originated from metallic Ni (JCPDS card 04-0850), which was assigned to (111) of metal Ni.^{32,33}

The surface morphology of the NiMoO_x/MC catalysts was investigated by SEM, as shown in Fig. 2. The length and width of NiMoO_x/MC-PL were ~1075 nm and ~350 nm, and those of the NiMoO_x/MC-LR catalyst were ~820 nm and ~230 nm,

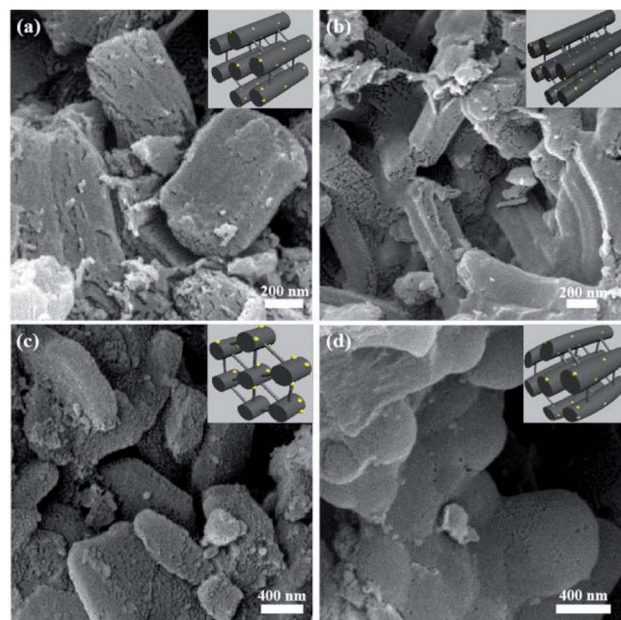


Fig. 2 SEM images of as-prepared NiMoO_x/MC catalysts with different morphologies. Note: (a) hexagonal prisms, (b) long rods, (c) platelets, and (d) sphericals.

respectively. Moreover, the length and width of NiMoO_x/MC-HP were ~755 nm and ~530 nm, respectively. For NiMoO_x/MC-SP, the particle diameter was measured to be ~745 nm. From Fig. 2, it can be seen that these four catalysts retained the original

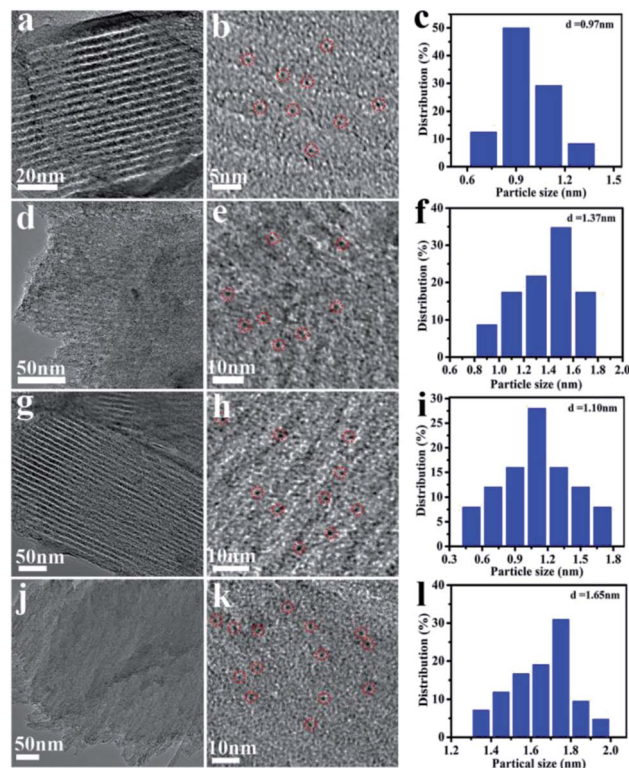


Fig. 3 TEM images and NiMoO_x particles size distribution histograms of NiMoO_x/MC. Note: (a–c) NiMoO_x/MC-HP, (d–f) NiMoO_x/MC-PL, (g–i) NiMoO_x/MC-LR and (j–l) NiMoO_x/MC-SP.



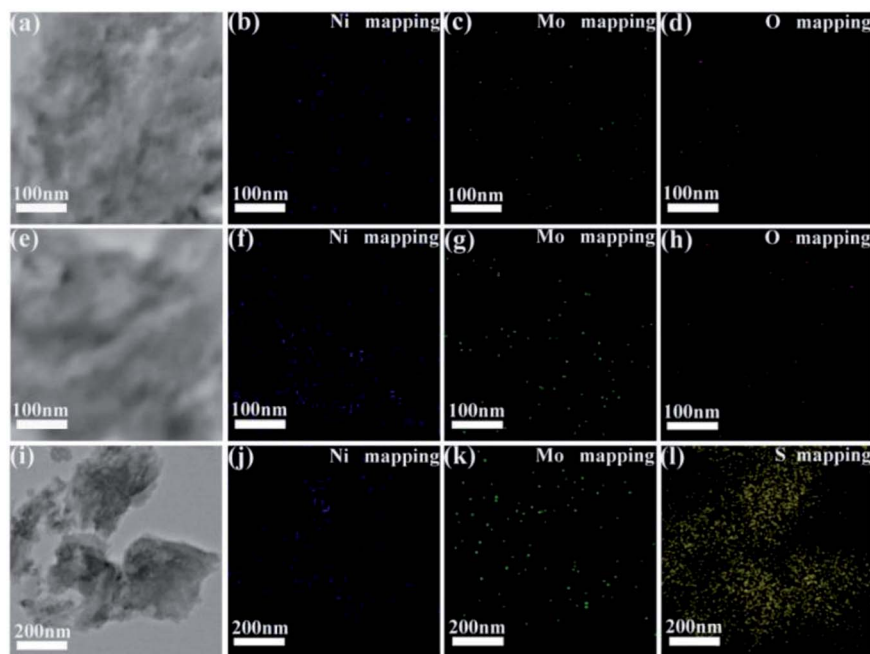


Fig. 4 HRTEM images and the corresponding EDS elemental mapping images of (a–d) NiMoO_x/MC-PL, (e–h) NiMoO_x/MC-PL-450 and (i–l) NiMoS/MC-PL.

morphology of their corresponding NiMoO_x/SBA-15, indicating that the high calcination temperature did not destroy the structures of NiMoO_x/SBA-15 during the replication process.

