RSC Advances

PAPER

Cite this: RSC Adv., 2016, 6, 105940

Monodisperse AgPd alloy nanoparticles as a highly active catalyst towards the methanolysis of ammonia borane for hydrogen generation

Daohua Sun,* Pengyao Li, Bin Yang, Yan Xu, Jiale Huang and Qingbiao Li

Monodisperse AgPd nanoparticles (NPs) were successfully synthesized by coreducing silver nitrate and palladium chloride in the presence of oleylamine. Characterization techniques such as transmission electron microscopy, energy-dispersive X-ray spectroscopy, and X-ray diffraction were employed to verify the alloy nature of the obtained NPs. These AgPd NPs, supported on carbon, were found to be active catalysts for the generation of hydrogen from the methanolysis of ammonia borane at room temperature. The catalytic activity depended on the alloy composition, and the $Ag_{30}Pd_{70}$ NPs displayed the greatest activity among the tested catalysts, with a very high total turnover frequency value of 366.4 min⁻¹ and activation energy of 37.5 kJ mol⁻¹. In a reusability test, the Ag₃₀Pd₇₀/C catalyst retained 67% of its initial activity and 100% conversion after eight cycles. **PAPER**
 Solution
 Solutio

Received 30th August 2016 Accepted 26th October 2016

DOI: 10.1039/c6ra21691a

www.rsc.org/advances

1. Introduction

Storage and transport of hydrogen are the widely accepted technological barriers to the development of the so-called hydrogen economy.¹ As the lightest element in the periodic table, hydrogen in the gas form is extremely difficult to store. Considering economically viable and safe practices, efforts have been directed towards the development of chemical hydrides as a potential onboard storage form of hydrogen.^{2,3} Ammonia borane (AB) has been considered to be the most promising candidate among the chemical hydrides.4,5 This molecule is stable in the solid state at ambient temperatures and has shown an impressive gravimetric capacity of 19.6 wt% H_2 , potentially capable of meeting the 2015 target of the U.S. Department of Energy (9 wt% hydrogen for a material to be practically applicable).^{6,7} Hydrogen stored in AB can be produced and obtained by pure thermal decomposition, hydrolysis and methanolysis using a suitable catalyst. Considering the mild reaction conditions and favorable kinetics, solvolysis is preferred.⁸ The methanolysis or hydrolysis of AB can release 3.0 equivalents of hydrogen in the presence of a suitable catalyst. However, the hydrolysis of AB in concentrated solutions can cause liberation of ammonia gas, which can poison Pt-based fuel cells, and the hydrolysis product ammonium metaborate is difficult to recycle, which can pose enormous challenges to practical applications.9,10 In a previous study by Ramachandran, when methanol was used instead of water, the catalytic methanolysis of AB yielded hydrogen gas without ammonia evolution,

and yielded the recyclable methanolysis product ammonium tetramethaoxyborate.⁶ In addition, the methanolysis reaction can even be initiated at temperatures below 0° C through use of a suitable catalyst, which satisfies applications in cold weather.¹¹

Since methanolysis of AB can achieve a complete release of hydrogen in the presence of a suitable catalyst, finding the best catalyst for this process is very important. Heterogeneous catalysts were the first ones to be tested for the methanolysis of AB. RuCl₃ and RhCl₃ have been shown to exhibit excellent catalytic activity, but they are difficult to isolate and they display low stability, features that seriously restrict their practical application.⁶ Cu@Cu₂O, Cu₂O,¹² Co-Co₂B, Ni-Ni₂B, Co-Ni-B,¹³ and Cu-Cu₂O-CuO/C¹⁴ are cheaper transition metal-based catalysts, but suffer from low activity. The performance of the catalyst in the methanolysis of AB appears to depend on the identity of the metal. Thus, recent studies have been directed toward bimetallic alloy nanoparticles (NPs), as their catalytic properties have been found to be superior to those of either component metal alone due to synergistic effects. Our previous studies have also indicated monodisperse CoPd NPs¹⁵ and CuPd NPs¹⁶ to be more active than Pd-rich NPs in catalyzing the AB methanolysis reaction, with total turnover frequency (TOF) values of 27.7 and 53.2 min^{-1} , respectively, having been attained.

