Synthesis of palladium-rhodium bimetallic nanoparticles for formic acid dehydrogenation

Ilaria Barlocco, Sofia Capelli, Elisa Zanella, Xiaowei Chen, Juan J. Delgado, Alberto Roldan, Nikolaos Dimitratos, Alberto Villa

1. Introduction

Noble metals alloyed nanoparticles (NPs) are fundamental for several essential industrial processes. Although academia is switch-ing its attention on cheaper and abundant transition metals, indus-tries continue to employ noble metal-based catalysts due to their higher activity, stability and their well comprehended chemistry [1,2]. For these reasons a key challenge is to modify these cata-lysts, increasing their catalytic properties such as activity, selectiv-ity and stability. In particular, palladium-based catalysts are widely employed in different types of heterogeneous catalytic reactions, such as hydrogenations [3] and oxidations [4,5].

The addition of a second metal to Pd (Au, Ag, Ni, etc.) can tune the catalytic properties of the material, often enhancing activity, selectivity and stability of the catalyst due to the formation of new electronic states (ligand effect) combined with the strain of the lattice (steric effects) [6,7]. Rhodium also plays a fundamental role in many reactions such as the C–H activation [8]. The Pd-Rh system is barely studied. However, it was reported that alloying Rh to Pd it is possible to enhance the activity and stability of pure Pd in different reactions, including hydrogenation of cyclohexene [9], oxida-tion of o-phenylenediamine [10] and ethanol steam reforming [11]. The problem is that it is difficult to obtain well-defined bimetallic Pd-Rh structure. Indeed, Pd and Rh are immiscible at the thermal equilibrium bulk state over the whole composition range, therefore the alloy is in a segregated state at room temperature [12]. For this reason, alloys formed by these two metals have been ob-tained only in a metastable state by quenching the mixture at ex-tremely high temperature, above 1000 K [13]. In the last decade, it

Keywords:
- Palladium
- Rhodium
- Alloy
- Formic acid
- Hydrogen
- Muconic acid

Herein, we report for the first time the synthesis of preformed bimetallic Pd-Rh nanoparticles with different Pd:Rh ratios (nominal molar ratio: 80–20, 60–40, 40–60, 20–80) and the corresponding Pd and Rh monometallic ones by sol immobilization using polyvinyl alcohol (PVA) as protecting agent and NaBH₄ as reducing agent, using carbon nanofibers with high graphitization degree (HHT) as the desired support. The synthesized catalysts were characterized by means of Transmission Electron Microscopy (TEM) and inductively coupled plasma optical emission spectroscopy (ICP-OES). TEM shows that the average particle size of the Pd-Rh nanoparticles is in the range of 3–4 nm, with the presence of few large agglomerated nanoparticles. For bimetallic catalysts, EDX-STEM analysis of individual nanoparticles demonstrated the presence of random-alloyed nanoparticles even in all cases Rh content is lower than the nominal one (calculated Pd:Rh molar ratio: 90–10, 69–31, 49–51, 40–60). The catalytic performance of the Pd-Rh cat-alysts was evaluated in the liquid phase dehydrogenation of formic acid to H₂. It was found that Pd-Rh molar ratio strongly influences the catalytic performance. Pd-rich catalysts were more active than Rh-rich ones, with the highest activity observed for Pd₀Rh₁₀ (1792 h⁻¹), whereas Pd₉₀Rh₁₀ (921 h⁻¹) resulted the most stable during recycling tests. Finally, Pd₀Rh₁₀ was chosen as a representative sample for the liquid-phase hydrogenation of muconic acid using formic acid as hydrogen donor, showing good yield to adipic acid.
was report that Pd-Rh alloyed system can be obtained at low tem-perature, mixing the two metal at the atomic level, using a poly-ols method where poly(N-vinyl-2-pyrrolidone) (PVP) acts as stabi-lizer and ethylene glycol as reducer [9,14]. Ham and co-workers used theoretical calculations to demonstrate that the addition of Rh to Pd enhance the activity of pure Pd in the formic acid dehy-drogenation, enhancing the selectivity to H₂ and avoiding the for-mation of CO which acts as poison for Pd catalysts [15]. The ben-eﬁcial effect obtained by introducing Rh to Pd was attributed to a contraction of the Pd-Pd distance and the increase in the elec-tron density in surface Pd atoms compared to pure Pd [15]. Formic acid, that can be produced from lignocellulosic biomass, has at-tracted extensive attention as a promising renewable liquid-phase hydrogen donor due to properties such as high volumetric hydro-gen content (4.4%), liquid state at room temperature, high stability, environmental benignity and non-toxicity [16,17]. Formic acid de-composition in the liquid phase has been studied using a range of heterogeneous catalysts [18]. Heterogeneous catalysts, based on supported metal nanoparticles, such as, Pd, Au and the correspond-ing bimetallic systems have been reported in both the past and re-cent years [19,20]. However, the preparation of an active, selective and above all stable catalyst still represents a challenge.

