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## Boosting the activity of PdAg alloy nanoparticles during H<sub>2</sub> production from formic acid induced by CrO<sub>x</sub> as an inorganic interface modifier†

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Interfacial modification of PdAg nanoparticles supported on mesoporous carbon (MSC) functionalized with weakly basic phenylamine groups was performed using an amorphous CrO<sub>x</sub> phase. The resulting PdAgCr/amine-MSC catalyst was found to promote the efficient dehydrogenation of formic acid (HCOOH) serving as a liquid organic hydrogen carrier. A maximum turnover frequency of 6898 h<sup>-1</sup> (based on the mass of Pd employed) was achieved, which was 1.6 times larger than the value previously obtained using a PdAg catalyst. Physicochemical characterization and density functional theory calculations indicated that electronic interactions with the CrO<sub>x</sub> phase induced a significant electronic gap. This effect, in turn, generated unique ensemble sites on the PdAg nanoparticle surfaces at which electron-deficient Ag<sup>δ+</sup> and electron-rich Pd<sup>δ-</sup> atoms were adjacent. Kinetic analyses and theoretical investigations demonstrated that O–H bond dissociation was assisted by amine groups on the support surface. Enhanced C–H bond dissociation and H<sub>2</sub> desorption resulting from the cooperative action at the ensemble sites were found to play a crucial role in increasing the catalytic activity. The present system is easily synthesized, does not require additives, can be recycled and effectively suppresses CO formation, meaning that this material is a suitable candidate for practical applications in fuel cells.

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### Broader context

Formic acid (FA, HCOOH) has emerged as one of the most promising hydrogen storage compounds because it is non-flammable, relatively nontoxic, and contains 4.4 wt% (53.4 g L<sup>-1</sup>) hydrogen. There have been extensive studies in developing catalysts targeted for efficient H<sub>2</sub> delivery. Our own group has demonstrated that the dispersed PdAg nanoparticles supported on phenylamine-functionalized mesoporous carbon display an extremely high turnover frequency (TOF) of 5638 h<sup>-1</sup> during the dehydrogenation of FA, and this work has provided advanced insights into the architecture of catalytically-active sites for this target reaction (*ACS Catal.*, 2018, 8, 2277). In this study, interfacial modification of PdAg nanoparticles was performed using highly dispersed CrO<sub>x</sub>. The resulting ternary PdAgCr/amine-MSC is proven to display a maximum TOF of 6898 h<sup>-1</sup>, which is 1.6 times larger than that with our previously reported binary catalyst. On the basis of kinetic and theoretical investigations, we demonstrated that the significant enhancement of activity is attributed to the interplay of unique ensemble sites on the PdAg nanoparticle surface, where electron-deficient Ag<sup>δ+</sup> and electron-rich Pd<sup>δ-</sup> are adjacent. We are sure that these findings are of sufficiently immediate interest to a general chemistry research readership.

## 1. Introduction

Hydrogen (H<sub>2</sub>) has a high gravimetric energy density of 33.3 kW h kg<sup>-1</sup>, which is three times the value for gasoline, and produces only water as a by-product when converted into energy in an internal combustion engine or a fuel cell.<sup>1,2</sup> Hence,

H<sub>2</sub> is a potentially important energy source. However, the development of safe and efficient methodologies for the storage and transportation of hydrogen gas is required to realize a sustainable hydrogen-based economy.<sup>3,4</sup> Conventional storage technologies, including pressurization and/or cryogenic liquefaction, suffer from a tradeoff relationship between storage density and efficiency.<sup>5</sup> As such, chemical hydrogen storage in liquid organic hydrogen carriers (LOHCs) may be a viable alternative to physical hydrogen storage.<sup>6–8</sup> In an LOHC system, hydrogen is stored by converting a hydrogen-lean compound into its hydrogen-rich counterpart *via* a catalytic hydrogenation reaction, while hydrogen is released by dehydrogenation of the hydrogen-rich compound. There are several LOHC systems,

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including cyclohexane–benzene, methylcyclohexane–toluene, decalin–naphthalene and perhydro-*N*-ethylcarbazole–*N*-ethylcarbazole, that have been recognized as suitable candidates.<sup>9–13</sup>

Formic acid (FA, HCOOH) has emerged as a prominent LOHC component over the last decade. FA is of interest because of its high hydrogen content (4.4 wt%; 53.4 g L<sup>-1</sup>), low toxicity and low flammability (flashpoint of 69 °C, which is much higher than the values for methanol (12 °C) and gasoline (-40 °C)) under ambient conditions.<sup>14,15</sup> The use of FA could allow economical CO<sub>2</sub>-mediated hydrogen storage energy cycling based on the regeneration of FA through the hydrogenation of CO<sub>2</sub> with H<sub>2</sub>.<sup>16</sup> The selective dehydrogenation of FA is favored thermodynamically ( $\Delta G = -48.4$  kJ mol<sup>-1</sup>), which is necessary to ensure that pure H<sub>2</sub> is generated without the formation of CO and H<sub>2</sub>O *via* the competitive dehydration pathway ( $\Delta G = -28.5$  kJ mol<sup>-1</sup>). This factor is especially important because the CO byproduct is toxic to Pt-based fuel-cell catalysts. For all these reasons, numerous investigations have been conducted in attempts to identify catalysts, both homogeneous and heterogeneous, capable of promoting FA dehydrogenation under mild conditions.<sup>17–23</sup>

To ensure that practical systems are developed, current research has focused on the use of reliable heterogeneous catalysts such as Au, Pt and especially Pd, which exhibits excellent catalytic activity. Alloying these elements with additional metals that possess superior resistance to CO poisoning is another promising strategy.<sup>24–28</sup> Our group previously developed binary nanoparticle (NP) catalysts based on Pd combined with coinage metals such as Au, Ag or Cu that exhibited superior activity compared with monometallic catalysts.<sup>29–31</sup> The synergistic effect provided by these materials is attributed to the electronic activation of Pd species by charge transfer resulting from the different work functions of the two metals. This prior work suggested possible design strategies for producing catalytically active sites promoting the dehydrogenation of FA.

Research regarding the nano-engineering of catalysts for efficient dehydrogenation of FA has indicated that precise tuning of the NP architecture at the angstrom level is required to provide the appropriate activity and selectivity. Controlling particle size can provide large surface-area-to-volume ratios and also optimize the relative proportions of low- and high-coordination surface atoms within regular arrangements of the NPs.<sup>32,33</sup> The catalytic behavior of NPs is also affected by morphology due to differences in the preferential exposure of particular crystallographic planes.<sup>34</sup> Many supports, including zeolites, silica, carbon and metal–organic frameworks, have been investigated in attempts to obtain strong metal–support interactions, with frequent reports of unique synergistic effects at the interfaces between NPs and supports.<sup>35–41</sup> Surface modification with organic molecules can also inhibit the agglomeration of NPs, thus allowing control over the adsorption kinetics of reactants or intermediates through specific geometric phenomena or metal–ligand electronic effects.<sup>42,43</sup>