Despite the significant progress made in the synthesis of bimetallic alloy NPs,^{17,18} size-controlled fabrication of nanostructures has become one of the most important and challenging aspects of modern nanotechnology. In the widely used method, the metal NPs are generated from the reduction of metal ions in solution in the presence of a stabilizer or supporting materials.¹⁹ To obtain uniformly sized, shapecontrolled, and highly dispersed monodisperse NPs, "hot

Department of Chemical and Biochemical Engineering, College of Chemistry and Chemical Engineering, National Laboratory for Green Chemical Productions of Alcohols, Ethers and Esters, Xiamen University, Xiamen, 361005, P. R. China. E-mail: sdaohua@xmu.edu.cn; Fax: +86 592 2184822; Tel: +86 592 2183088

injection" techniques, which utilize homogeneous nucleation to synthesize monodisperse nanocrystals in organic solutions, are commonly used. For example, when using this technique, Sun²⁰ prepared CoPd NPs with diameters of 5-12 nm and with a standard deviation of the diameter of less than 7%. Hyeon²¹ even succeeded in the incremental 1 nm size controlled synthesis of iron oxide NPs ranging from 6 to 13 nm. To allow precise control over size and monodispersity, capping agents such as oleic acid and organic metal salt precursor (e.g., acetylacetonate) have usually been employed. $22-24$ These agents, however, increase the costs of the catalytic process. Therefore, it is of crucial importance to develop rational cost-effective approaches to synthesize monodisperse alloy NPs.

Herein, we report the synthesis of monodisperse AgPd alloy NPs from inorganic salt silver(I) nitrate and palladium(I) chloride in oleylamine, and the evaluation of their performance in catalyzing the methanolysis of AB at ambient conditions. We demonstrated that the activity of these AgPd NPs can be enhanced by properly controlling the Ag/Pd molar ratio. In addition, these AgPd NPs showed a very high catalytic activity and prolonged lifetime.

2. Experimental

2.1 Chemicals

AgNO₃, PdCl₂, isopropanol, hexane and methanol were of analytical levels and used as received without any further purification. The borane–ammonia complex (90%) and oleylamine (80–90%) were purchased from Sigma-Aldrich and Aladdin, respectively. All pieces of glassware were washed several times with aqua regia and ordinary distilled water.

2.2 Preparation of AgPd NPs

Typically, under the protection of a constant nitrogen flow, 0.2 mmol AgNO₃ and 0.2 mmol PdCl₂ were added into a fournecked flask with 15 mL of oleylamine. The mixture was heated to 110 °C gradually and kept at this temperature for 0.5 h in order to remove oxygen and moisture from the flask, and then the solution was heated to 210 \degree C at a heating rate of 4 \degree C min^{-1} and maintained at this temperature for 1 h before being cooled to room temperature. Afterwards, the mixture and 40 mL of isopropanol were transferred to centrifuge tubes, and the product was separated by centrifugation at 10 000 rpm for 10 min. Finally, the product was dispersed in hexane.

For comparison, Ag NPs and Pd NPs were also, separately, synthesized and obtained. The Ag NPs were prepared by carrying out thermal decomposition of $AgNO₃$ in the presence of oleylamine. The Pd NPs were prepared by carrying out thermal decomposition of $PdCl₂$ in oleylamine. The procedures used to synthesize the Ag NPs and Pd NPs were similar to that used to synthesize AgPd NPs.

2.3 Preparation of the catalyst

Typically, accurately weighed 10 mg of AgPd nanoparticles were redispersed in 5 mL hexane and 5 mL acetone in a vial with a volume of 20.0 mL, and then a mass of 30 mg of Ketjen carbon

 $({\sim}750~\text{m}^2~\text{g}^{-1})$ was added. The resulting mixture was sonicated for 2 h to ensure that the nanoparticles were completely loaded onto the support. Next, the black solid residue AgPd/C catalyst was obtained after the hexane and acetone were vaporized as a result of a constant flow of nitrogen.