Our group has a good experience in the preparation of bimetal-lic systems (e.g. Au-Pd, Au-Pt, Au-Ru) using the immobilization of preformed metal nanoparticles with an alloy structure [21]. In this work, we utilized this technique to prepare Pd-Rh nanoparticles using polyvinyl alcohol (PVA) as protecting agent and NaBH₄ as reducing agent. We choose highly graphitized carbon nanofibers (HHT) as support. Indeed, we have recently reported that the uti-lization of these materials can improve the catalytic activity of sup-port Pd nanoparticles compared to other carbonaceous supports [22]. The synthesized catalysts were tested in the decomposition of formic acid to produce hydrogen in liquid phase reaction condi-tions. The structure of the catalysts before and after reaction was carefully studied by means of electron microscopy. The most ac-tive catalysts were then tested in the liquid-phase hydrogenation of muconic acid in presence of formic acid as hydrogen donor. This is an important reaction to obtain bio-adipic aci d starting from re-
newable resources and using the concept of in situ hydrogen gener-ation using high content H donors like formic acid, at mild reaction conditions [23,24].

2. Experimental

2.1. Materials and chemicals

Rhodium(II) acetate (Rh(CO₂CH₃)₂) was purchased from Alfa Aesar (99.99%) and sodium tetrachloropalladate (II) (Na₂PdCl₄, 99.99%), sodium borohydride (NaBH₄, 99.99%) and poly(vinyl al-cohol) (PVA, average molar weight 10,000, 87%-89% hydrolyzed) were used without any pre-treatment for the catalysts synthesis and they were purchased from Sigma-Aldrich (Haverhill, MA, USA). All the catalytic tests were carried out using Formic acid (98.5%, Sigma-Aldrich) and deionized water and ethanol (95%, Sigma-Aldrich) as the substrate and the solvent, respectively. CNFs PR24-HHT (High Heat Treated carbon nanofiber) were obtained from Applied Science Company (Cedarville, OH, USA).

2.2. Catalysts preparation

In a typical sol immobilization procedure [25], the catalyst was prepared adding the solution of the metal precursor salts (Na₂PdCl₄, [Rh(CO₂CH₃)₂]₂; Table 1) and the capping agent PVA (M/PVA=1/0.5 w/w) in 0.5 L of H₂O/EtOH=1/1 vol/vol. Then, a fresh aqueous solution of NaBH₄ (M/NaBH₄=1/8 (mol/mol)) was added at once. The colloidal metal solution was supported on HHT.

<table>
<thead>
<tr>
<th>Pd-Rh ratio</th>
<th>Pd (mol mol⁻¹)</th>
<th>Rh (mol mol⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pd</td>
<td>4.70</td>
<td>–</td>
</tr>
<tr>
<td>Pd₈Rh₂</td>
<td>3.78</td>
<td>0.95</td>
</tr>
<tr>
<td>Pd₆Rh₆</td>
<td>2.86</td>
<td>1.90</td>
</tr>
<tr>
<td>Pd₄Rh₈</td>
<td>1.92</td>
<td>2.88</td>
</tr>
<tr>
<td>Pd₂Rh₁₀</td>
<td>0.97</td>
<td>3.86</td>
</tr>
<tr>
<td>Rh</td>
<td>–</td>
<td>4.86</td>
</tr>
</tbody>
</table>

CNFs (0.495 g) under vigorous stirring. By using sulphuric acid, the suspension was acidified to pH 2 and stirred for 30 min in order to ensure the full immobilization of the nanoparticles on the support. The solid was filtered, washed with 0.5 L of H₂O and dried in oven at 80 °C for 1 day.