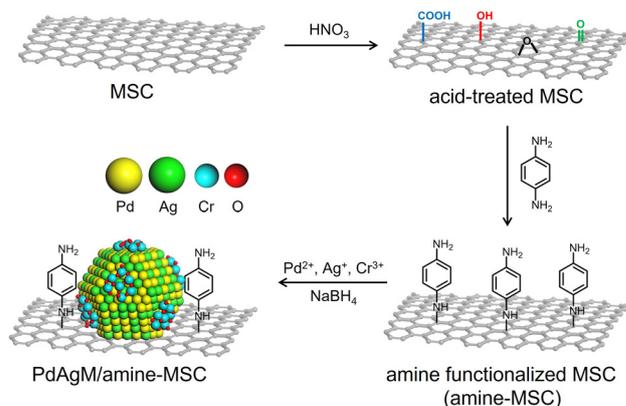
We have previously reported that the PdCuCr ternary NPs within a macroreticular basic resin that possesses -N(CH<sub>3</sub>)<sub>2</sub> functional groups are responsible for the efficient production

of H<sub>2</sub> from formic acid, which is attributed to the stabilization effect of surface Cr clusters and a synergistic effect to boost the C–H bond dissociation step.<sup>44</sup> Inspired by our previously obtained insight, the present study utilized a highly dispersed amorphous CrO<sub>x</sub> phase as an inorganic interface modifier to alter the electronic state of PdAg NPs supported on mesoporous carbon (MSC) functionalized with a weakly basic phenylamine. Unlike organic ligands, which exhibit Lewis basicity, metal centers connected with oxide (O<sub>2</sub><sup>-</sup>) groups in CrO<sub>x</sub> are positively charged, while those in the periphery of the Cr<sup>3+</sup> atoms are highly negatively charged. Thus, the modification of these materials with CrO<sub>x</sub> improves catalytic activity during dehydrogenation of FA compared with a pure binary PdAg catalyst. These electronic interactions generate unique ensemble sites at which electron-deficient Ag<sup>δ+</sup> and electron-rich Pd<sup>δ-</sup> atoms are adjacent, and ultimately lead to a rate-determining step involving the dissociation of C–H bonds in FA, as demonstrated by kinetics analyses and density functional theory (DFT) calculations.

## 2. Results and discussion

### 2.1. Characterization of catalysts

The synthetic procedure employed to produce catalysts comprising PdAgCr supported on phenylamine-functionalized MSC (denoted herein as PdAgCr/amine-MSC) is illustrated in Scheme 1. In this procedure, the MSC was treated with an aqueous nitric acid solution, which resulted in the formation of surface O-based functional groups such as -COOH and -OH (Fig. S1A, ESI<sup>†</sup>). The surface of the acid-treated MSC was subsequently functionalized with *p*-phenylenediamine to afford amine-MSC. During this step, the surface O-based functional groups reacted with the amine groups to generate amide and C–N bonds. The concentration of amine groups grafted onto the MSC was determined to be approximately 0.57 mmol g<sup>-1</sup> based on CHN elemental analysis (Table S1, ESI<sup>†</sup>), in good agreement with the value of 0.53 mmol g<sup>-1</sup> obtained from thermogravimetric analysis (Fig. S1B, ESI<sup>†</sup>). Pd, Ag and Cr were deposited by mixing the amine-MSC support with aqueous



Scheme 1 Schematic illustration of the synthesis of the PdAgCr-supported amine-MSC catalyst.



solutions containing  $\text{Pd}(\text{NO}_3)_2$ ,  $\text{AgNO}_3$  and  $\text{Cr}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ . Finally, each sample was reduced using  $\text{NaBH}_4$ , affording PdAgCr/amine-MSc.

The  $\text{N}_2$  adsorption–desorption isotherm for each specimen displayed a characteristic type IV curve with sharp capillary condensation steps and an H1-type hysteresis loop over the relative pressure ( $p/p_0$ ) range of 0.6–0.9, indicating the presence of ordered and well-preserved cylindrical mesopores (Fig. S2, ESI†). These results confirmed that the original pore structure of the MSC was maintained despite the addition of the modifier and metals. The Brunauer–Emmett–Teller surface areas ( $S_{\text{BET}}$ ) and pore volumes ( $V_p$ ) calculated from the  $\text{N}_2$  adsorption–desorption isotherms are summarized in Table S2 (ESI†). After amine functionalization to generate amine-MSc ( $S_{\text{BET}} = 153 \text{ m}^2 \text{ g}^{-1}$ ,  $V_p = 0.39 \text{ cm}^3 \text{ g}^{-1}$ ), both  $S_{\text{BET}}$  and  $V_p$  decreased compared with the values for the original MSC ( $S_{\text{BET}} = 320 \text{ m}^2 \text{ g}^{-1}$ ,  $V_p = 0.60 \text{ cm}^3 \text{ g}^{-1}$ ). A further slight decrease was observed after deposition of the metals ( $S_{\text{BET}} = 104 \text{ m}^2 \text{ g}^{-1}$ ,  $V_p = 0.24 \text{ cm}^3 \text{ g}^{-1}$ ).

Fig. 1 presents a high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) image of PdAgCr/amine-MSc, together with energy dispersive X-ray spectroscopy (EDX) maps. The NPs had a mean diameter ( $d_{\text{ave}}$ ) of 1.5 nm and were well dispersed (Fig. 1a). This value was comparable with that obtained for PdAg/amine-MSc ( $d_{\text{ave}} = 1.2 \text{ nm}$ ) and Pd/amine-MSc ( $d_{\text{ave}} = 1.8 \text{ nm}$ ) specimens, as shown in Fig. S3 and S4 (ESI†). However, a comparison of the particle sizes clearly demonstrated that the addition of the second and third metals resulted in the formation of smaller NPs. Energy-dispersive X-ray spectroscopy maps indicated that the bright particles were composed of Pd, Ag and Cr atoms that are highly dispersed throughout the amine-functionalized material (Fig. 1b–e).

The electronic state of each metal was investigated by X-ray photoelectron spectroscopy (XPS). In the case of Pd/amine-MSc, the Pd 3d<sub>5/2</sub> and 3d<sub>3/2</sub> electron binding energy values were 335.9 and 338.0 eV, and 341.0 and 343.3 eV, respectively. The peaks at lower binding energies were assigned to the Pd<sup>0</sup> state, while the higher ones were attributed to Pd<sup>2+</sup> (Fig. 2A). The presence of Pd<sup>2+</sup> was presumably due to the oxidation of the surface Pd NPs by  $\text{CrO}_x$  or strong interaction with surface amine groups. The Pd 3d peaks generated by PdAg/amine-MSc

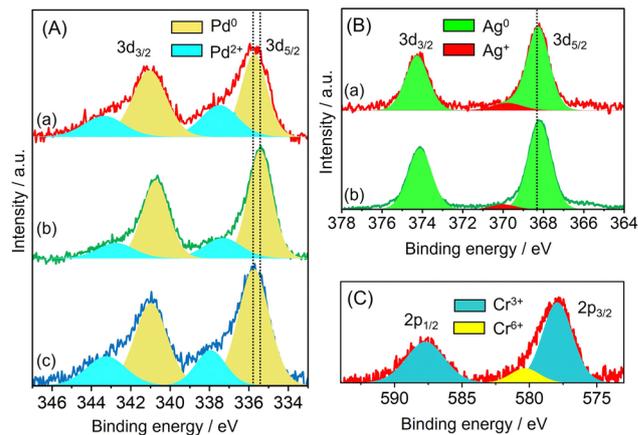


Fig. 2 (A) Pd 3d XPS spectra of (a) PdAgCr/amine-MSc, (b) PdAg/amine-MSc and (c) Pd/amine-MSc; (B) Ag 3d XPS spectra of (a) PdAgCr/amine-MSc and (b) PdAg/amine-MSc; and (C) Cr 2p XPS spectrum of PdAgCr/amine-MSc.