The TEM images and NiMoO_x NPs size distribution histograms of NiMoO_x/MC are shown in Fig. 3. From Fig. 3(a, d, g and j), it can be seen that all catalysts had a highly-ordered mesoporous structure, which was consistent with the small-angle XRD patterns. Importantly, ultrafine NPs were highly dispersed without any aggregation and well-encapsulated into the mesopores. As seen in the size distribution histograms, the average sizes of the NiMoO_x NPs of NiMoO_x/MC-HP, NiMoO_x/MC-LR, NiMoO_x/MC-PL, and NiMoO_x/MC-SP were ~ 0.97 , ~ 1.10 , ~ 1.37 and ~ 1.65 nm, respectively. Furthermore, the corresponding elemental mapping images of Ni, Mo, and O for

the NiMoO_x/MC-PL and NiMoO_x/MC-PL-450 catalysts are shown in Fig. 4, which reveal that the three elements were homogeneously distributed in the structure of the catalyst.³⁴ The molar ratio of Ni to Mo was about 2, which was obtained from the TEM-EDS of NiMoS/MC-PL (Fig. S2†).

The nitrogen adsorption–desorption isotherms, pore size distribution, and the corresponding parameters of the catalysts with different morphologies are displayed in Fig. 5 and Table 1. The nitrogen adsorption–desorption isotherms exhibited typical type IV patterns with hysteresis loops and capillary condensation, which indicated the presence of mesopores.^{35,36} The pore size distributions of NiMoO_x/MC-PL and NiMoO_x/MC-SP were about 4.0 nm, which indicated that the as-synthesized mesoporous materials had a uniform pore size distribution.³⁷

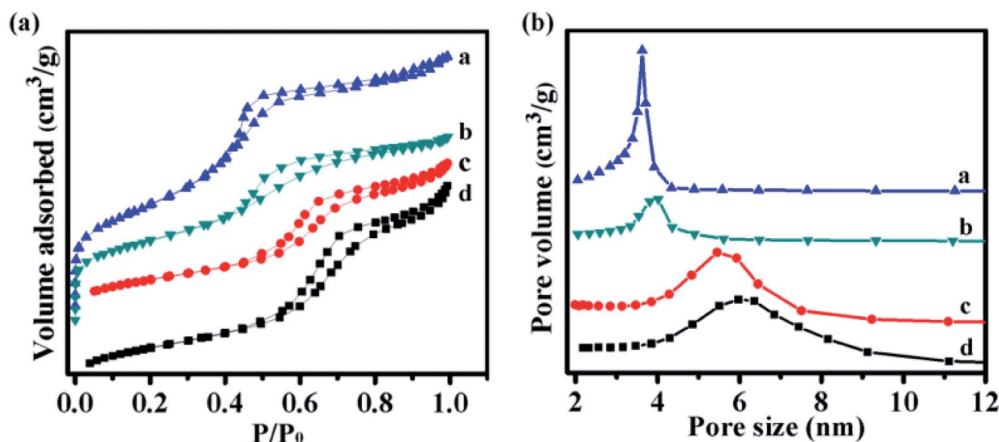


Fig. 5 The N₂ adsorption–desorption isotherms and pore size distribution of (a) NiMoO_x/MC-PL, (b) NiMoO_x/MC-SP, (c) NiMoO_x/MC-HP, and (d) NiMoO_x/MC-LR.



Table 1 The structural characteristics of mesoporous NiMoO_x/MC

| Samples | Dimensions (ca. nm) | S_{BET}^a (m ² g ⁻¹) | S_{mic}^b (m ² g ⁻¹) | V_{mic}^c (cm ³ g ⁻¹) | V_{mes}^c (cm ³ g ⁻¹) | d_{BJH}^d (nm) |
|---------------------------|---------------------|--|--|---|---|-------------------------|
| NiMoO _x /MC-PL | 1075 × 350 | 995 | 42 | 0.02 | 0.98 | 4.0 |
| NiMoO _x /MC-LR | 820 × 230 | 795 | 360 | 0.19 | 0.81 | 5.0 |
| NiMoO _x /MC-SP | 755 × 530 | 762 | 185 | 0.09 | 0.66 | 3.9 |
| NiMoO _x /MC-HP | 745 × 745 | 673 | 298 | 0.15 | 0.61 | 4.5 |

^a BET surface area. ^b Calculated using the *t*-plot micropore area. ^c Calculated using the *t*-plot micropore volume. ^d Pore diameter was calculated with the BJH method.

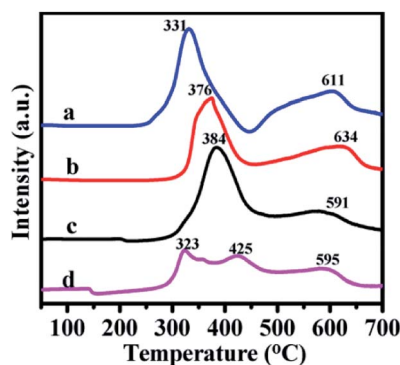


Fig. 6 H₂-TPR profiles of catalysts (a) NiMoO_x/MC-PL, (b) NiMoO_x/MC-LR, (c) NiMoO_x/MC-HP and (d) NiMoO_x/MC-SP.

As shown in Table 1, the surface area of NiMoO_x/MC-PL was ~995 m² g⁻¹, which was larger than those of NiMoO_x/MC-HP, NiMoO_x/MC-LR, and NiMoO_x/MC-SP. Therefore, the NiMoO_x/MC-PL catalyst provided a large surface area for the dispersion of the active sites. The average pore size of the NiMoO_x/MC catalyst followed the order: NiMoO_x/MC-LR (5.0 nm) > NiMoO_x/MC-HP (4.5 nm) > NiMoO_x/MC-PL (4.0 nm) > NiMoO_x/MC-SP (3.9 nm).