2.3. Catalytic tests

Formic acid dehydrogenation: Liquid-phase Formic acid decom-position was carried out in a two-necked 100 mL round-bottom flask placed in a water bath equipped with a magnetic stirrer and a reflux condenser.

Typically, 10 mL of an aqueous solution 0.5 M of HCOOH was placed in the reactor and the reaction temperature was set-up at 30 °C. The formic acid/metal molar ratio is 2000:1 and once the solution reached the desired temperature, the solution was stirred at 1400 rpm. At these experimental conditions we have shown pre-viously we operate under kinetic regime [22]. Recycle tests were performed on the most active catalysts using 25 mL of an aqueous solution of HCOOH 0.5 M in a 250 mL round-bottom flask at the same experimental conditions of the reaction previously discussed.

Muconic acid (MA) hydrogenation: 25.3 mg of MA were placed in a round bottom flask with 15 mL of distilled water. Then 32 mg were added in 10 mL flask to have a MA/FA ratio of 1/4 (mol/mol). Water was used as solvent for both the solutions. FA was therefore added to MA solution. The final concentration of MA was 0.007 M. Finally, the catalyst was added with a MA/metal ratio of 200/1 (mol/mol). The reactions were performed at 30 °C with a stirring rate of 1400 rpm. The reactor was left open to facilitate FA decom-position toward the production of H₂. Periodic withdrawals were performed at different reaction times and the products were anal-ysed by High Performance Liquid Chromatography (HPLC).

Product analysis: Formic acid and muconic acid conversion were analysed using high-performance liquid chromatography (HPLC). In particular an H+ chromatographic column was used (Alltech OA- 10,308, 300 mm 7.8 mm) with UV detector settled at 210 nm. Liquid samples were withdrawn periodically (200 µL) and diluted to 5 mL with H₃PO₄ solution (0.1 wt%) which was also the eluent of the analysis. The isocratic eluent flow was set at 0.4 mL min⁻¹. The gas phase was analyzed by gas chromatography (Agi-lent, mod. 7890) equipped with a TCD detector and a proper set up for the quantification of H₂ and CO.

2.4. Catalyst characterization

Transmission electron microscopy (TEM) experiments were per-formed on a double Cs aberration-corrected FEI Titan™ Themis 60–300 microscope equipped with a monochromator, a X-FEG gun and a high efficiency XEDS ChemiSTEM, which consists of a 4-windowless SDD detectors. HR-STEM imaging was performed at 200 kV and using a high-angle annular dark-field (HAADF) detect-tor with a camera length of 11.5 cm. The HAADF-STEM technique is sensitive to the atomic number of the elements, whose inten-sity is roughly proportional to the square of the atomic number (Z²), and it makes possible to distinguish small nanoparticles sup-ported on light supports. XEDS mappings were performed using
Table 2. Nominal and calculated Pd-Rh molar ratio and average particle size for the Pd-Rh catalysts.

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>Nominal Pd-Rh molar ratio</th>
<th>Calculated Pd-Rh molar ratio (EDX-TEM)</th>
<th>Calculated Pd-Rh molar ratio (ICP)</th>
<th>Average particle size (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pd</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>3.2 ± 0.8</td>
</tr>
<tr>
<td>Pd used</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>4.5 ± 1.3</td>
</tr>
<tr>
<td>Pd₈ Rh₂</td>
<td>80:20</td>
<td>90:10</td>
<td>87–13</td>
<td>2.9 ± 0.9</td>
</tr>
<tr>
<td>Pd₈ Rh₂ used</td>
<td>80:20</td>
<td>95:5</td>
<td>93–7</td>
<td>4.5 ± 1.0</td>
</tr>
<tr>
<td>Pd₆ Rh₄</td>
<td>60:40</td>
<td>69:31</td>
<td>73–27</td>
<td>2.5 ± 0.6</td>
</tr>
<tr>
<td>Pd₆ Rh₄ used</td>
<td>60:40</td>
<td>81:19</td>
<td>84–16</td>
<td>3.0 ± 0.7</td>
</tr>
<tr>
<td>Pd₄ Rh₆</td>
<td>40:60</td>
<td>49:51</td>
<td>47–53</td>
<td>3.4 ± 1.3</td>
</tr>
<tr>
<td>Pd₄ Rh₆</td>
<td>20:60</td>
<td>40:60</td>
<td>38–62</td>
<td>2.6 ± 0.7</td>
</tr>
<tr>
<td>Rh</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>3.5 ± 1.1</td>
</tr>
</tbody>
</table>