were shifted to lower binding energies by 0.4 eV compared with those obtained from the corresponding monometallic Pd catalyst. This shift was attributed to the ability of Ag atoms to donate electrons to Pd atoms upon alloying, as a consequence of the net difference in the ionization potentials of the two elements (Ag: 7.57 eV, Pd: 8.34 eV). In contrast, the addition of Cr atoms shifted the Pd 3d peaks toward higher binding energies compared with PdAg/amine-MSc. No differences were observed in the Ag 3d XPS spectra acquired from PdAg/amine-MSc and PdAgCr/amine-MSc (Fig. 2B), suggesting that the presence of Ag<sup>+</sup> was minimal. The Cr 2p spectra could be deconvoluted into two components (Fig. 2C), comprising a main peak at approximately 577.5 eV assigned to Cr<sup>3+</sup> and a small peak at 580 eV attributed to Cr<sup>6+</sup>. These results indicate that the Ag atoms were preferentially located in the core regions of the NPs, while Pd atoms were typically situated on NP surfaces and were partially oxidized by the amorphous chromium oxide ( $\text{CrO}_x$ ) and/or chromium hydroxide ( $\text{Cr}(\text{OH})_3$ ).

X-ray absorption analyses were also performed to elucidate the local structures of the specimens. The shapes of the Pd K-edge X-ray absorption near-edge structure (XANES) spectra of the three Pd-based samples all differed from that of Pd foil but resembled the spectrum of PdO (Fig. 3A). The Fourier-transform extended X-ray absorption fine structure (FT-EXAFS) spectra of all materials exhibited two peaks associated with Pd–O(N) and contiguous metallic Pd–Pd bonds, appearing at 1.6 and 2.6 Å, respectively (Fig. 3B), with no significant shift from the Pd–Pd distance in Pd foil. The first peak, at 1.6 Å, is believed to be partly due to Pd–N bonds originating from interactions with the amine-MSc support. The inverse FT of PdAgCr/amine-MSc was well fitted using Pd–O(N) and Pd–Pd shells with a coordination number (CN) and an interatomic distance ( $R$ ) of CN = 4.7 and  $R = 2.00 \text{ Å}$  and CN = 6.8 and  $R = 2.74 \text{ Å}$ , respectively (Table 1 and Fig. S5A, ESI†). Assuming that the contribution of Pd–N for all samples is almost the same since the mean diameter for all samples is comparable, the ratio of the contribution of Pd–O bonds to that

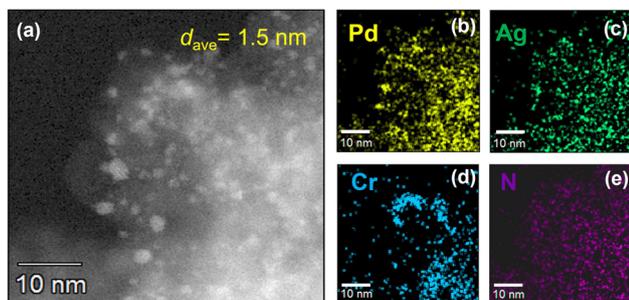
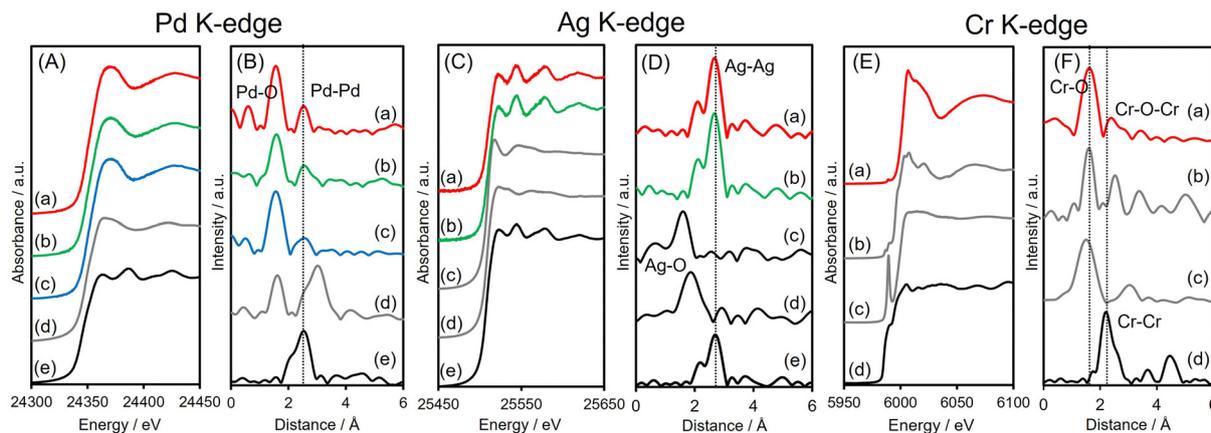


Fig. 1 (a) HAADF-STEM image of PdAgCr/amine-MSc. EDX mapping of (b) Pd, (c) Ag, (d) Cr, and (e) N over the region shown in (a).





**Fig. 3** (A) Pd K-edge XANES and (B) FT-EXAFS spectra of (a) PdAgCr/amine-MSC, (b) PdAg/amine-MSC, (c) Pd/amine-MSC, (d) PdO and (e) Pd foil; (C) Ag K-edge XANES and (D) FT-EXAFS spectra of (a) PdAgCr/amine-MSC, (b) PdAg/amine-MSC, (c) Ag<sub>2</sub>O, (d) AgNO<sub>3</sub> and (e) Ag foil; and (E) Cr K-edge XANES and (F) FT-EXAFS spectra of (a) PdAgCr/amine-MSC, (b) Cr<sub>2</sub>O<sub>3</sub>, (c) CrO<sub>3</sub>, and (d) Cr foil.

**Table 1** Curve fitting results determined by Pd, Ag, and Cr K-edge FT-EXAFS data

	K-edge	Shell	CN	$R/\text{Å}$	$\sigma^2$
PdAgCr/amine-MSC	Pd	Pd-O(N)	4.7	2.00	0.0067
		Pd-Pd	6.8	2.74	0.0065
	Ag	Ag-Ag	11.3	2.89	0.0077
	Cr	Cr-O	4.6	2.01	0.0032
PdAg/amine-MSC	Pd	Pd-O(N)	4.8	2.02	0.0089
		Pd-Pd	9.3	2.76	0.015
	Ag	Ag-Ag	11.3	2.82	0.0073
Pd/amine-MSC	Pd	Pd-O(N)	5.0	2.01	0.0078
		Pd-Pd	9.6	2.73	0.015

of Pd-Pd bonds ( $\text{CN}_{\text{Pd-O}}/\text{CN}_{\text{Pd-Pd}} = 0.69$ ) in the case of PdAgCr/amine-MSC was larger than those for PdAg/amine-MSC (0.52) and Pd/amine-MSC (0.52) (Table 1). These results are in good agreement with the Pd 3d XPS data and suggest that the Pd was partially oxidized following modification with CrO<sub>x</sub> and/or Cr(OH)<sub>3</sub>.