The H₂-TPR technique was used to study the reducibility of the Ni and Mo oxides on the support. The profiles are listed in Fig. 6. The profiles of the as-prepared catalysts exhibited two wide peaks in the 320–430 °C and 590–640 °C ranges.

The first reduction peak was derived from nickel oxide and a portion of the molybdenum oxide. The second peak was ascribed to the second step reduction of MoO_x. The different peak reduction temperatures were ascribed to the differences in the reducibility of the oxide. Compared to the Mo species, the Ni species was easier to reduce. According to the reference, the reduction peak for pure NiO (Ni²⁺ to Ni⁰) was centered around 350–400 °C³⁸ and the reduction peak for MoO₃ (Mo⁶⁺ to Mo⁴⁺) appeared at about 430 °C.³⁹ Compared to pure NiO, the reduction peak of the different catalysts for the Ni species migrated to a lower temperature. This result was attributed to the phenomenon of hydrogen overflow from molybdenum oxide to nickel, which promoted the reduction of nickel oxide.⁴⁰

XPS is a surface analytical technique that reveals information on the chemical state and surface composition of the catalysts. The XPS spectra and related parameters for NiMoO_x/MC-PL, NiMoO_x/MC-PL-450, NiMoO_x/MC-PL-750, and NiMoS/MC-PL are displayed in Fig. 7 and Table S1.† The Ni 2p spectra showed four main peaks, which could be assigned to Ni⁰, NiO, Ni(OH)₂, and the satellite signal.^{41,42} The ratio defined as Ni⁰/

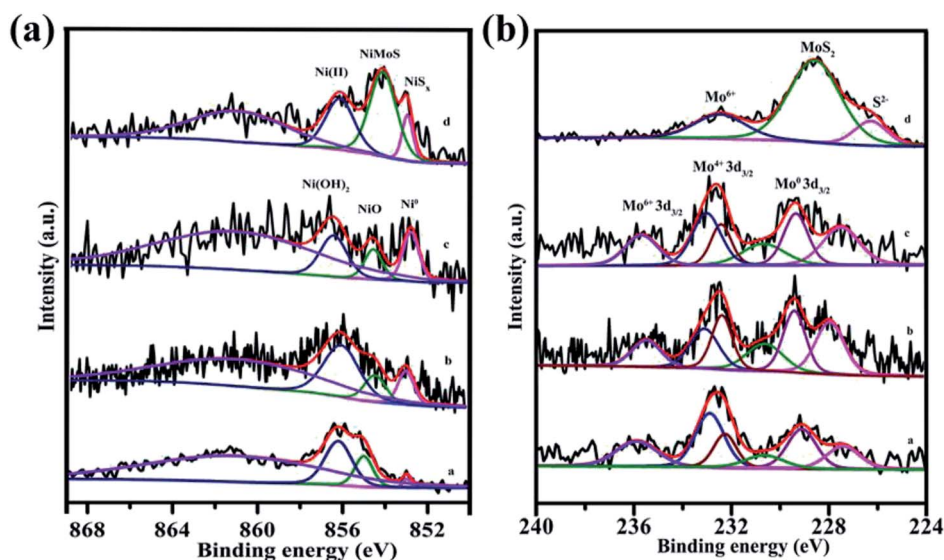


Fig. 7 XPS spectra of Ni 2p and Mo 3d for different catalysts. Note: (a) NiMoO_x/MC-PL, (b) NiMoO_x/MC-PL-450, (c) NiMoO_x/MC-PL-750, and (d) NiMoS/MC-PL.



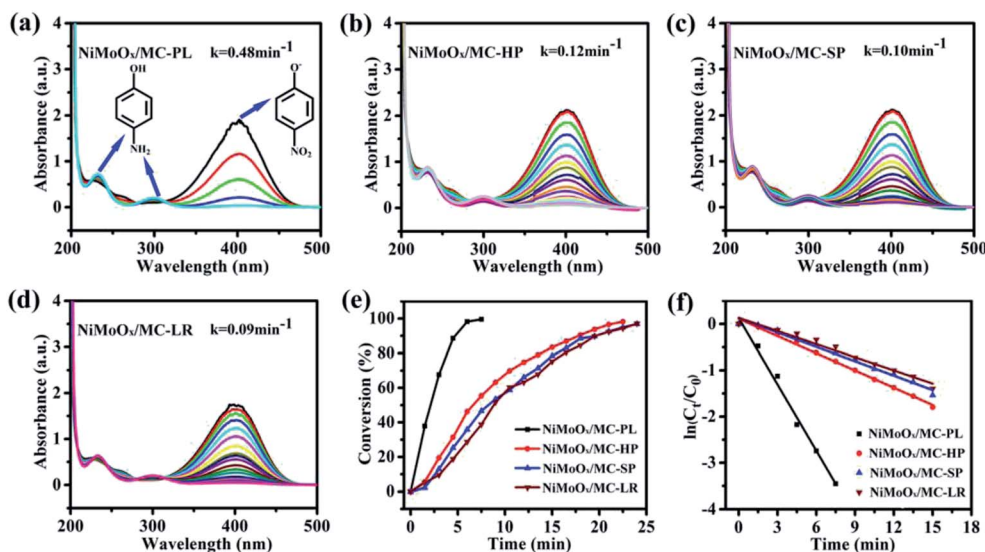


Fig. 8 The reduction of 4-NP recorded every 1.5 min over NiMoO_x/MC-PL (a), NiMoO_x/MC-HP (b), NiMoO_x/MC-SP (c), and NiMoO_x/MC-LR (d). The relationship between conversion and reaction time (e). Plots of $\ln(C_t/C_0)$ versus reaction time (f).