Metal loading and metal leaching were analyzed by inductively coupled plasma optical emission spectroscopy (ICP-OES) using a Perking Elmer Optima 8000 emission.

3. Results

Pd₈Rh₆/HHT bimetallic catalysts with different Pd:Rh atomic ratio, and the corresponding Pd and Rh monometallic ones were prepared by sol immobilization method using polyvinyl alcohol (PVA) as protective agent (Table 2). Compared to the classical preparation method [22] where pure water is used as solvent, a mixture of ethanol and water was used due to the low solubility of [Rh(CO₂CH₃)]₂ in water.

3.1. TEM-STEM results

The catalysts were characterized at the sub-nanometric scale by transmission electron microscopy. TEM image of the monometallic Pd/HHT was reported elsewhere [22] and a STEM image of Rh/HHT is reported in Fig. S1 in the supporting information. Both monometallic catalysts showed a similar average particle size of 3.2 and 3.5 nm, respectively (Table 2) with particles well dispersed on the surface of the carbon nanofibers. Figs. 1–4 show representative results obtained by STEM and investigation of the Pd-Rh bimetallic systems. The catalysts show a similar average particle size in the range of 2.5–3.2 nm, Table 2, with a narrow particle size distribution, as shown in Figs. 1(c)–4(c). STEM-XEDS analysis was performed on individual particles to analyse their com-position (Figs. 1–4, Table 2). Line profiles of single particles are also included to illustrate the formation of bimetallic nanoparticles. Regarding sample with noPd₂Rh₆, (Fig. 1) an average particle size of 2.6 nm was obtained, although some big agglomeration of tiny crystallites was found in the sample (Fig. S2). Interestingly, the particles included in these agglomerations have a low content of Rh (<10%) or even they are pure Pd (Fig. S2b). The small particles are always made of both Rh and Pd, with a Pd-Rh ratio different from particle to particle and average Pd-Rh molar ratio of 40:60 (Table 2, Fig. 1b).

Pd₄Rh₆ consists of particles with an average particle size of 3.2 nm (Table 2), with a broader particle distribution (Fig. 2). All the particles analysed were bimetallic with an average molar ratio of 49:51 (Pd:Rh), which indicates an enrichment of Pd, if we compare with the nominal composition (40:60). This composition also varies from particle to particle, (Fig. 2b). In addition, in this case, large aggregates were observed (Fig. S3a) with a lower content of Rh (Pd-Rh ratio of 79:21) in comparison with the one obtained for the small particles, (Fig. S3b).

Particles with an average size of 2.5 nm were found for Pd₆Rh₄ catalyst (Table 2), with the presence of large aggregations (Fig. S4). STEM-XEDS analysis (Fig. 3) confirms the presence of Pd-Rh bimetallic particles with an average molar ratio of 69:31, which again shows a higher Pd content that what it was expected. More-over, for this sample the Pd-Rh relative content varies from particle to particle (Fig. 3b).

Pd₄Rh₂ shows highly and homogeneously dispersed metal nanoparticles with an average size of 2.9 nm and no aggregates were observed (Fig. 4). Particles exhibit an enrichment of Pd, with an average Pd:Rh atomic ratio of 90:10, which is also higher than the nominal one of 80:20.