The Ag K-edge XANES spectra obtained from PdAgCr/amine-MSC and PdAg/amine-MSC were similar to that of Ag foil and their FT-EXAFS spectra showed a single intense peak ascribed to contiguous Ag-Ag bonds at approximately 2.6–2.8 Å without any noticeable evidence of Ag-O bonds (Fig. 3C and D). This peak is shifted to a slightly shorter interatomic distance compared with that for pure Ag foil, indicating the presence of Pd-Ag bonds. The inverse FT was well fitted using Ag-Ag shells with CN = 11.3 and  $R = 2.89$  Å for PdAgCr/amine-MSC, and CN = 11.3 and  $R = 2.82$  Å for PdAg/amine-MSC, indicating that Ag formed metallic NPs in both samples (Table 1 and Fig. S5B, ESI†). These results are in agreement with the Ag 3d XPS data.

The Cr K-edge XANES spectrum of CrO<sub>3</sub> displayed an intense pre-edge peak at 5992 eV which is characteristic of terminal Cr<sup>6+</sup>=O bonds in a tetrahedral coordination (Fig. 3E).<sup>44,45</sup> The XANES spectrum obtained for PdAgCr/amine-MSC was different from that for CrO<sub>3</sub> but resembled that for Cr<sub>2</sub>O<sub>3</sub>, which

contains Cr<sup>3+</sup>. The edge energy for PdAgCr/amine-MSC was higher than that for Cr foil, confirming that the Cr in this sample was in the +3 oxidation state. The length of the first Cr-O bond in PdAgCr/amine-MSC as determined from its FT-EXAFS spectrum was comparable to that in Cr<sub>2</sub>O<sub>3</sub> but longer than that in CrO<sub>3</sub>. These data suggest the absence of Cr=O double bonds in PdAgCr/amine-MSC (Fig. 3F). A curve-fitting analysis also confirmed that the interatomic distance for the first Cr-O bond was 2.01 Å, in conjunction with a CN of 4.6 (Table 1 and Fig. S5C, ESI†). Moreover, the intensity of the peak associated with the second coordination sphere for PdAgCr/amine-MSC was significantly weaker than the intensity obtained for Cr<sub>2</sub>O<sub>3</sub>. In the case of the PdAgCr specimen, this peak was attributable to contiguous Cr-O-M (M: Cr or Pd) bonds because of the absence of evidence for Ag-O bonds in the Ag K-edge FT-EXAFS spectrum. The reduced intensity in this region was ascribed to the formation of small, structurally disordered nanoclusters. No evidence for metallic bonds, which were detectable at approximately 2.3 Å in the case of Cr foil, was observed. Because the reduction potentials for Cr<sup>3+</sup> ions ( $E^\circ(\text{Cr}^{3+}/\text{Cr}^{2+}) = -0.42$  V,  $E^\circ(\text{Cr}^{2+}/\text{Cr}^0) = -0.90$  V vs. NHE) are more negative than those for Pd<sup>2+</sup> and Ag<sup>+</sup> ions ( $E^\circ(\text{Pd}^{2+}/\text{Pd}^0) = +0.99$  V,  $E^\circ(\text{Ag}^+/\text{Ag}^0) = +0.80$  V vs. NHE), the reduction of Cr<sup>3+</sup> ions was inhibited to a greater extent than for Pd<sup>2+</sup> and Ag<sup>+</sup> ions, and Cr ions preferentially constitute extremely small clusters.

Based on these data, a reasonable structural model for PdAgCr/amine-MSC was devised. In this model, Ag atoms are preferentially located in the core regions of the NPs, while Pd atoms tend to be situated in the shells. In addition, the amorphous CrO<sub>x</sub> phase including Cr(OH)<sub>3</sub> is partially located on the surfaces of these Pd-rich shells as a result of the formation of Pd-O-Cr bonds, as illustrated in Scheme 1. This configuration is based on the EXAFS analysis showing that the Pd atoms in PdAgCr/amine-MSC were partially oxidized, either because of the interaction with Cr atoms or the coordination with N atoms in the surface amine groups by the exposure to the surface.



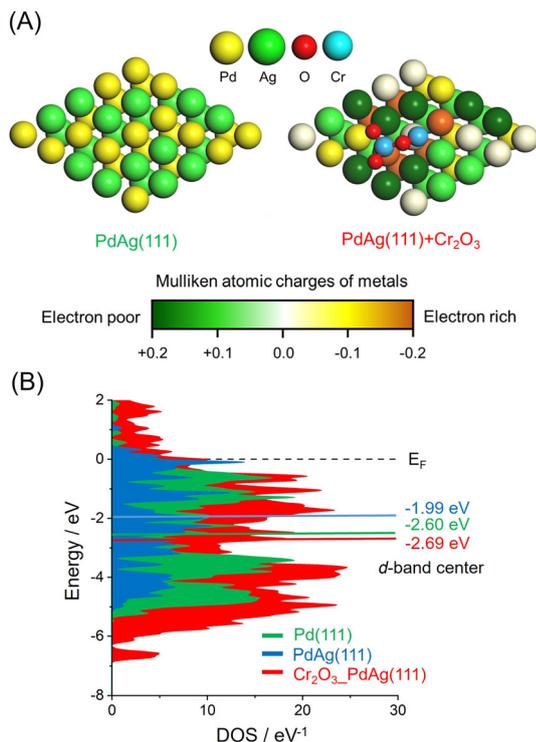


Fig. 4 (A) The optimized structure of pristine PdAg(111) and PdAg(111) associated with one stoichiometric Cr<sub>2</sub>O<sub>3</sub> cluster with a color overlay of representative Mulliken atomic charge determined using density functional theory calculations. (B) Calculated DOS plots.

The effect of the CrO<sub>x</sub> phase on the electronic state of the Pd and Ag atoms was assessed by calculating atomic charges using DFT. These calculations were based on supercell slab models consisting of 5 × 5 surface unit cells including PdAg(111) interacting with Cr<sub>2</sub>O<sub>3</sub> clusters together with pristine PdAg(111). The optimized structures are shown in Fig. 4A, in which the colors of Pd atoms indicate the calculated Mulliken atomic charges. These investigations indicated that the integration of Pd with Ag induced charge transfer from Ag atoms to Pd atoms to produce electron-rich Pd<sup>δ-</sup> and electron-deficient Ag<sup>δ+</sup>. In addition, Pd atoms in the vicinity of the Cr atoms in Cr<sub>2</sub>O<sub>3</sub> were determined to have a large negative charge as a consequence of electron transfer from the Cr atoms. In contrast, Ag atoms bound to the O atoms of Cr<sub>2</sub>O<sub>3</sub> were highly positive, suggesting electron transfer from the Ag atoms to the O atoms. These electronic interactions generated unique ensemble sites and a significant electronic gap based on the adjacent electron-deficient Ag<sup>δ+</sup> and electron-rich Pd<sup>δ-</sup>. Density of states (DOS) calculations showed that the d-band center of PdAg(111) associated with Cr<sub>2</sub>O<sub>3</sub> was located at -2.69 eV and so had a similar energy to that of -2.60 eV for PdAg(111). However, this value was more negative than that of -1.99 eV for Pd(111) (Fig. 4B). These findings confirmed that modification with CrO<sub>x</sub> did not change the electronic structure of the catalyst in the bulk but rather caused uneven charge distribution on the surface of the material.