(Ni⁰+Ni²⁺) followed the order of NiMoO_x/MC-PL-750 > NiMoO_x/MC-PL-450 > NiMoO_x/MC-PL, which correlated well with the H₂-TPR spectra. The results showed that the content of metallic nickel and the active site increased with the increase in the reduction temperature. The Mo 3d XPS spectra for the different catalysts showed several pairs of corresponding peaks, which could be assigned to Mo⁶⁺, Mo⁴⁺, and metal Mo⁰. Compared to NiMoO_x/MC-PL, the binding energy of Mo⁴⁺ and Mo⁶⁺ in NiMoO_x/MC-PL-450 and NiMoO_x/MC-PL-750 shifted to lower values, respectively, indicating that more unsaturation sites were generated. From Table S1,[†] the percentage of MoO_x species, defined as (Mo⁶⁺ + Mo⁴⁺)/(Mo⁰ + Mo⁴⁺ + Mo⁶⁺), exhibited a significant change in different catalysts; it dropped from an initial of 88% to 63.7% with the increase in the reduction temperature. These results indicated that the molybdenum oxide on the surface of NiMoO_x/MC was not completely reduced, even when the temperature was relatively high. In the case of NiMoS/MC-PL, the XPS peaks of Ni and Mo demonstrated the formation of the NiMoS nanocrystal.

3.2 Catalytic activity towards the reduction of 4-nitrophenol

The catalytic performance of the catalysts was evaluated by the reduction of 4-nitrophenol (4-NP) in the presence of NaBH₄. The reduction process of 4-NP was monitored by measuring the UV-Vis absorbance of the solution. According to the reference, the aqueous solution of 4-NP showed a strong absorption band at 317 nm. The absorption band shifted to 400 nm, when NaBH₄ was added into the above solution, which indicated the formation of the nitrophenolate ion.⁴³ The catalytic properties of NiMoO_x/MC with different morphologies towards the reduction of 4-NP were comparatively studied, as displayed in Fig. 8. With the prolongation of the reaction time, the intensity of absorption peak at 400 nm decreased while another absorption band at 300 nm appeared.⁴⁴ After the completion of the reaction, the peak corresponding to the nitrophenolate ion was not observed, revealing that 4-NP was completely reduced. From Fig. 8(e), it can be observed that the reaction times for the conversion of 50% of 4-NP to 4-AP in the presence of NiMoO_x/MC-PL, NiMoO_x/MC-HP, NiMoO_x/MC-SP, and NiMoO_x/MC-LR

Table 2 The kinetic constants, turnover frequencies (TOF), and metal Ni loading with different catalysts

| Samples | k (min ⁻¹) | K (min ⁻¹ g ⁻¹) | Ni loading ^a (wt%) | TOF ^b × 10 ³ (s ⁻¹) |
|-------------------------------|--------------------------|--|-------------------------------|---|
| NiMoO _x /MC-PL | 0.48 | 960 | 4.05 | 175.8 |
| NiMoO _x /MC-HP | 0.12 | 240 | 3.2 | 69.3 |
| NiMoO _x /MC-SP | 0.10 | 200 | 2.78 | 64.7 |
| NiMoO _x /MC-LR | 0.09 | 180 | 3.37 | 49.9 |
| NiMoO _x /MC-PL-450 | 0.65 | 1300 | 4.05 | 241.5 |
| NiMoO _x /MC-PL-750 | 0.18 | 360 | 4.05 | 53.8 |
| NiMoS/MC-PL | 0.03 | 60 | 4.05 | — |
| NiO/MC-PL | 0.16 | 320 | 4.05 | 56.2 |
| MoO ₃ /MC-PL | 0.002 | 4 | — | — |
| NiMoO _x | 0.17 | 2073 | 24.7 | 103.8 |
| MC-PL | 0.001 | 2 | — | — |

^a Determined by ICP-MS. ^b Calculated on the basis of total catalyst and a conversion rate of 50%.



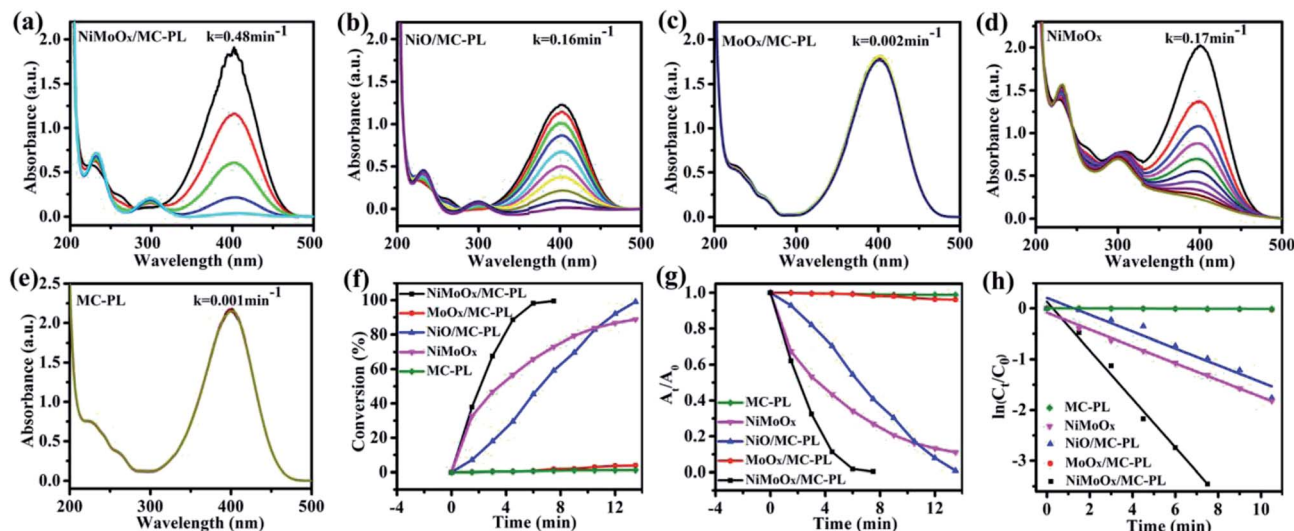


Fig. 9 Catalytic reduction of 4-NP for a time period of 1.5 min over NiMoO_x/MC-PL (a), NiO/MC-PL (b), MoO₃/MC-PL (c), NiMoO_x (d), and MC-PL (e). The conversion of 4-NP with reaction time over different catalysts (f). The relationship between A_t/A_0 and time (g). Plots of $\ln(C_t/C_0)$ versus time (h).