STEM-XEDS analysis performed on individual particles demonstrated that almost all the particles were Pd-Rh bimetallic, although the content of Rh is lower than the nominal one in all the prepared catalysts. Additionally, monometallic Rh and Pd particles were not observed. In order to demonstrate the formation of the bimetallic nanoparticles, EDS line profiles of isolated nanoparticles were collected (Figs. 1–4g–h)). In all the analysed nanoparticles, we can observe the signals of Rh and Pd inside of the individual particles. Although the presence of really small particle size of 2–6 nm complicated the analysis, the obtained line profiles may indicate a relatively homogeneous distribution of both metal and that there is not any preferential distribution of the elements such core-shell structure. To understand the discrepancy between nominal and calculated Pd:Rh atomic ratio, we utilized the inductively coupled plasma optical emission spectroscopy (ICP-OES) to analyse the metal loading of the synthesised catalysts (Table 2). The analysis performed on the filtrated solution, after the immobilization of the metal colloid on the carbon nanofibers, revealed the presence of Rh in solution. The same problem has been detected for the monometallic Rh catalyst which shows a calculated leading of 0.91% instead of the nominal one of 1%.

3.2. Catalytic activity

The catalytic performance of the Pd-Rh catalysts was evaluated in the liquid-phase decomposition of formic acid as a model reaction for hydrogen generation (Reaction conditions, 0.5 M HCOOH in water, 30 °C, stirring rate of 1400 rpm, substrate/metal molar ratio of 2000). The decomposition of formic acid follows two main pathways, (i) dehydrogenation (HCOOH → CO₂ + H₂, G=−48.4 kJ mol⁻¹) and (ii) dehydration (HCOOH → CO + H₂O, G=−28.5 kJ mol⁻¹). The reaction conditions, in particular stirring rate and substrate/metal molar ratio were optimised in previous reports, in or-der that the reaction is taking place in chemical kinetic regime [22]. Fig. 5 presents the catalytic activity of the catalysts prepared by varying the Pd-Rh atomic ratio. To better correlate the effect...
of the metal ratio and the activity, the calculated Pd-Rh instead of the nominal molar ratio will be now considered. The activity is expressed as mol of formic acid reacted per total mol of metal per hour and calculated after 5 min of reaction. The high-est initial activity was obtained using Pd$_{90}$Rh$_{10}$ (1793 h$^{-1}$), fol-lowed by monometallic Pd and Pd$_{69}$Rh$_{31}$ which showed similar results (979 and 921 h$^{-1}$, respectively). Increasing the Rh con-tent, the activity progressively decreased and the monometallic Rh showed very low activity (Fig. 5). However, this trend was not fol-lowed by Pd$_{40}$Rh$_{60}$ which showed an activity two times higher (696 h$^{-1}$) than Pd$_{48}$Rh$_{52}$ (341 h$^{-1}$). We could not find the reason to fully justify the observed catalytic trend. Indeed, the two cat-alysts showed similar bimetallic structure, even the particle size is slightly larger for Pd$_{48}$Rh$_{52}$ (3.4 nm) than the other bimetallic catalysts (2.4–2.9 nm (Table 2). In general, the influence of particle size in catalyst activity for this series of catalysts seems to be negligible compared to that of Pd-Rh composition. Table 3 presents a comparison of the activity of various Pd based bimetallic catalysts for the liquid phase dehydrogenation of formic acid under mild conditions. Pd$_{90}$Rh$_{10}$ shows an activity better of most of the catalysts presented in the literature. Fig. 6 showed simi-
sis performed on the filtrated solution revealed the leaching of 5% of Pd during the reaction. Moreover, Pd active sites can be poisoned by the CO produced (11–15 ppm, in the different cycles). It is well known, indeed, that CO can poison Pd catalysts even when present at ppm level [26]. A similar trend was observed for Pd90Rh10, the most active catalyst. STEM analysis evidenced the coalescence of the particles with growing of the average particle size from 2.9 to 4.5 nm (Fig. 8c). The STEM-XEDS analysis showed an enrichment in Pd of the particles, which have an average composition of 95:5 with the presence of segregated monometallic Pd particles (Table 2). ICP results demonstrated that this is due to the partial leaching of Rh in the reaction solution, whereas no leaching of Pd was observed. On the contrary, Pd69Rh31 showed a good stability during the 6 cycles (Fig. 7c). STEM analysis of the used catalyst evidenced a limited growing of the average particle size from 2.5 to 3.0 nm (Table 2). The analysis of the atomic composition of the single nanoparticles showed an enrichment in Pd with a Pd-Rh atomic ratio of 81:19, (Fig. 9b), due to the leaching of Rh in the solution. The high stability might be attributed to a cata-lyst with a Pd-Rh composition similar to the most active ones. The higher activity of the new phase is offset by the leaching of metal in the solution, and the conversion was constant during the 6 cy-cles. The partial Rh leaching observed for the bimetallic catalysts can be due to the fact that Pd-Rh structures are not stable upon hydrogen absorption/desorption, leading to a partial segregation of the two metals [13,27]. Probably, segregated Rh is more prone to leaching than when alloyed.