2.4 Evaluation of the catalytic activity

To evaluate the catalytic activity of the AgPd/C catalyst for methanolysis of AB, the classic water-displacement method involving determining the rate of hydrogen generation was performed.¹⁵ Typically, a volume of 10 mL of a methanol suspension of the AgPd/C catalyst was transferred into a reaction flask (25 mL) containing a Teflon-coated stir bar placed on a magnetic stirrer. The reaction temperature was maintained at 25 °C using a constant temperature bath during the process, and the reaction flask was connected to a burette filled up with water to measure the volume of hydrogen. Weighed AB was then added into the reaction flask quickly with stirring at a rate of 800 rpm. The volume of hydrogen generated from methanolysis of AB in the flask was monitored and measured by recording the displacement of the water level at various intervals of time. The reaction was considered to be complete when the generation of more gas was no longer observed. Paper

Distribution in the content of the

2.5 Characterizations

Samples used for TEM and HRTEM analyses were prepared in advance by sonicating AgPd in hexane at a proper concentration and then depositing a drop of the resulting dispersion on a carbon-coated copper grid. The morphologies of the various types of NPs were visualized by using a TEM instrument operating at 300 kV, and an elemental analysis of the AgPd NPs was performed by carrying out an energy-dispersive X-ray spectroscopy (EDX) experiment. The dried and powdered samples were analyzed via X-ray diffraction (XRD) by using an X-ray diffractometer (Phillips, Netherlands) with Cu Ka radiation (40 kV, 30 mA). The composition and metallic content of the NPs were determined by performing inductively coupled plasma mass spectrometry (ICP-MS) (Agilent 7700x). The dried NPs were dissolved in $HNO₃$ to assure the complete dissolution of the metal into the acid, and the obtained solution was then diluted with 2% HNO₃. X-ray photoelectron spectroscopy (XPS) measurements were taken with an X-ray photoelectric spectrophotometer, and all binding energies were referenced to the C 1s peak, at 284.8 eV.

3. Results and discussion

3.1 Synthesis and characterization of the monodisperse AgPd NPs

Zhang has reported the synthesis of AgPd alloy NPs with dimensions of 2.2 nm by coreducing Ag(Ac) and Pd(acac)₂ in oleylamine, oleic acid and 1-octadecene at 180 $^{\circ}$ C.²² We replaced the above precursors with inorganic metal salts to yield a procedure that is more cost-effective but for which obtaining monodisperse nanoparticles is more challenging. Specifically, AgPd NPs were synthesized from $AgNO₃$ and PdCl₂ in

RSC Advances **RSC Advances** Paper **RSC Advances** Paper **Paper Paper Paper Paper** Paper Pap

oleylamine at 220 $^{\circ}$ C under the protection of a constant flow of nitrogen. The morphology and structure of the as-synthesized AgPd NPs were observed by performing TEM. For a molar ratio of Ag to Pd of about $1:1$, the composition of the assynthesized NPs was determined using ICP-MS to be $Ag_{36}Pd_{64}$.

The as-synthesized NPs were observed to be spherical, with a mean diameter of 6.2 nm (Fig. 1(a)). These NPs were monodisperse, with a standard deviation of \sim 5%. STEM-EDX images of the $Ag_{36}Pd_{64}$ NPs are shown in Fig. 1(b)-(e). Close inspection of several as-synthesized NPs suggested the Ag (Fig. 1(d)) and Pd

Fig. 1 (a) TEM image and size histogram (inset) of the AgPd NPs. (b) Distributions of Ag and Pd components along the cross-sectional line profiles of two AgPd NPs. (c) High-magnification STEM image of AgPd NPs. (d and e) EDX elemental maps for Ag (d, green) and Pd (e, yellow) concentrations within four individual AgPd NPs.

Fig. 2 Representative TEM images of (a) Ag, (b) Ag₇₁Pd₂₉, (c) Ag₄₄Pd₅₆, (d) Ag₃₆Pd₆₄, (e) Ag₃₀Pd₇₀, (f) Ag₁₂Pd₈₈ and (g) Pd NPs. (h) Plot of AgPd NPs size versus mole fraction of Ag.

(Fig. 1(e)) atoms to be evenly distributed throughout the NP. Moreover, EDX elemental line scanning of two NPs (Fig. 1(b)) also verified the presence of the alloy structure in these NPs.