catalysts were 2.0 min, 6.6 min, 8.2 min, and 8.7 min, respectively. To qualify the reactivity, the reaction was regarded as a pseudo-first-order reaction. From Fig. 8(f), it can be observed that the reaction time is linear with respect to $\ln(C_t/C_0)$, which was consistent with first order kinetics. In order to compare the catalytic performances of different catalysts, the rate constants, k (min^{-1}) and K ($\text{min}^{-1} \text{g}^{-1}$), and the value of TOF (s^{-1}) of the catalysts were calculated. It can be found (Table 2) that the morphology of the catalysts had an important effect on the K and TOF values. Among the four different morphologies of the catalysts, the NiMoO_x/MC-PL catalyst exhibited the highest K ($960 \text{ min}^{-1} \text{g}^{-1}$) and TOF (0.176 s^{-1}) values. According to the HRTEM results, the size of the NiMoO_x NPs of the catalysts was similar. Therefore, the difference in the catalytic activities of

NiMoO_x/MC-PL, NiMoO_x/MC-HP, NiMoO_x/MC-LR, and NiMoO_x/MC-SP was mainly attributed to their different pore diameters, channel lengths, and surface areas. The NiMoO_x/MC-PL catalyst had short channels, which shortened the diffusion route for the reactants in the pores. This made the reactants to have easier contact with the active sites, thus favoring 4-NP reduction. In addition, the NiMoO_x/MC-PL catalyst exhibited a super catalytic performance owing to its large specific surface area ($\sim 995 \text{ m}^2 \text{g}^{-1}$) and short channel length ($\sim 351 \text{ nm}$). A larger specific surface area provided more attachment sites for NiMoO_x NP and then increased the active sites. However, NiMoO_x/MC-HP, NiMoO_x/MC-LR, and NiMoO_x/MC-SP catalysts had long channel lengths, which made 4-NP to

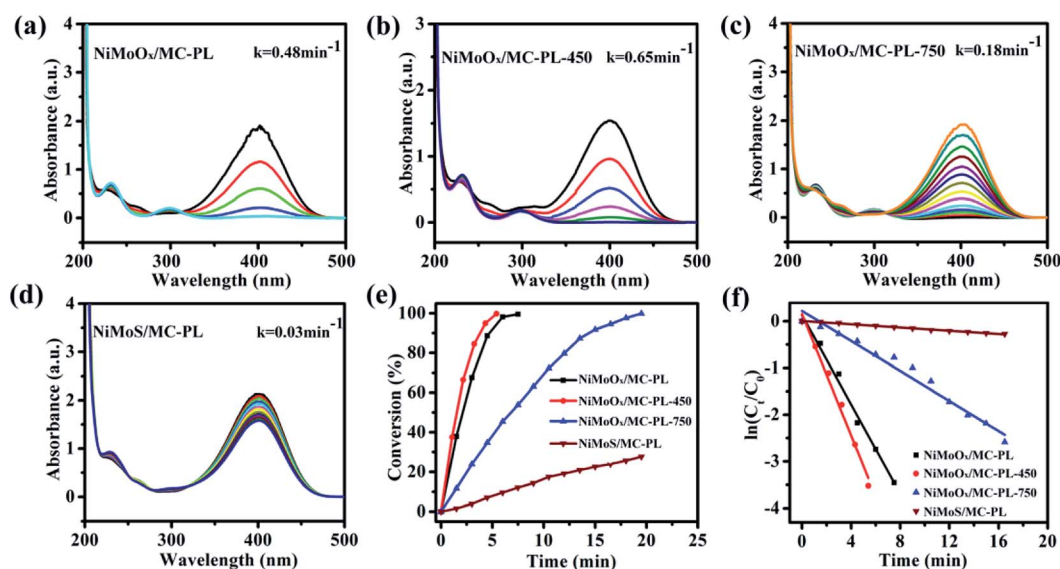


Fig. 10 Catalytic reduction of 4-NP over NiMoO_x/MC-PL (a), NiMoO_x/MC-PL-450 (b), NiMoO_x/MC-PL-750 (c), and NiMoS/MC-PL (d). Plots of the conversion of 4-NP versus time (e). The relationship of $\ln(C_t/C_0)$ and the reaction time (f).



Table 3 The reduction of nitroarenes over the NiMoO_x/MC-PL-450 catalyst^a

$$4 \text{ O}_2\text{N}-\text{C}_6\text{H}_4\text{R} + 3 \text{ NaBH}_4 \xrightarrow{\text{NiMoO}_x \text{ catalyst}} 4 \text{ H}_2\text{N}-\text{C}_6\text{H}_4\text{R} + 3 \text{ NaBO}_2 + 2 \text{ H}_2\text{O}$$

| Entry | Substrate | Product | Time (min) | Conv. (%) | <i>k</i> (min ⁻¹) | <i>K</i> (min ⁻¹ g ⁻¹) | TOF ^b × 10 ³ (s ⁻¹) |
|-------|-----------|---------|------------|-----------|-------------------------------|---|---|
| 1 | | | 10.0 | 97.2 | — | — | 37.6 |
| 2 | | | 40.0 | 68.1 | 0.039 | 78 | 6.6 |
| 3 | | | 10.0 | 97.5 | — | — | 37.7 |
| 4 | | | 20.0 | 88.4 | 0.128 | 256 | 17.1 |
| 5 | | | 10.0 | 99 | — | — | 38.3 |
| 6 | | | 40 | 99 | 0.039 | 78 | 9.6 |
| 7 | | | 40 | 61.5 | 0.018 | 36 | 6.0 |
| 8 | | | 10.0 | 66.5 | 0.287 | 574 | 25.7 |

^a Reaction conditions: catalyst (0.5 g L⁻¹, 1 mL), nitroarenes (1 mmol L⁻¹, 8 mL), NaBH₄ (0.4 mol L⁻¹, 0.75 mL), 600 rpm, room temperature. ^b The turnover frequencies (TOF) denote the moles of 4-NP converted per mole of Ni or Mo per second, calculated on the basis of total Ni and Mo loading in this research system.

diffuse for a longer period of time in contact with the active sites and therefore exhibited relatively inferior catalytic activities.