Many studies have reported in the last 10 years the utilization of formic acid as hydrogen source for the hydrogenation of biomass derived molecules [16]. In this work we utilized formic acid for the upgrade of muconic acid (MA), a molecule which can be obtained from cellulose and lignin to produce bio-adipic acid (Scheme 1). The reaction proceeds in two-steps as we have recently shown [23]. In the first one, monounsaturated acids are produced ((2E)-hexenedioic acid and (3E)-hexenedioic acid), followed by the hydrogenation of the second C=C bond to obtain adipic acid.

Pt and Pd based catalysts have been used for this reac-tion [23,37], using molecular hydrogen. However, Pd based ones showed the best activity even under mild conditions. Therefore, we combine the ability of Pd catalyst to generate hydrogen from formic acid (H-source) and at the same time to activate H2 under mild conditions to hydrogenate muconic acid to bio-adipic acid. Muconic acid hydrogenation was performed under similar reaction conditions utilized for the dehydrogenation of formic acid (Reac-tion conditions, 0.007 M MA in water, MA/Formic acid ratio of 1/4 (mol/mol), 30 °C, stirring rate of 1400 rpm, substrate/metal molar ratio of 200). The two most active bimetallic catalysts, Pd90Rh10 and Pd69Rh31 were compared to monometallic Pd. In Fig. 10 it is
Fig. 3. (a) STEM-HAADF image of Pd₆Rh₄ catalyst, (b) particle size distribution, (c-f) STEM-XEDS of single nanoparticles (Pd-Rh atomic ratio: (1) 40:60, (2) 62:38, (3) 89:11, (4) 75:25, (5) 67:33, (6) 62:38) and (g-h) line profile of the particles 2 and 3, respectively.

Table 3. Comparison of catalytic activity of Pd-based bimetallic catalysts for the liquid phase dehydrogenation of formic acid under mild conditions.