The composition and size of AgPd bimetallic NPs could be facilely controlled by changing the ratio of $AgNO₃$ to $PdCl₂$. In our study, the total amounts of $AgNO₃$ and $PdCl₂$ were each set at 0.4 mmol, and ICP analyses indicated that as-synthesized NPs with the formulas $Ag_{71}Pd_{29}$, $Ag_{44}Pd_{56}$, $Ag_{36}Pd_{64}$, $Ag_{30}Pd_{70}$ and $Ag₁₂Pd₈₈$ were obtained when using the molar ratios of AgNO₃ to PdCl₂ of 9 : 1, 7 : 3, 5 : 5, 3 : 7, and 1 : 9, respectively. The TEM images of each of the above five NPs are shown in Fig. $2(b)$ –(f), with an image of the Ag NPs shown in Fig. 2(a) and that of the Pd NPs in Fig. 2(g). All of the as-synthesized NPs were observed to be spherical in shape and well-dispersed, except for the $Ag₁₂Pd₈₈$ and Pd NPs. The results may be attributed to the smaller particles more easily agglomerating during the synthetic process. As shown in Fig. 2(h), when the mole fraction

of Ag was increased from 30% to 100%, the dimensions of the resulting particle increased on average from 5.4 nm to 12.3 nm. The influence of the molar ratio of the metals on the size of the bimetallic alloy NPs has also been reported in earlier studies.^{25,26} The dependence of the size of the alloy NPs on the composition has been indicated to be related to the collision energy, sticking coefficient, rate of nucleation and growth.²⁷

XRD patterns of the monometallic and bimetallic NPs are shown in Fig. 3(b). Each pattern exhibited several diffraction peaks, which can be indexed to the (111), (200), (220), (311) and (222) planes of the fcc structure of Ag and Pd, representing five diffraction rings in the selected area electron diffraction (SAED) pattern (Fig. 3(a)). Fig. 3(c) shows a plot of the diffraction angle versus the Pd molar fraction. All of these XRD peaks gradually shifted to larger angles as the Pd content was increased. As shown in the inset of Fig. 3(c), an intense peak resulted from each of the AgPd NPs produced using $AgNO₃$ to PdCl₂ ratios of

Fig. 3 (a) SAED image from Ag₃₆Pd₆₄ nanoparticles (NPs). (b) XRD patterns and (c) (111) diffraction peak positions of the (a) Ag, (b) Ag₇₁Pd₂₉, (c) Ag₄₄Pd₅₆, (d) Ag₃₆Pd₆₄, (e) Ag₃₀Pd₇₀, (f) Ag₁₂Pd₈₈ and (g) Pd NPs.

Fig. 4 X-ray photoelectron spectroscopy (XPS) patterns showing the (a) Ag 3d region for the Ag NPs, (b) Pd 3d region for the Pd NPs, (c) Ag 3d and (d) Pd 3d regions for the AgPd alloy NPs.

Table 1 X-ray photoelectron spectroscopy (XPS) peak positions of Ag Pd and AgPd NPs

	Binding energy (eV)			
	Ag		Pd	
NPS	$3d_{5/2}$	$3d_{3/2}$	$3d_{5/2}$	$3d_{3/2}$
Ag	368.66	374.65		
$Ag_{44}Pd_{56}$	368.36	374.37	340.96	335.69
$Ag_{36}Pd_{64}$	368.29	374.29	341.20	335.91
$Ag_{30}Pd_{70}$	368.28	374.27	341.42	336.12
Pd			341.28	336.00

Fig. 5 Plots of reaction time versus volume of hydrogen released from the methanolysis of AB catalyzed by AgPd/C catalysts with different compositions (10 mg NPs, [AB] = 100 mM, $T = 25 \pm 1$ °C).

 $9: 1, 7: 3, 5: 5, 3: 7, 1: 9$, respectively. The curves in this figure indicated that the positions of the (111) diffraction peaks from the AgPd NPs were indeed between that from a pure Ag crystal at 38.23° and that from a pure Pd crystal at 39.23°. These results further confirmed the alloy nature of the products synthesized from the various initial molar ratios of $AgNO₃$ to $PdCl₂$.