To further investigate the effect of composition on the reduction of 4-NP, the NiO/MC-PL, MoO₃/MC-PL, NiMoO_x and MC-PL catalysts were also prepared and compared with NiMoO_x/MC-PL. For NiMoO_x/MC-PL, NiO/MC-PL, and NiMoO_x, the reaction times for the conversion of 50% of 4-NP were 2.0, 6.5 and 3.5 min in (Fig. 9), respectively. However, 4-NP displayed almost no change over the MoO₃/MC-PL and MC-PL catalysts with the reaction time of 14.0 min, indicating that the sole Mo species and MC exhibited no activity for the 4-NP reduction. As shown in Table 2, the composition of the catalyst played an important effect on the values of *K* and TOF. The TOF values for these catalysts followed the order of NiMoO_x/MC-PL (0.176 s⁻¹) > NiMoO_x (0.104 s⁻¹) > NiO/MC-PL (0.056 s⁻¹) > MoO₃/MC-PL (0 s⁻¹) = MC-PL (0 s⁻¹). It was evident that the NiMoO_x/MC-PL

catalyst had the largest TOF, indicating that the Ni species in the NiMoO_x/MC was the main active site, while the Mo species contributed to the performance of the Ni catalyst. Compared with NiMoO_x/MC-PL, the NiMoO_x catalyst exhibited almost no catalytic activity. The enhanced catalytic properties of the NiMoO_x/MC-PL catalyst could be attributed to the synergistic effect of Mo and Ni species, as well as the strong interaction between the metal species and MC.

To further study the influence of valence-states of Ni and/or Mo on the catalytic performance, the NiMoO_x/MC-PL catalyst was reduced by hydrogen at different temperatures.

NiMoS/MC-PL was also prepared for comparison. It can be concluded from Fig. 10(e) that the conversion of 4-NP followed the order of NiMoO_x/MC-PL-450 > NiMoO_x/MC-PL > NiMoO_x/MC-PL-750 > NiMoS/MC-PL. From Table 2, it can be found that the valence-states of the metal also played a great role in the *K*



and TOF. The NiMoO_x/MC-PL-450 catalyst had the largest K (1300 min⁻¹ g⁻¹) and TOF (0.242 s⁻¹) values. Among these four catalysts, NiMoO_x/MC-PL-450 exhibited the highest TOF value, which was attributed to the excellent synergistic effect between Ni and Mo species. According to the results of H₂-TPR and XPS, after reduction at temperature 450 °C, most of the Ni species is converted to metallic nickel, while the MoO_x species is still present. The results demonstrate that the metallic Ni had more activity than NiO towards the reduction of 4-NP. Moreover, metallic Ni and MoO_x possessed the best synergistic effect for the reduction of 4-NP. With regard to the catalyst reduced at 750 °C, the proportion of Mo⁰ and Mo⁴⁺ species increased. However, the activity of NiMoO_x/MC-PL-750 was lower than that of NiMoO_x/MC-PL-450, clarifying that MoO₃ could contribute more to the reduction properties of Ni compared to metallic Mo and MoO₂. In addition, NiMoS/MC-PL showed a poor reaction rate, which indicated that the NiMoS species was less active than the corresponding metallic and oxidized NiMoO_x species.

3.3 The effect of reaction parameters on the reduction of 4-NP

It is well known that the reaction rate of a chemical reaction is related to the concentration of reactants and other factors. In this research, the amount of catalyst and 4-NP as well as the concentration of NaBH₄ were systematically studied. The concentration of NaBH₄ was varied, while all other parameters were kept constant (the concentration of 4-NP was 12 mM and the amount of catalyst was 0.5 mg). From Fig. S3 and Table S2,[†] it can be seen that the reaction rate increased as the concentration of NaBH₄ increased. Interestingly, the reaction rate increased rapidly as the concentration of NaBH₄ ranged from 0.005 to 0.02 M, whereas the rate increased slowly as the concentration of NaBH₄ ranged from 0.02 to 0.04 M. As we can see from the Table S2,[†] the variation tendency of TOF was similar to that of the rate constant as the concentration of NaBH₄ changed. It is concluded that the reduction rate and TOF were closely related to the concentration of NaBH₄.

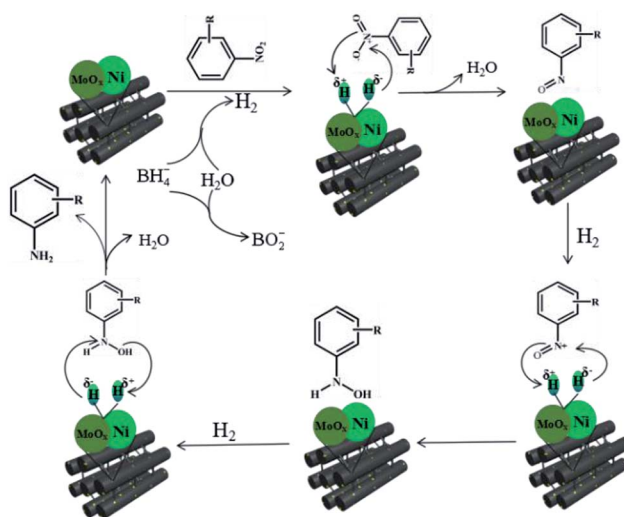
The effect of the amount of catalyst on the reduction of 4-NP is shown in Fig. S4 and Table S3.[†] It can be observed that k increased as the amount of catalyst increased. As seen in Fig. S4(h),[†] k was almost linear with respect to the amount of catalyst in the reaction system. This indicated that the increase in the catalyst amount resulted in more surface active sites. More chemical reactions then occurred, thus increasing the overall rate. The effect of the dosages of 4-NP on this reaction is shown in Fig. S5 and Table S4.[†] From Fig. S5(h),[†] it can be found that the reaction rate had an indirect relationship with the amount of 4-NP. The reaction rate decreased as the amount of 4-NP increased. This phenomenon can be ascribed to the competition between the reactants and free sites on the surface of NiMoO_x NPs. When reactants were adsorbed on the surface of the catalyst, the reaction occurred. In addition, adsorption of excessive reactants reduced the whole reaction rate.