## 2.2. Catalytic activity during FA dehydrogenation

Dehydrogenation of FA was conducted in an aqueous solution containing HCOOH and HCOONa at a molar ratio of 9/1. No induction period was observed during periodic monitoring of the reaction (Fig. 5A), and H<sub>2</sub> and CO<sub>2</sub> were generated in a molar ratio of almost 1, indicating complete dehydrogenation of the FA. PdAgCr/amine-MSC displayed the highest activity among the various catalysts and exhibited substantially higher performance than the monometallic Pd and binary PdAg materials.<sup>46</sup> Using the optimal FA concentration (Fig. S6, ESI<sup>†</sup>), a turnover frequency (TOF) of 6898 h<sup>-1</sup> (based on the mass of Pd employed) was achieved, which was 1.6 times larger than the value for the PdAg catalyst. The catalytic activity attained by PdAgCr/MSC without amine modification was quite low, confirming that the amine functional group plays a crucial role in determining the elementary steps in the catalytic dehydrogenation of FA, especially in the O–H bond dissociation step, as discussed elsewhere.<sup>29,31,46</sup> In addition, the reaction essentially did not occur when using pure Ag or Cr. The addition of Mn, Ni or Co was found to slightly increase the catalytic activity, while the presence of Fe, Zn, Ga or Cu reduced the dehydrogenation performance (Fig. 5B). These results indicate that a significant synergistic effect was obtained from the ternary NPs. The effect of the composition of the PdAgCr catalyst was investigated by simply varying the initial molar ratio of the metal precursors. Fig. 5C shows the effect of the Cr mole fraction relative to the combined moles of Pd and Ag on the Pd-based TOF and indicates that the activity was maximized at a Cr mole fraction of 0.5. This volcano type activity relationship verified that the unusual synergistic effect observed in the present work originated from the combination of the three metals and also

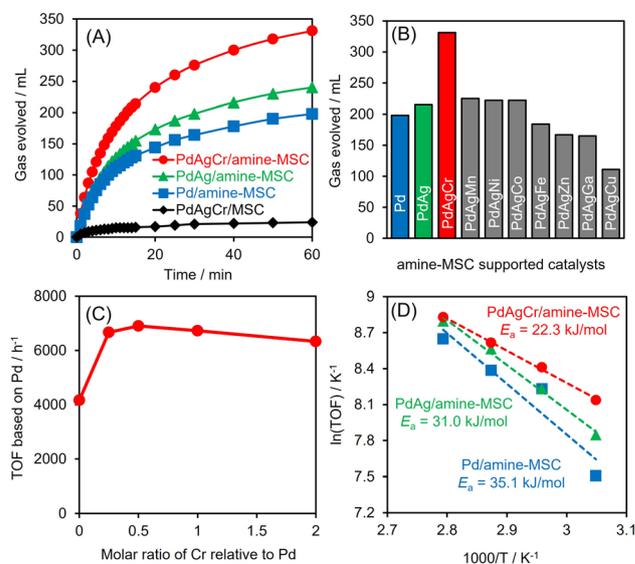


Fig. 5 (A) Time course of the reaction in the dehydrogenation of FA using various amine-MSC supported catalysts. (B) Effect of the addition of third metals to the Pd-based catalysts. (C) TOF versus mole fraction of Cr for the PdAg/amine-MSC catalyst. (D) Arrhenius plots in the dehydrogenation of FA.



demonstrated that an excess of  $\text{CrO}_x$  can suppress access to the active Pd centers.

The Arrhenius plot in Fig. 5D demonstrates that the activation energy ( $E_a$ ) for the dehydrogenation of FA decreased significantly with increasing number of catalyst components, with values of  $22.3 \text{ kJ mol}^{-1}$  for PdAgCr/amine-MSC,  $31.0 \text{ kJ mol}^{-1}$  for PdAg/amine-MSC and  $35.1 \text{ kJ mol}^{-1}$  for Pd/amine-MSC. These results indicate that modification with  $\text{CrO}_x$  boosted the reaction rate in the elementary step during the dehydrogenation of FA. Notably, PdAg/ $\text{Cr}_2\text{O}_3$  prepared using the same impregnation method followed by  $\text{NaBH}_4$  reduction showed almost no catalytic activity, indicating that  $\text{Cr}_2\text{O}_3$  itself does not act as an efficient support. The mean particle size determined by TEM analysis was  $7.5 \text{ nm}$ , which is larger than that of PdAgCr/amine-MSC (Fig. S7, ESI†). The Pd 3d peaks in XPS analysis generated by PdAg/ $\text{Cr}_2\text{O}_3$  shifted to higher binding energies by  $1 \text{ eV}$  compared with those obtained from PdAgCr/amine-MSC (Fig. S8A, ESI†). Similarly, the shift toward higher binding energies by  $1 \text{ eV}$  can be observed in Ag 3d XPS spectra (Fig. S8B, ESI†). These shifts were attributed to the ability of  $\text{Cr}_2\text{O}_3$  to withdraw electrons from PdAg alloy NPs, as a consequence of the relatively low electronegativity among metal oxides.<sup>47</sup>

It should be noted that the present catalytic system efficiently suppressed the undesirable formation of CO *via* the reaction  $\text{HCOOH} \rightarrow \text{H}_2\text{O} + \text{CO}$ . No CO was detected by the gas chromatography analysis method (the detection limit is approximately less than  $2 \text{ ppm}$ ). This concentration of CO was significantly lower than that generated during conventional gas reforming from methanol, ethanol and methane, and meets the requirement provided in the PEMFC standard, meaning a CO concentration below  $10 \text{ ppm}$ .<sup>48</sup> Upon completion of the reaction, the heterogeneous catalyst was readily separated from the reaction system and analysis of the filtrate by inductively coupled plasma confirmed that leaching of the metals from the catalyst was negligible. From these results it may be deduced that the reaction occurred on the solid surfaces of the material rather than in solution. The recovered catalyst could be recycled at least three times while retaining  $90\%$  of its original activity (Fig. S9, ESI†). TEM observations showed that the particle size remained virtually unchanged after the reaction without the formation of large agglomerated NPs (Fig. S10, ESI†). XAFS analyses also indicated a lack of structural changes in the K-edge spectrum obtained from the recovered catalyst relative to that for the fresh material (Fig. S11, ESI†). Unfortunately, the catalytic activity decreased to almost half of its original activity after the fourth recycling experiment. A critical reason for the decreased activity is unclear so far. Moreover, the maximum turnover frequency achieved in this study is  $6898 \text{ h}^{-1}$ , which is comparable to or higher than those of other more recently reported catalytic systems,<sup>49–52</sup> but still lower than those of the most pronounced catalytic systems.<sup>53,54</sup> Thus, the further improvement of activity as well as the durability of the present catalytic system is now under investigation.