The Ag 3d and Pd 3d regions of the XPS spectra of three representative AgPd alloy NPs are shown in Fig. 4(c) and (d), respectively. For comparison, Fig. 4(a) and (b) show the XPS spectra of the Ag 3d and Pd 3d regions of monometallic Ag NPs and Pd NPs, respectively. The differences (Δ values) between the positions of the $3d_{5/2}$ and $3d_{3/2}$ peaks of Ag were observed to be 6.0 eV, and the differences between the positions of the $3d_{5/2}$ and $3d_{3/2}$ peaks of Pd were observed to be 5.3 eV. These values were the same as the reported values for zero-valent Ag and Pd, respectively.28,29 Hence, Ag and Pd atoms existed in the zerovalent state in all of the prepared alloy samples. The positions of the XPS peaks are listed in Table 1 for further reference. The binding energy of Ag 3d was clearly shifted to lower energy and the binding energy of Pd 3d shifted to higher energy when the Pd content was increased, indicating that Ag and Pd formed an alloy cluster.³⁰ The formation of interfacial metal–metal bonds for a bimetallic alloy can produce large perturbations to the Hamiltonian, binding energy density, and hence the catalytic properties.³¹ Sun has mentioned that the polarization of AgPd shifts the valence density of state to generate excessive electrons, which makes the AgPd alloy serve as a charge donor in the catalytic reaction.³²

3.2 Methanolysis of AB catalyzed by monodisperse AgPd NPs

The as-synthesized NPs supported on Ketjen carbon acted as catalysts for the generation of hydrogen from the methanolysis of the AB solution. As shown in Fig. 5, pure Ag NPs, pure Pd NPs and 5 compositions of AgPd NPs supported on Ketjen carbon exhibited different levels of catalytic activity for the methanolysis of AB. Interestingly, among the tested catalysts, $Ag_{30}Pd_{70}/C$ NPs displayed the highest catalytic activity with a completion time of 15 s for 100 mM AB, obviously better than those of pure Pd NPs and Ag NPs, which indicated the existence of a synergistic effect between the Ag and Pd. The metal content of the oleylamine-coated $Ag_{30}Pd_{70}$ was determined, using ICP-AES, to be 34.6 wt%. The catalytic activity of $Ag_{30}Pd_{70}/C$ for the methanolysis of AB was also assessed by determining its TOF, which was found to be 366.4 mol of $H₂$ (mol of catalyst $\mathrm{min})^{-1}.$ The very high activity of Ag $_{30}\mathrm{Pd}_{70}$ NPs probably resulted from their small dimensions and regular shapes. **EXAMELIZER ARTISES ARTISES**

OPEN ACCESS ARTISES ARTISES ARTISES ARTISES ARTISES ARTISES ARTISES ARTISES AREAS ARTISES ARTISES AREAS ARTISES AREAS ARTISES AREAS ARTISES AREAS ARTISES AREAS ARTISES AREAS AREAS AREAS AREAS

3.3 Methanolysis kinetics of the $Ag_{30}Pd_{70}/C$ catalyst

As the $Ag_{30}Pd_{70}/C$ catalyst exhibited the highest activity, we chose it for further study of its kinetics. First, the methanolysis reaction was carried out using various dosages of the $Ag_{30}Pd_{70}$ NPs, specifically at 0.5 mg, 2 mg, 5 mg, and 10 mg of the NPs

Fig. 6 (a) Plot of reaction time versus volume of hydrogen generated from the methanolysis of AB catalyzed by various dosages of the Ag₃₀Pd₇₀/ C catalyst. (b) Plot of the natural logarithm of the hydrogen generation rate versus the natural logarithm of the catalyst concentration (rate = mL of H₂ per s; [AB] = 100 mM, $T = 25 \pm 1$ °C).

Fig. 7 (a) Plot of reaction time versus volume of hydrogen generated from methanolysis of AB catalyzed by the Ag₃₀Pd₇₀/C catalyst at various temperatures. (b) Arrhenius plot, i.e., a plot of ln k versus $1/T$ ([AB] = 100 mM, [AgPd] = 3.2 mM).