3.4 The universal applicability of the NiMoO_x/MC-PL-450 catalyst towards the reduction of nitroarene

In order to demonstrate the universal applicability of the NiMoO_x catalyst in reducing the nitroarenes to aniline, various nitroarenes with different substituents were selected as the substrates.⁴⁵ NiMoO_x/MC-PL-450 was used as the catalyst and the results are listed in Table 3. NiMoO_x/MC-PL-450 exhibited an efficient catalytic activity to selectively reduce nitroarenes with different functionalities. Almost no by-products were detected. The nitroarenes were converted to the corresponding amines in the yield of >84.0% (Table 3, entries 1, 3, 4, 5 and 6), while 4-nitroanisole and 2-amino-4-nitrotoluene showed a relatively low conversion (less than 70%) under the same conditions. The NiMoO_x/MC-PL-450 catalyst demonstrated a high performance towards the reduction of nitroarenes under the optimized conditions. Therefore, the NiMoO_x/MC-PL-450 catalyst is of great significance for the reduction of nitroarenes with different functional groups.

3.5 The proposed mechanism towards the reduction of nitroarenes

According to the above experimental results, the NiMoO_x/MC-PL-450 catalyst exhibited a high performance, which was mainly ascribed to the synergistic effect between metal Ni and MoO_x species. A possible mechanism was proposed towards the reduction of nitroarenes in the presence of NaBH₄ over the NiMoO_x/MC-PL-450 catalyst in Scheme 2.⁴⁶ The typical steps are described below. First, the hydrolysis of NaBH₄ produced active hydrogen and electrons that were adsorbed on the surface of the NiMoO_x catalyst. Then, the active hydrogen was transferred from the surface of the NiMoO_x catalyst to nitroarenes, which were adsorbed on the catalyst surface. Second, the nitroso intermediates were formed after the loss of water molecules. The intermediates were then rapidly reduced to a stable hydroxylamine. Finally, the final product amine was obtained



Scheme 2 The mechanism towards the reduction of nitroarenes over the NiMoO_x/MC-PL-450 catalyst.



by dehydroxylation of hydroxylamine and then desorption from the catalyst, which was the rate-determining step.