In a preliminary experiment, the activity of PdAgCr/amine-MSC was evaluated in the hydrogenation of  $\text{CO}_2$  with  $\text{H}_2$  to give

FA using a  $1.0 \text{ M}$  aqueous  $\text{NaHCO}_3$  solution at  $\text{pH} = 8.5$  under a total pressure of  $2.0 \text{ MPa}$  ( $\text{H}_2:\text{CO}_2 = 1:1$ , volume ratio) at  $100^\circ\text{C}$ . PdAgCr/amine-MSC showed a TON of 893 after 24 h with  $>99\%$  selectivity, which is larger than those obtained with PdAg/amine-MSC (TON = 830) and Pd/amine-MSC (TON = 310), indicating the synergistic effect between Pd and Ag as well as the modification with  $\text{CrO}_x$ . It can be concluded that unique ensemble sites on the PdAg nanoparticle surfaces, at which electron-deficient  $\text{Ag}^{\delta+}$  and electron-rich  $\text{Pd}^{\delta-}$  atoms were adjacent, are also crucial for attaining high activity during hydrogenation of  $\text{CO}_2$  with  $\text{H}_2$ . Thus, the electron-deficient  $\text{Ag}^{\delta+}$  facilitates the adsorption of  $\text{HCO}_3^-$  ions, whereas the electron-rich  $\text{Pd}^{\delta-}$  accelerates not only  $\text{H}_2$  dissociation but also the attack of dissociated H atoms on C atoms in  $\text{HCO}_3^-$  ions.<sup>55</sup>

### 2.3. FA dehydrogenation mechanism

The dehydrogenation reaction proceeds exclusively to afford  $\text{H}_2$  and  $\text{CO}_2$  in a molar ratio of  $1:1$ . The presence of amine functional groups on supports significantly boosted the activity. The addition of  $\text{HCOONa}$  to the reaction mixture increased  $\text{H}_2$  productivity, indicating the participation of the Pd–formate species as a reaction intermediate. Upon consideration of these preliminary insights as well as the results of previous studies, a possible reaction mechanism for the dehydrogenation of FA assisted by weakly basic surface  $-\text{NH}_2$  functional groups in the vicinity of metal NPs was devised and is presented in Fig. 6A.<sup>31,46,56–59</sup> In this process, the  $-\text{NH}_2$  groups act as proton scavengers to promote the dissociation of O–H bonds in FA molecules and generate metal–formate species along with  $-\text{NH}_3^+$  groups (Step 1). Each formate intermediate subsequently undergoes isomerization (Step 2), followed by C–H bond dissociation, to produce  $\text{CO}_2$  and a metal hydride (Step 3). Finally, the hydride reacts with a neighboring  $-\text{NH}_3^+$  group to form

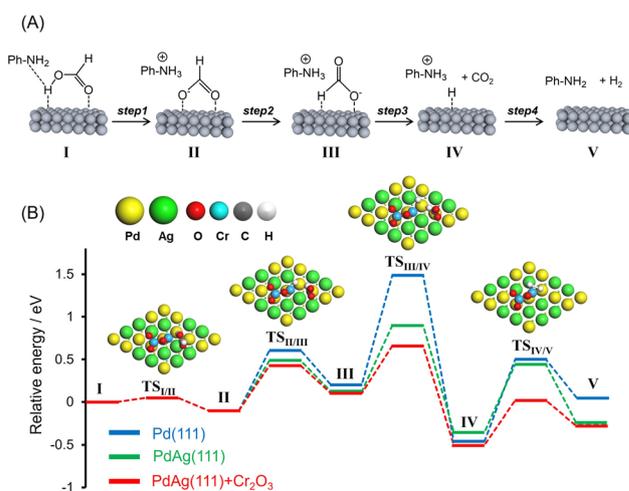


Fig. 6 (A) Possible reaction mechanism and (B) potential energy profiles for dehydrogenation of formic acid calculated using pristine Pd(111), PdAg(111), and PdAg(111) associated with one stoichiometric  $\text{Cr}_2\text{O}_3$  cluster in the absence of amine molecules. The representative TS structures obtained in the calculation of PdAg(111) +  $\text{Cr}_2\text{O}_3$  are shown.



molecular hydrogen (Step 4) with the simultaneous regeneration of the initial active species.

It is likely that FA molecules will encounter amine groups in the vicinity of the NPs prior to adsorption on the surface of the catalyst, as a result of the acid–base interaction. The present DFT calculations demonstrate that the optimized structure of the resulting acid–base pair has a *cis*-M(O)–M(O–C)-bridged configuration (M = Pd or Ag) in which the carbonyl oxygen and non-carbonyl oxygen interact with Ag and Pd atoms, respectively. The O–H bond length (1.075 Å) in this optimized configuration is larger than the value of 0.985 Å for free *trans*-HCOOH, suggesting that the O–H bond of the FA undergoes a spontaneous dissociation without any transition state when an incoming HCOOH molecule is coupled with a phenylamine group. Kinetic isotope effect (KIE) studies using HCOOH and HCOOD show that the  $k_{\text{H}}/k_{\text{D}}$  ratios obtained from competitive reactions using the Pd-based catalysts supported on amine-MSC were quite small (approximately 1) compared with that obtained using PdAgCr/MSc without the amine modification (Table 3, entries 1–3 vs. 4). These results provide clear evidence that the addition of –NH<sub>2</sub> groups promoted cleavage of the O–H bonds in the FA.

To better understand the role of interfacial CrO<sub>x</sub> modification in catalyzing the FA dehydrogenation, potential energy profiles were generated using DFT calculations employing Pd(111), PdAg(111) and PdAg(111) slab models together with a Cr<sub>2</sub>O<sub>3</sub> cluster in the absence of amine molecules (Fig. 6B). The calculated activation energies ( $E_{\text{a}}$ ) are summarized in Table 2. The *trans*-M(O)–M(H–O)-bridged configuration (I, M = Pd or Ag) was determined to be the most energy favorable HCOOH adsorption structure. In the case of PdAg(111) and PdAg(111) models associated with a Cr<sub>2</sub>O<sub>3</sub> cluster, the adsorbed molecule undergoes O–H bond dissociation and forms HCOO<sup>−</sup> as an intermediate (II) *via* transition state TS<sub>I/II</sub> with an energy barrier of 0.05 eV (Step 1). Subsequently, this species is isomerized to a *trans*-M(H)–Pd(O)-bridged HCOOH configuration (III) *via* transition state TS<sub>II/III</sub> with an energy barrier of 0.53 eV (Step 2). Following this, C–H bond scission occurs *via* TS<sub>III/IV</sub> to form CO<sub>2</sub> and a Pd–H species with an energy barrier of 0.56 eV (Step 3). Finally, the catalytic cycle is completed with the release of H<sub>2</sub> *via* TS<sub>IV/V</sub> with an energy barrier of 0.52 eV (Step 4).