Table 2 Activities of catalysts in the methanolysis of AB at room temperature

Paper				RSC Advances
	(a) Volume of Hydrogen/mL 70 60 50 40 30 20 10 $\mathbf 0$	10 20 30 Time/s	0° C 10° C 20° C -30° C 50 60 40	(b) 2.4 2.2 2.0 1.8 \times 1.6 ≤ 1.4 1.2 1.0 0.8 $Ink = -4528.98(1/T) + 17.21$ 0.6 0.0035 0.0036 0.0037 0.0033 0.0034 $1/T(K^{-1})$
				Fig. 7 (a) Plot of reaction time versus volume of hydrogen generated from methanolysis of AB catalyzed by the Ag ₃₀ Pd ₇₀ /C catalyst at various
				temperatures. (b) Arrhenius plot, i.e., a plot of ln k versus $1/T$ ([AB] = 100 mM, [AgPd] = 3.2 mM).
				tration of 100 mM at 25 \pm 1 °C. The plots of hydrogen genera
Catalyst	TOF/min^{-1}		Ref.	tion from the methanolysis of the AB solution in the presence of different dosages of $Ag_{30}Pd_{70}/C$ NPs are shown in Fig. 6(a). The
		$E_{\rm a}/{\rm kJ}~{\rm mol}^{-1}$		slope of the linear initial portion of each plot was chosen as the
RuCl ₃	150		6	rate of hydrogen generation corresponding to the respective
NiCl ₂	2.9		6	catalyst concentration. As shown in Fig. 6(b), a plot of the
PdCl ₂	1.5		6	natural logarithm of the rate of hydrogen generation versus the
Cu@Cu ₂ O	0.16		12	
$Co-Ni-B$	10.0		13	natural logarithm of the concentration of $Ag_{30}Pd_{70}$ was found to
PVP-Pd	22.3 30	35 ± 2 40.04	33 34	be nearly linear, with a slope of 1.16. The linear nature of this
Rh/zeolite Ru/MMT	90.9	23.8 ± 1.2	9	plot indicated that the methanolysis of AB followed nearly first
PVP-Ru	66.95	58 ± 2	35	order kinetics with respect to the concentration of $Ag_{30}Pd_{70}/C$.
CoPd/C	27.7	25.5	15	To obtain the activation energy of the methanolysis of AE
Mesoporous Cu	2.41	34.2 ± 1.2	36	catalyzed by AgPd/C NPs, reactions at various temperatures
CuPd/C	53.2	35.4	16	
$Cu-Cu2O-CuO/C$	24	67.9	14	from 0-30 °C were carried out. The values of the rate constant λ
Ru/graphene	99.4	54.1 \pm 2	11	at these temperatures were calculated from the slope of the
PVP-Ni	12.1	63 ± 2	37	linear part of each plot shown in Fig. $7(a)$. The Arrhenius plot
Rh/nano HAP	147	56 ± 2	38	i.e., ln k vs. $1/T$, for the catalyst is shown in Fig. 7(b), from which
Rh/nanosilica	168	49 ± 2	39	the apparent activation energy was determined to be approxi- mately 37.5 kJ mol ⁻¹ . Table 2 summarizes performance features

To obtain the activation energy of the methanolysis of AB catalyzed by AgPd/C NPs, reactions at various temperatures from 0–30 °C were carried out. The values of the rate constant k at these temperatures were calculated from the slope of the linear part of each plot shown in Fig. 7(a). The Arrhenius plot, *i.e.*, $\ln k \nu s$. 1/*T*, for the catalyst is shown in Fig. 7(b), from which the apparent activation energy was determined to be approximately 37.5 kJ mol $^{-1}$. Table 2 summarizes performance features of the various catalysts towards the methanolysis of AB at room temperature. The $Ag_{30}Pd_{70}/C$ catalyst was determined to be the

Fig. 8 TEM images of the Ag₃₀Pd₇₀/C catalyst before (a) and after (b) eight catalytic runs. (c) Catalytic performance versus the number of catalytic runs for the Ag₃₀Pd₇₀/C catalyst ([AgPd] = 3.2 mM, [AB] = 100 mM).

most active catalyst, even better than Ru¹¹- and Rh^{38,39}-based catalysts based on comparing their TOF values.