4. Conclusions

In summary, the inexpensive NiMoO_x catalysts confined in the mesoporous carbon with different morphologies and compositions were prepared using SBA-15 as a hard template with the replication method. For the catalysts with different morphologies, NiMoO_x/MC-PL contributed to the dispersion of NiMoO_x NPs and the diffusion of 4-NP owing to its large specific surface area and shorter channel length, exhibiting outstanding activity. For the catalysts with different metal valence states, the NiMoO_x/MC-PL-450 catalyst showed the highest k (0.650 min⁻¹) and TOF (0.242 s⁻¹) values towards the reduction of 4-NP owing to the excellent synergistic effects of metallic Ni and MoO_x species. The as-prepared NiMoO_x/MC with efficient accessibility and suitable composition of the active sites illustrated a promising way for other reactions to design a novel non-noble catalyst.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- N. Sahiner, S. Yildiz and H. Al-Lohedan, *Appl. Catal., B*, 2015, **166–167**, 145–154.
- M. Zhu, C. Wang, D. Meng and G. Diao, *J. Mater. Chem. A*, 2013, **1**, 2118–2125.
- Z. Jiang, J. Xie, D. Jiang, X. Wei and M. Chen, *CrystEngComm*, 2013, **15**, 560–569.
- Y. Peng, W. Leng, B. Dong, R. Ge, H. Duan and Y. Gao, *Chin. J. Catal.*, 2015, **36**, 1117–1123.
- M. A. Oturan, J. Peiroten, P. Chartrin and A. J. Acher, *Environ. Sci. Technol.*, 2000, **34**, 3474–3479.
- H. Hu, J. H. Xin, H. Hu and X. Wang, *Nano Res.*, 2015, **8**, 3992–4006.
- W. Xiong, D. Sikdar, L. W. Yap, P. Guo, M. Premaratne, X. Li and W. Cheng, *Nano Res.*, 2016, **9**, 415–423.
- U. Kurtan, M. Amir and A. Baykal, *Chin. J. Catal.*, 2015, **36**, 705–711.
- C.-C. Yeh and D.-H. Chen, *Appl. Catal., B*, 2014, **150–151**, 298–304.
- J. Liang, X. Zhang, L. Jing and H. Yang, *Chin. J. Catal.*, 2017, **38**, 1252–1260.
- J. P. Xiao, Y. Xie, R. Tang, M. Chen and X. B. Tian, *Adv. Mater.*, 2001, **13**, 1887–1891.
- Q. Wang, W. Jia, B. Liu, A. Dong, X. Gong, C. Li, P. Jing, G. Xu and J. Zhang, *J. Mater. Chem. A*, 2013, **1**, 12732–12741.
- S. Jian and Y. Li, *Chin. J. Catal.*, 2016, **37**, 91–97.
- W. She, T. Qi, M. Cui, P. Yan, S. W. Ng, W. Li and M. Li, *ACS Appl. Mater. Interfaces*, 2018, **10**, 14698–14707.
- W. Gao, W. Li, Z. Xue, M. Pal, Y. Liu, C. Wang, J. Wang, S. Wang, X. Wan, Y. Liu and D. Zhao, *New J. Chem.*, 2016, **40**, 4200–4205.
- Y.-F. Jiang, C.-Z. Yuan, X. Xie, X. Zhou, N. Jiang, X. Wang, M. Imran and A.-W. Xu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 9756–9762.
- H. Huang, X. Wang, X. Li, C. Chen, X. Zou, W. Ding and X. Lu, *Green Chem.*, 2017, **19**, 809–815.
- J. Fang, Y. Zhang, Y. Zhou, S. Zhao, C. Zhang, C. Yang, W. Chen, M. Huang and Y. Gao, *Carbon*, 2017, **121**, 602–611.
- J. Zhang, G. Chen, M. Chaker, F. Rosei and D. Ma, *Appl. Catal., B*, 2013, **132–133**, 107–115.
- S. Zhu, X. Lian, T. Fan, Z. Chen, Y. Dong, W. Weng, X. Yi and W. Fang, *Nanoscale*, 2018, **10**, 14031–14038.
- K. Xu, W. Li, Q. Liu, B. Li, X. Liu, L. An, Z. Chen, R. Zou and J. Hu, *J. Mater. Chem. A*, 2014, **2**, 4795–4802.
- D. Liu, X. Y. Quek, W. N. E. Cheo, R. Lau, A. Borgna and Y. Yang, *J. Catal.*, 2009, **266**, 380–390.
- A. Ungureanu, B. Dragoi, A. Chiriac, C. Ciotonea, S. Royer, D. Duprez, A. S. Mamede and E. Dumitriu, *ACS Appl. Mater. Interfaces*, 2013, **5**, 3010–3025.
- H. Cheng, Y.-Z. Su, P.-Y. Kuang, G.-F. Chen and Z.-Q. Liu, *J. Mater. Chem. A*, 2015, **3**, 19314–19321.
- Z. Weng, W. Liu, L.-C. Yin, R. Fang, M. Li, E. Altman, Q. Fan, F. Li, H.-M. Cheng and H. Wang, *Nano Lett.*, 2015, **15**, 7704–7710.
- M. J. Mostazo-López, R. Ruiz-Rosas, A. Castro-Muñiz, H. Nishihara, T. Kyotani, E. Morallón and D. Cazorla-Amorós, *Carbon*, 2018, **129**, 510–519.
- X. Duan, M. Xiao, S. Liang, Z. Zhang, Y. Zeng, J. Xi and S. Wang, *Carbon*, 2017, **119**, 326–331.
- D. Gao, A. Duan, X. Zhang, Z. Zhao, E. Hong, J. Li and H. Wang, *Appl. Catal., B*, 2015, **165**, 269–284.
- Z. Zhao, Y. Dai, J. Lin and G. Wang, *Chem. Mater.*, 2014, **26**, 3151–3161.
- S. Badoga, K. C. Mouli, K. K. Soni, A. K. Dalai and J. Adjaye, *Appl. Catal., B*, 2012, **125**, 67–84.
- S. Wang, J. Wang, M. Zhu, X. Bao, B. Xiao, D. Su, H. Li and Y. Wang, *J. Am. Chem. Soc.*, 2015, **137**, 15753–15759.
- H. Jung, B. Kim, S. Kim, H. Jang, N.-S. Lee, W. Kim and M. H. Kim, *Mater. Lett.*, 2017, **204**, 173–176.
- W. Lu, Y. Song, M. Dou, J. Ji and F. Wang, *Chem. Commun.*, 2018, **54**, 646–649.
- Y. Sun, I. Sinev, W. Ju, A. Bergmann, S. Dresch, S. Köhl, C. Spöri, H. Schmies, H. Wang, D. Bernsmeier, B. Paul, R. Schmack, R. Kraehnert, B. R. Cuenya and P. Strasser, *ACS Catal.*, 2018, **8**, 2844–2856.
- D. Saikia, Y.-Y. Huang, C.-E. Wu and H.-M. Kao, *RSC Adv.*, 2016, **6**, 35167–35176.



- 36 Q. Wang, Z. Wang, T. Zheng, X. Zhou, W. Chen, D. Ma, Y. Yang and S. Huang, *Nano Res.*, 2016, **8**, 2294–2302.
- 37 Y. Chi, L. Zhao, Q. Yuan, X. Yan, Y. Li, N. Li and X. Li, *J. Mater. Chem.*, 2012, **27**, 13571–13577.
- 38 A. Alsalmeh, N. Alzaqri, A. Alsaleh, M. R. H. Siddiqui, A. Alotaibi, E. F. Kozhevnikova and I. V. Kozhevnikov, *Appl. Catal., B*, 2016, **182**, 102–108.
- 39 Q. Bkour, K. Zhao, L. Scudiero, D. J. Han, C. W. Yoon, O. G. Marin-Flores, M. G. Norton and S. Ha, *Appl. Catal., B*, 2017, **212**, 97–105.
- 40 N. Chen, S. Gong and E. W. Qian, *Appl. Catal., B*, 2015, **174**–**175**, 253–263.
- 41 Y. Sun, C. Li and A. Zhang, *Appl. Catal., A*, 2016, **522**, 180–187.
- 42 C. Chen, D. Yan, X. Luo, W. Gao, G. Huang, Z. Han, Y. Zeng and Z. Zhu, *ACS Appl. Mater. Interfaces*, 2018, **10**, 4662–4671.
- 43 A. Gangula, R. Podila, R. M. L. Karanam, C. Janardhana and A. M. Rao, *Langmuir*, 2011, **27**, 15268–15274.
- 44 C.-H. Liu, J. Liu, Y.-Y. Zhou, X.-L. Cai, Y. Lu, X. Gao and S.-D. Wang, *Carbon*, 2015, **94**, 295–300.
- 45 J. Hu, Y. Ding, H. Zhang, P. Wu and X. Li, *RSC Adv.*, 2016, **6**, 3235–3242.
- 46 R. Nie, J. Wang, L. Wang, Y. Qin, P. Chen and Z. Hou, *Carbon*, 2012, **50**, 586–596.