In the case of the Pd(111) and PdAg(111) models, the dissociation of the O–H bond occurs with a similar barrier to that obtained with the PdAg(111) model associated with a Cr<sub>2</sub>O<sub>3</sub> cluster. In sharp contrast, the activation energies for Steps 2, 3

**Table 2** Activation energies ( $E_{\text{a}}$ ; eV) for various steps during dehydrogenation from formic acid

Model	Step 1 (I → II)	Step 2 (II → III)	Step 3 (III → IV)	Step 4 (IV → V)
	O–H bond dissociation	Isomerization of formate	C–H bond dissociation	H <sub>2</sub> desorption
Pd(111)	0.05	0.71	1.29	0.96
PdAg(111)	0.05	0.49	0.78	0.79
PdAg(111) + Cr <sub>2</sub> O <sub>3</sub>	0.05	0.53	0.56	0.52

**Table 3** KIE in the dehydrogenation of FA for various Pd-based catalysts

Entry	Catalyst	Formic acid	$k_{\text{H}}/k_{\text{D}}$
1	Pd/amine-MSc	HCOOH HCOOD	1.14
2	PdAg/amine-MSc	HCOOH HCOOD	1.07
3	PdAgCr/amine-MSc	HCOOH HCOOD	1.00
4	PdAgCr/MSc	HCOOH HCOOD	1.67
5	Pd/amine-MSc	HCOOH DCOOH	2.20
6	PdAg/amine-MSc	HCOOH DCOOH	1.37
7	PdAgCr/amine-MSc	HCOOH DCOOH	1.03

and 4 were determined to increase in the order of PdAg(111) + Cr<sub>2</sub>O<sub>3</sub> < PdAg(111) < Pd(111). The lower energy barriers determined for the PdAg(111) model associated with a Cr<sub>2</sub>O<sub>3</sub> cluster as compared with the Pd(111) and PdAg(111) models confirmed the positive effect of the modification with Cr<sub>2</sub>O<sub>3</sub> in terms of boosting the FA dehydrogenation reaction. Even though this effect is evident with regard to several reaction steps, the effect on C–H bond dissociation was most pronounced (Step 3). This was also determined to be the rate-determining step in the case of pristine Pd(111).

These results agree with the data obtained from a KIE analysis performed using HCOOH and DCOOH. The  $k_{\text{H}}/k_{\text{D}}$  ratio found for PdAgCr/amine-MSc was 1.03, which was smaller than those observed for the monometallic Pd (2.20) and binary PdAg samples (1.37) (Table 3, entries 5–7). These results are consistent with the order of catalytic activity and indicate that the PdAg and PdAgCr catalysts facilitated C–H bond dissociation of the metal-formate intermediate. One possible reason for the improved catalytic activity of the binary PdAg catalyst could be the formation of an electron-rich Pd<sup>δ−</sup> species based on energy transfer from Ag to Pd atoms, as confirmed by the XPS analysis (Fig. 2B) and DOS calculations (Fig. 4). In this system, the metal-formate intermediate generated over electron-rich Pd<sup>δ−</sup> species was stabilized by significant back-donation from the metal to the formate that, in turn, facilitated selective C–H bond dissociation *via* a bidentate configuration.<sup>27</sup> The enhancement of the rate-determining C–H bond dissociation step in the case of the PdAgCr catalyst can be explained by considering the electronic state of reaction intermediate III in Fig. 6A. The DFT-optimized configurations are summarized in Fig. 7. In this scenario, the H atom of HCOO<sup>−</sup> adsorbed at an electron-deficient Ag<sup>δ+</sup> site over PdAg(111) interacting with a Cr<sub>2</sub>O<sub>3</sub> cluster is more positively charged because of the reduced distance of 2.37 Å compared with the values for Pd(111) (3.29 Å) and PdAg (111) (2.42 Å). Accordingly, the electronic charge of the H atom becomes negative, accompanied by elongation of the C–H bond, which is then more likely to undergo dissociation.

The synergistic effect involved in promoting the final H<sub>2</sub> desorption step was also assessed (Step 4). Reactions performed with a flow of H<sub>2</sub> and D<sub>2</sub> through the catalyst indicated



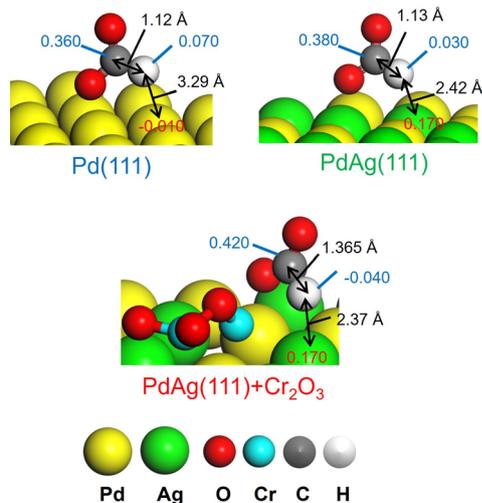


Fig. 7 Representative Mulliken atomic charges as determined by DFT calculations in reaction intermediate III for Pd(111), PdAg(111), and PdAg(111) associated with the  $\text{Cr}_2\text{O}_3$  cluster.

that the formation rate for HD increased in the order of monometallic Pd < binary PdAg < ternary PdAgCr (Fig. S12, ESI<sup>†</sup>), in agreement with the lower reaction barriers for the  $\text{H}_2$  desorption step calculated using DFT (Table 2). It has been suggested that the energy of an adsorbed atom is related to the intrinsic electronic properties of the metallic particles on the catalyst. Specifically, a greater downward shift of the d-band center from the Fermi level of a metal increases the possibility that anti-bonding states will be formed, leading to weaker binding of the adsorbed atom.<sup>60</sup> Thus, the binary PdAg catalyst exhibited a lower activation energy for  $\text{H}_2$  desorption than the monometallic Pd catalyst because the d-band center of PdAg(111) is deeper than that of Pd(111) (Fig. 4B). The participation of multiple interactions induced by the modification with  $\text{CrO}_x$  in this process was also apparent. The dissociated hydride species at electron-rich  $\text{Pd}^{\delta-}$  sites were determined to be negatively charged, while those at electron-deficient  $\text{Ag}^{\delta+}$  sites were found to be positive. These hydrides having different charges generated at sites provided by  $\text{CrO}_x$  modification are believed to have increased the production of hydrogen molecules.

### 3. Conclusions

This work increased the activity of a binary PdAg catalyst during dehydrogenation of FA based on interfacial modification with a highly dispersed amorphous  $\text{CrO}_x$  phase. The combined effects of the weakly basic amine groups on the material surface and the synergistic catalysis provided by PdAgCr NPs explain the improved activity of this catalyst. The basic amine groups on the surface of the support promoted the dissociation of O–H bonds in FA molecules, while the  $\text{CrO}_x$  phase enhanced the rate-determining C–H bond dissociation step of the metal-formate intermediate as well as the  $\text{H}_2$  desorption step. These effects resulted from the formation of unique ensemble sites

involving adjacent electron-deficient  $\text{Ag}^{\delta+}$  and electron-rich  $\text{Pd}^{\delta-}$  atoms. These findings provide vital information that will assist in the future design of catalytically active centers for dehydrogenation of FA. They also emphasize the importance of interface design for further improvement of catalytic performance as a means of realizing environmentally-friendly hydrogen release systems.