3.4 Reusability of the $\text{Ag}_{30}\text{Pd}_{70}/\text{C}$ catalyst

The reusability of a catalyst is crucial for practical applications. After the first run of methanolysis of AB (1 mmol) catalyzed by 3.2 mM NPs, the catalyst was kept in the solution and another equivalent amount of AB was added to the reaction system. The gas generation was monitored and the same procedure was repeated. Fig. 8(c) shows the hydrogen productivity versus reaction time for the generation of hydrogen from the AB solution catalyzed by $Ag_{30}Pd_{70}/C$ at 8 recycles under ambient conditions. The activity of $Ag_{30}Pd_{70}/C$ catalysts decreased with the number of recycles, and the TOF value dropped from 366.4 to 248.9 min⁻¹ after the eight runs. The TEM images shown in Fig. 8(a) and (b) indicated that there was no noticeable change either in NP size or morphology, so the decreased activity of the $Ag_{30}Pd_{70}/C$ catalysts may be attributed to the loss of metal from the surface of the carbon or the deactivation effect of the increasing metaborate concentration during the methanolysis of AB. To examine this possibility, supernatant liquid after the reusability tests was filtered to remove the support. ICP analysis of the solution indicated the presence of a small amount of Ag. RSC Advances

This causing is earlier catalytic condition and μ_0 . This article is the resulting on the causing intermediation and μ_0 . This article is the resulting of the taggethout the sumption of the resulting o

4. Conclusions

In summary, we have reported the synthesis of monodisperse AgPd alloy NPs by the coreduction of silver nitrate and palladium chloride in oleylamine. These NPs exhibited notable activity towards hydrogen generation from the methanolysis of AB at room temperature due to the uniform size of the particles on the support and the synergistic effects between Ag and Pd. The catalytic activity was found to be closely related to the alloy composition, and the $Ag_{30}Pd_{70}$ NPs were shown to be the most active, with a TOF value of 366.4 min^{-1} . The kinetic studies that we carried out on these alloy NPs indicated the catalytic methanolysis of AB to be first-order with respect to the catalyst concentration. The activation energy was determined to be 37.5 kJ mol $^{-1}$. Moreover, the Ag $_{30}$ Pd $_{70}$ /C catalyst retained 67% of its initial activity after 8 catalytic cycles. Their tunable catalytic properties shown here indicate that these AgPd alloy NPs have great potential in developing AB as a hydrogen storage material for fuel cell applications.

Acknowledgements

This work was supported by NSFC project numbers 21476187 and 21536010.