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>T(°C)</th>
<th>Reagent</th>
<th>TOF (h⁻¹)</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Initial</td>
<td>2 h</td>
</tr>
<tr>
<td>Pd₆₉Rh₁₁/HHT</td>
<td>30</td>
<td>Formic acid (0.5 M)</td>
<td>1793</td>
<td>This work</td>
</tr>
<tr>
<td>Pd₆₉Rh₁₁/HHT</td>
<td>30</td>
<td>Formic acid (0.5 M)</td>
<td>921</td>
<td>This work</td>
</tr>
<tr>
<td>Pd/HHT</td>
<td>21</td>
<td>Formic acid (0.5 M)</td>
<td>979</td>
<td>This work</td>
</tr>
<tr>
<td>Pd/C</td>
<td>21</td>
<td>Formic acid (1.33 M)</td>
<td>18</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>15</td>
<td></td>
</tr>
<tr>
<td></td>
<td>30</td>
<td></td>
<td>48</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>28</td>
<td></td>
</tr>
<tr>
<td>Au₄Ag₄Pd₆₉/C</td>
<td>35</td>
<td>Formic acid (1 M)</td>
<td>230</td>
<td>[29]</td>
</tr>
<tr>
<td>Ag/Pd alloy (1:1)</td>
<td>30</td>
<td>Formic acid (1 M)</td>
<td>156</td>
<td>[30]</td>
</tr>
<tr>
<td>Ag/Pd alloy (1:1)</td>
<td>20</td>
<td></td>
<td>144</td>
<td>[30]</td>
</tr>
<tr>
<td>Ag₄AuPd₆₉</td>
<td>50</td>
<td>Formic acid (1 M)</td>
<td>382</td>
<td>[31]</td>
</tr>
<tr>
<td>Pd-MnOₓ/SiO₂ –NH₂</td>
<td>20</td>
<td>Formic acid (0.265 M)</td>
<td>140</td>
<td>[32]</td>
</tr>
<tr>
<td></td>
<td>50</td>
<td></td>
<td>1300</td>
<td></td>
</tr>
<tr>
<td>Ag₄AuPd₆₉/GO</td>
<td>25</td>
<td>Formic acid</td>
<td>105</td>
<td>[33]</td>
</tr>
<tr>
<td>PdNiAg/C</td>
<td>50</td>
<td>Formic acid (0.175 M)</td>
<td>85</td>
<td>[34]</td>
</tr>
<tr>
<td>Pd₅₀Cu₅₀/resin</td>
<td>75</td>
<td>HCOOH/HCOONa=9:1</td>
<td>810</td>
<td>[35]</td>
</tr>
<tr>
<td>20 wt% PdAu/C–CeO₂</td>
<td>102</td>
<td></td>
<td>832</td>
<td>[36]</td>
</tr>
</tbody>
</table>
plotted the concentration of the reactant MA, the mono unsatu-rated intermediates and adipic acid as a function of reaction time. All the catalysts were highly active in the first step, almost quan-titatively converting MA to the intermediates after 1 h of reaction. However only the two bimetallic catalysts were able to convert the intermediates to adipic acid, in particular in the case of Pd$_{69}$Rh$_{31}$. The lower activity of Pd can be attributed to the lower activity and stability in the generation of hydrogen from formic acid compared to bimetallic counterparts. However, it can be observed that after 2 h of reaction the concentration of the products remains stable. Future studies will be performed to improve the catalytic perfor-mance of the present catalysts and to understand the reason of de-
activation in the muconic acid hydrogenation in presence of formic acid as hydrogen donor.

4. Conclusions

A series of preformed colloidal Pd$_x$Rh$_y$ nanoparticles and the corresponding monometallic supported Pd and Rh nanoparticles were prepared by sol immobilization method and deposited on highly graphitized carbon nanofibers (HHT). Using this technique we report for the first time, the ability to prepare preformed bimetallic random alloyed Pd-Rh nanoparticles with average particle size in the range of 2–4 nm even at low temperature as confirmed by STEM-HAADF and STEM-XEDS analyses. However, the composition of the particles is not homogeneous from particle to particle and a general enrichment in Pd was observed compared to the nominal Pd-Rh ratio. These results were attributed to the unsuccessful quantitative immobilization of Rh, as evidenced by ICP analysis. Bimetallic catalysts showed good activity in the dehydrogenation of formic acid in liquid phase, in particular for Pd-rich bimetallic systems (Pd$_{90}$Rh$_{10}$ and Pd$_{69}$Rh$_{31}$). Comparing the prod-
uct analysis of liquid and gas phase products, we concluded that the major reaction pathway towards the production of CO$_2$ and H$_2$ was obtained for all the catalysts, with minor formation of CO in the range of 5–15 ppm. However, in the case of monometallic Pd, traces of CO (11–15 ppm) were observed. A good stability during 6 consecutive runs was observed for Pd$_{60}$Rh$_{31}$ compared to monometallic Pd which rapidly deactivate. The low stability of this latter one was attributed to the growing of particle size due to coalescence of the particle, partial Pd leaching and presence of CO which acts as poison for Pd-based catalysts. Furthermore, monometallic Pd, Pd$_{60}$Rh$_{10}$ and Pd$_{60}$Rh$_{31}$ were tested in the mu-conic acid hydrogenation using formic acid as hydrogen donor, under mild conditions. The catalysts showed good activity for the first step of the reaction, towards the production of monounsaturated products. Only in the case of using bimetallic catalysts the second step of the reaction towards the formation of adipic acid was enhanced.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

References