## 4. Experimental section

### 4.1. Materials

Acetone, nitric acid ( $\text{HNO}_3$ ), *p*-phenylenediamine,  $\text{AgNO}_3$ ,  $\text{Cr}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ , sodium hydroxide (NaOH), sodium borohydride ( $\text{NaBH}_4$ ), formic acid (FA) and sodium formate (SF) were purchased from Nakalai Tesque, Inc. Pd(II) nitrate solution was obtained from Wako Pure Chemical Ind. Co., Ltd. Mesoporous carbon (MSC, average pore diameter:  $100 \pm 10 \text{ \AA}$ ) was obtained from Aldrich Chemical Co. All commercially available compounds were used as received. Distilled water was employed as the reaction solvent.

### 4.2. Synthesis of amine functionalized MSC (amine-MSC)

The detailed procedure is described elsewhere.<sup>46</sup> Briefly, the MSC (1.0 g) was mixed with aqueous nitric acid solution (60%, 50 mL) at  $80 \text{ }^\circ\text{C}$  for 5 h. The sample was collected by filtration, washed with distilled water, and dried under vacuum overnight. The acid treated MSC (1.0 g) was added into 500 mL of distilled water and sonicated for 2 h. The aqueous dispersion solution was mixed with acetone (50 mL) containing *p*-phenylenediamine (PDA, 3.24 g) and stirred at room temperature for 48 h. The resultant sample was separated by filtration and washed with acetone and distilled water until no color was confirmed in the filtrate, followed by drying under vacuum overnight.

### 4.3. Synthesis of the PdAgCr supported amine-MSC catalyst (PdAgCr/amine-MSC)

The amine-MSC (0.3 g) was mixed with an aqueous solution (80 mL) containing  $\text{Pd}(\text{NO}_3)_2$  (0.606 mL, Pd 47 mM),  $\text{AgNO}_3$  (2.85 mL, 10 mM), and  $\text{Cr}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$  (1.425 mL, 10 mM). After mixing, NaOH aqueous solution (4.2 mL, 1M) was added to adjust the pH around 10 and stirred for 30 min. Subsequently, the sample was reduced by  $\text{NaBH}_4$  under an Ar atmosphere, collected by filtration, and dried under vacuum overnight, giving PdAgCr/amine-MSC (Pd 1 wt%, Pd:Ag:Cr = 1:1:0.5). Monometallic Pd, binary PdAg, and ternary PdAgM (M = Mn, Co, Ga, Ni, Zn, Fe, and Cu) supported on amine-MSC catalysts were also prepared by the same method. Metal loadings were determined using inductively coupled plasma (ICP) analysis.

### 4.4. Characterization

Powder XRD patterns, BET surface area, ICM-ES, XPS spectra, and TEM images were acquired according to the procedure described previously.<sup>46</sup> Quantification of the loading amount of amine functionality in samples was performed by two methods.



One is the CHN elemental analysis using a Micro Corder JM-10 (J-Science Lab Co., Ltd). Another is thermogravimetric (TG) analysis using a Rigaku Thermo Plus EVO2 TG8121 system from room temperature to 700 °C at a heating rate of 10 °C min<sup>-1</sup> in air flow (10 mL min<sup>-1</sup>). STEM images and elemental maps were obtained using a JEOL-ARM 200F apparatus equipped with a Kve energy-dispersive X-ray detector (JED-2300T) operating at 200 kV. Pd, Ag, and Cr K-edge XAFS spectra were recorded using a fluorescence yield collection technique with a Si(111) monochromator at the 01B1 beamline station in the SPring-8 facility, JASRI, Harima, Japan (proposal no. 2020A0523 and 2020A1062). Data reduction was performed using the REX2000 software program (Rigaku). To obtain radial structure functions, Fourier transformation of the  $k^3$ -weighted normalized EXAFS data was carried out over the range  $3.0 \text{ \AA} < k/\text{\AA}^{-1} < 12 \text{ \AA}$ . The H<sub>2</sub>-D<sub>2</sub> exchange reaction was monitored using a BELCAT II system (MicrotracBEL Corp.). After pretreatment under He flow at 323 K for 30 min, the H<sub>2</sub>-D<sub>2</sub> exchange reaction was performed at 323 K using 25% H<sub>2</sub>/He and 25% D<sub>2</sub>/He at a flow rate of 25 mL min<sup>-1</sup>. The products (H<sub>2</sub>, HD and D<sub>2</sub>) were analyzed with an online mass spectrometer. The rate constant was determined from the produced HD amount.

#### 4.5. DFT calculations

All DFT calculations were performed with the CASTEP program in the Materials Studio 17.2 software package.<sup>61,62</sup> The generalized gradient approximation exchange–correlation functional proposed by Perdew, Burke, and Ernzerhof was combined with the double numerical basis set plus polarization functions. A supercell slab consisting of a surface  $5 \times 5$  unit cell with three atomic (111) surface layers was adopted. For the calculation, periodic boundary conditions were applied to the slabs. The slab was separated by a vacuum space with a height of 20 Å. The reactant atoms and their adjacent atoms were relaxed during geometry optimizations, and the other layers were fixed at the corresponding bulk positions. Transition states (TSs) were determined using the nudged elastic band method, and the activation energy was defined by the energy difference between the TS and reactant.

#### 4.6. General procedure for the catalytic dehydrogenation of formic acid

The catalyst was placed in a reaction vessel (30 mL) equipped with a reflux condenser and a gas burette. After purging with N<sub>2</sub>, FA : SF = 9 : 1 aqueous solution (1 M, 10 mL) was added to the reaction vessel and reacted at 75 °C with magnetic stirring. TOF values [h<sup>-1</sup>] were determined according to the equation:

$$\text{TOF} = P_{\text{atm}} V_{\text{H}_2} / RT N_{\text{Pd}} t$$

in which  $P_{\text{atm}}$  is the atmospheric pressure,  $V_{\text{H}_2}$  is the generated volume of H<sub>2</sub>,  $R$  is the gas constant,  $T$  is room temperature,  $N_{\text{Pd}}$  is the mole number of Pd or surface exposed Pd, and  $t$  is the reaction time.

#### 4.7. General procedure for the catalytic hydrogenation of CO<sub>2</sub>

CO<sub>2</sub> hydrogenation to FA was conducted with a batch reactor system (stainless steel autoclave (60 mL)) using a catalyst (50 mg) and 1.0 M aqueous NaHCO<sub>3</sub> solution (10 mL) under a total pressure of 2.0 MPa (H<sub>2</sub>:CO<sub>2</sub> = 1:1, volume ratio) at 100 °C. After 24 h, disodium succinate (external standard) was added and the yield of formic acid was determined by HPLC using a Shimadzu HPLC instrument equipped with a Bio-Rad Aminex HPX-87H Ion Exclusion Column (300 mm × 7.8 mm). 5 mm H<sub>2</sub>SO<sub>4</sub> (0.500 mL min<sup>-1</sup>) was used as a mobile phase. TON values were determined by dividing the quantity of FA produced after 24 h by the moles of total Pd.

### Author contributions

K. M. supervised the project, performed all the calculations and wrote the manuscript. T. F. performed the catalyst preparation, characterization, and catalytic reactions. H. Y. helped supervise the project.

### Conflicts of interest

There are no conflicts to declare.

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