References

- 1 L. Schlapbach and A. Züttel, Nature, 2001, 414, 353-358.
- 2 U. Eberle, M. Felderhoff and F. Schueth, Angew. Chem., Int. Ed., 2009, 48, 6608–6630.
- 3 J. Yang, A. Sudik, C. Wolverton and D. J. Siegel, Chem. Soc. Rev., 2010, 39, 656–675.
- 4 T. B. Marder, Angew. Chem., Int. Ed., 2007, 46, 8116–8118.
- 5 F. H. Stephens, V. Pons and R. T. Baker, Dalton Trans., 2007, 25, 2613–2626.
- 6 P. V. Ramachandran and P. D. Gagare, Inorg. Chem., 2007, 46, 7810–7817.
- 7 S. Satyapal, J. Petrovic, C. Read, G. Thomas and G. Ordaz, Catal. Today, 2007, 120, 246–256.
- 8 M. Diwan, D. Hanna and A. Varma, Int. J. Hydrogen Energy, 2010, 35, 577–584.
- 9 H. B. Dai, X. D. Kang and P. Wang, Int. J. Hydrogen Energy, 2010, 35, 10317–10323.
- 10 K. V. Kordesch and G. R. Simader, Chem. Rev., 1995, 95, 191– 207.
- 11 S. G. Peng, J. C. Liu, J. Zhang and F. Y. Wang, Int. J. Hydrogen Energy, 2015, 40, 10856–10866.
- 12 S. B. Kalidindi, U. Sanyal and B. R. Jagirdar, Phys. Chem. Chem. Phys., 2008, 10, 5870–5874.
- 13 S. B. Kalidindi, A. A. Vernekar and B. R. Jagirdar, Phys. Chem. Chem. Phys., 2009, 11, 770–775.
- 14 M. Yurderi, A. Bulut, I. E. Ertas, M. Zahmakiran and M. Kaya, Appl. Catal., B, 2015, 165, 169–175.
- 15 D. H. Sun, V. Mazumder, Ö. Metin and S. H. Sun, ACS Catal., 2012, 2, 1290–1295.
- 16 P. Y. Li, Z. L. Xiao, Z. Y. Liu, J. L. Huang, Q. B. Li and D. H. Sun, Nanotechnology, 2015, 26, 025401.
- 17 D. S. Wang and Y. D. Li, Adv. Mater., 2011, 23, 1044–1060.
- 18 R. G. Chaudhuri and S. Paria, Chem. Rev., 2012, 112, 2373– 2433.
- 19 M. Zahmakıran and S. Özkar, Nanoscale, 2011, 3, 3462-3481.
- 20 V. Mazumder, M. F. Chi, M. N. Mankin, Y. Liu, O. Metin, D. H. Sun, K. L. More and S. H. Sun, Nano Lett., 2012, 12, 1102–1106.
- 21 J. Park, E. Lee, N. M. Hwang, M. Kang, S. C. Kim, Y. Hwang, J. G. Park, H. J. Noh, J. Y. Kim, J. H. Park and T. Hyeon, Angew. Chem., Int. Ed., 2005, 44, 2872–2877.
- 22 S. Zhang, Ö. Metin, D. Su and S. H. Sun, Angew. Chem., Int. Ed., 2013, 52, 3681–3684.
- 23 O. Metin, S. F. Ho, C. Alp, H. Can, M. N. Mankin, ¨ M. S. Gültekin, M. F. Chi and S. H. Sun, Nano Res., 2013, 6, 10–18.
- 24 Ö. Metin, X. L. Sun and S. H. Sun, Nanoscale, 2013, 5, 910– 912.
- 25 D. H. Chen and C. J. Chen, J. Mater. Chem., 2002, 12, 1557– 1562.
- 26 Y. T. Ji, S. C. Yang, S. W. Guo, X. P. Song, B. J. Ding and Z. M. Yang, Colloids Surf., A, 2010, 372, 204–209.
- 27 F. F. Lu, D. H. Sun, J. L. Huang, M. M. Du, F. Yang, H. M. Chen, Y. L. Hong and Q. B. Li, ACS Sustainable Chem. Eng., 2014, 2, 1212–1218.
- 28 E. V. Carino, M. R. Knecht and R. M. Crooks, Langmuir, 2009, 25, 10279–10284.
- 29 S. W. Han, Y. Kim and K. Kim, J. Colloid Interface Sci., 1998, 208, 272–278.
- 30 D. S. Lee and Y. W. Chen, J. Taiwan Inst. Chem. Eng., 2013, 44, 40–44.

- 31 P. Kamakoti, B. D. Morreale, M. V. Ciocco, B. H. Howard, R. P. Killmeyer, A. V. Cugini and D. S. Sholl, Science, 2005, 307, 569–573. Paper

11 P. Kamislovi, B. D. Mortrelle. M. V. Clocko, B. H. Howard, 35 H. Bridgean, Ö. Metin and S. Galar, Cetal. 70day, 2011, 170

R. P. Clocko, P. V. Clocko, P. Clocko, P. Clocko, M. Kharingi, 2. S. Clock and Licence. 2
	- 32 C. Q. Sun, Y. Wang, Y. G. Nie, B. R. Mehta, M. Khanuja, S. M. Shivaprasad, Y. Sun, J. S. Pan, L. K. Pan and Z. Sun, Phys. Chem. Chem. Phys., 2010, 12, 3131–3135.
	- 33 H. Erdogan, Ö. Metin and S. Özkar, Phys. Chem. Chem. Phys., 2009, 11, 10519–10525.
	- 34 S. Caliskan, M. Zahmakiran and S. Özkar, Appl. Catal., B, 2010, 93, 387–394.
- 35 H. Erdogan, Ö. Metin and S. Özkar, Catal. Today, 2011, 170, 93–98.
- 36 Q. L. Yao, M. Huang, Z. H. Lu, Y. W. Yang, Y. X. Zhang, X. S. Chen and Z. Yang, Dalton Trans., 2014, 44, 1070–1076.
- 37 D. Özhava, N. Z. Kılıçaslan and S. Özkar, Appl. Catal., B, 2015, 162, 573–582.
- 38 D. Özhava and S. Özkar, Int. J. Hydrogen Energy, 2015, 40, 10491–10501.
- 39 D. Özhava and S. Özkar, Appl. Catal., B, 2016, 181, 716-726.